The homeodomain transcription factor Phox2 in the stellate ganglion of the squid *Loligo pealei*

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ABSTRACT

Homeodomain transcription factors regulate development of embryos and cellular physiology in adult systems. Paired-type homeodomain genes constitute a subclass that has been particularly implicated in establishment of neuronal identity in the mammalian nervous system. We isolated fragments of eight homeodomain genes of this subclass expressed in the stellate ganglion of the North Atlantic long finned squid *Loligo pealei* (lp). For reasons of uniformity and clarity, *Loligo pealei* is used here. Of the most abundant ones, we cloned a full length cDNA which encoded the squid ortholog of the paired-type homeodomain proteins Phox2a/b. The homology of lpPhox2 to invertebrate and mammalian Phox2 was limited to the homeodomain. In contrast to mouse Phox2b, lpPhox2 was unable to transactivate the dopamine beta-hydroxylase (DBH) promoter in a heterologous mammalian transfection system. In *vivo*, lpPhox2 was expressed in the developing stellate ganglion of stage 27 squid embryos and continued to be expressed in the adult stellate neurons where expression was confined to the giant fiber lobe containing the neurons that form the giant axons. The expression of lpPhox2 was similarly timed and distributed as the Fmr gene. Furthermore, the Fmr upstream region contained putative Phox2a/b binding sites. These results suggest a role of lpPhox2 in the developmental specification of neuronal identity and regulation of neurons of the squid giant axon.

KEY WORDS: Homeodomain transcription factors, Squid, Phox2, FMRFamide, Stellate ganglion, Squid giant axon, Brain development

INTRODUCTION

The development of the nervous system involves multiple stages of cellular commitment that lead via patterning, proliferation, and fate determination to defined neuronal identities. These processes are driven by complex regulatory networks of transcription factors and signalling molecules (Andersson et al., 2006; Dasen et al., 2005; Jessell, 2000; Smidt and Burbach, 2007). Neuronal identity is often marked by a neurotransmitter phenotype, as well as its specific connectivity to other neurons. These two properties allow neurons to function in neural networks and to contribute specific circuitry.

Several families of transcription factors have been demonstrated to play a key role in the development of the mammalian nervous system. Early stages of commitment to neural cell fates and proliferation involve basic helix-loop-helix factors, while homeodomain proteins provide patterning and neuronal fate specification (Goridis and Brunet, 1999; Guillemin, 2007; Smidt and Burbach, 2007). For example, homeodomain proteins encoded by the Hox clusters in spinal cord, that are expressed in a segmental fashion, are involved in pattern formation (Dasen et al., 2005; Pearson et al., 2005; Trainor and Krumlauf, 2001). Non-Hox homeodomain genes participate in further stages of differentiation in all parts of the nervous system, mostly confined to specific cell groups that share neuronal identity (Asbreuk et al., 2002; Molyneaux et al., 2007). Such factors often retain expression in the mature nervous system and adopt regulatory functions (Davis et al., 2010; Zhu et al., 2005a). In the mammalian nervous system processes driven by homeodomain factors result in the development of highly complex neural structures. In the more simple nervous systems of lower vertebrates and invertebrates similar cascades of transcription factors also operate during nervous system development (Hammock et al., 2010; Sen et al., 2013). This suggests that functions of homeodomain factors are generic and may play a role in all nervous systems of the animal kingdom (Marder, 2002).

In the mammalian nervous system neuronal diversity is enormous so as to produce a nervous system that can meet the complex demands of the organism. In order to evaluate homeodomain gene function in the establishment of neuronal identity and connectivity, we turned to neurons that are more homogenous in anatomy and function, namely, those of the squid giant axon (Marder, 2002). The squid giant axon is a classical preparation in which fundamental features of neurons, such as cellular anatomy, axonal transport, control of membrane potentials and ion currents, and neurotransmission were first explored (Hodgkin and Huxley, 1952; Llinas, 1988; Webb and Young, 1940; Young, 1939). The squid giant axon is a thousand-fold larger in diameter than mammalian axons because it is formed by the fusion of axons from multiple neurons located in the giant fiber lobe of the stellate ganglion (Young, 1939). The ganglion contains two types of neurons. One set of small neurons in the giant fiber lobe provides axons which fuse into the giant axons that innervate the mantle muscles. The axons of these neurons, when excited by axo-axonic contacts, drive jet propulsion as part of the escape response of the squid (Gilly et al., 1990). In addition, they control water flow over the gills, serving a respiratory function. The second set consists of large neurons that control fin movement and fine control of swimming. The anatomy of the stellate ganglion and its afferent connectivity with the central nervous system of the squid have been extensively detailed in the species *Loligo pealei* since the 1930s (Young, 1939, 1973).

The involvement of Hox clusters in the outline of the embryonic body plan of the squid has been demonstrated (Arnold, 1990; Lee et al., 2003; Martin, 1965). However, the molecular mechanisms of neuronal fate determination are unknown. To explore such
mechanisms in the squid, we here aimed to identify and characterize homeodomain transcription factors expressed in the squid stellate ganglion. To this end a RT-PCR cloning strategy was adopted to identify paired-like homeodomain transcription factors in the stellate ganglion (Asbreuk et al., 2002), and the most abundant one was cloned and characterized with respect to expression and transcriptional activity. The results show that this factor is the squid homologue of Phox2 and that it may have a potential function in developmental specification of neuronal identity and regulation in the squid stellate ganglion.

RESULTS
Identification of homeodomain transcripts
To screen for homeobox genes expressed in the stellate ganglion we employed a PCR cloning strategy used previously to identify paired-type homeobox genes expressed in rodent brain (Asbreuk et al., 2002). Sequencing of clones identified eight PCR fragments of 120 nucleotides with similarity to mammalian homeodomain sequences. Similarities indicated that these were squid orthologs of the homeobox genes Msx1/2/3, Pitx2/3, Phox2a/b, ARX/Gsc, Nkx1, zinc finger homeodomain protein-4, and POU6-2/RPF-1 (supplementary material). Screening of a cDNA phage library constructed from adult stellate ganglia by filter hybridization, using labelled PCR fragments as probes, resulted in the isolation of clones for RPF-1 and Phox2, but none of the other homeodomains.

Structural properties of lpPhox2
A clone with a ~1800 nt insert was isolated from an adult stellate ganglion cDNA phage library. In addition, a library of about 23000 EST sequences that we obtained from the stellate ganglion of Loligo pealei (DeGiorgis et al., 2011) contained one entry with identical sequence to this cDNA, spanning nucleotides 240 to 652. Transcriptome sequences of the stellate ganglion and giant fiber system of this species fully matched the cloned sequence. One variant was always present in transcriptomes of Loligo pealei, a transcript that lacks a stretch of eight Ts in the untranslated 3' end (underlined in Fig. 1). This cDNA had a predicted open reading frame from nucleotides 442 to 1443 (Fig. 1) as indicated by the translation initiation algorithm of NetStart 1.0 (http://www.cbs.dtu.dk/services/NetStart/). There was a variant Kozak sequence present at the translation initiation site 442 (CUCAUCAUGG) (Kozak, 1999). This protein had a length of 333 amino acids with the 60 amino acid homeodomain between residues 132 and 191 (Fig. 1).

Comparison of the 60-amino acid homeobox sequence to other invertebrates and vertebrates showed a high degree of conservation over a wide range of species, from 100% for Sepia officinalis to 92% for Homo sapiens (Fig. 2). The high degree of sequence similarity identified the predicted protein as the Phox2 ortholog of Loligo pealei, lpPhox2. The cephalopod Sepia officinalis contained a predicted protein (accession: AGC24169) with 96% overall identity to lpPhox2. Differences were confined to a stretch of repeated Gln residues (Fig. 2). Orthologs of Phox2 were found in the
cephalopodian mollusks *Euprymna scolopes* and *Lottia gigantea*. The mollusks *Aplysia californica* and *Lymnaea stagnalis* shared a 98% similarity in the homeodomain with lpPhox2. However, the overall homology was only 53% due to low similarity outside the homeodomain. Small patches of similarity were present in the N-terminal part of Phox2 proteins, specifically MEYxYLN (residues 1–7), GMDSS (residues 20–24) and TxGSCxxxxxRDH (residues 62–75). Compared to vertebrate Phox2 proteins, lpPhox2 shared a strong homology in the homeobox (Fig. 2B) and the four amino acids preceding it (EKRK). Homology was also found in the extreme N-terminal motif MEYxYLN. The C-terminal domain lacked significant similarities (Fig. 2).

A lower degree of similarity in the homeodomain (88%) was found with Phox2 of other invertebrates, e.g. *C. elegans* (ceh-17) (Fig. 2). In the genome of the ctenophore *Mnemiopsis leidyi*, which is considered to belong to the most primitive metazoans (Marlow and Arendt, 2014), no evidence for a Phox2 ortholog was found. The most similar homeobox sequence to lpPhox2 rather appeared as an Aristaless ortholog. While vertebrates generally have two Phox2 paralogs, Phox2a and Phox2b, a single Phox2 sequence exists in databases of invertebrates. By mining *Loligo pealei* transcriptome and genome sequences, no evidence was obtained for Phox2 paralogs.

These data indicated that the squid *Loligo pealei* expresses a Phox2 transcription factor that is highly conserved in the homeodomain. The homeodomain confers the DNA-binding properties of this class of transcription factors, suggesting that the specificity of DNA-binding is conserved, and, therefore, essential property for the function of Phox2 proteins.

**Transcriptional properties of lpPhox2**

In view of the sequence divergence, we tested whether lpPhox2 could transcriptionally activate the human dopamine beta-hydroxylase (*DBH*) gene. The DBH promoter is an established mammalian Phox2 target gene which contains Phox2 response elements (Hwang et al., 2005; Kim et al., 1998). It has been shown that Phox2b and Phox2a of mouse, human and rat are transcriptional activators of the *DBH* gene. When a *DBH*-luciferase reporter construct was co-transfected with a mouse Phox2b expression plasmid in a heterologous cell system using Neuro2A cells, the DBH promoter activity was increased 10-fold. A *DBH* promoter-reporter construct with mutated response elements showed no significant change in promoter activity. However, when the mouse Phox2b expression plasmid was replaced by one for lpPhox2, DBH promoter activity was not influenced. These data show that lpPhox2 in contrast to mouse Phox2b lacks transcriptional activity in this heterologous mammalian cell system.

**Expression of lpPhox2**

The expression of *lpPhox2* was investigated in squid embryos and in the adult stellate ganglion by *in situ* hybridization and immunohistochemistry. Whole mount *in situ* hybridization on squid embryo of Arnold stage 27 (Arnold, 1990) revealed a signal in the stellate ganglion, while a sense probe did not show this staining (Fig. 3A,C). Whole mount *in situ* hybridization showed background staining in other head and body structures (Fig. 3). No *lpPhox2* mRNA signals were detected at earlier stages, although the stellate ganglion is present from stage 26 onwards. This suggested that *lpPhox2* was expressed during late embryonic development of the stellate ganglion. The stellate ganglion and giant fiber system is morphologically and functionally mature by stage 27, shown by axons leaving the ganglion to innervate mantle muscles (Fig. 3E,F) and the expression of phosphorylated neurofilament proteins in the giant axons (Fig. 3G,I). In cryosections of stage 30 embryos weak signals of *lpPhox2* transcripts were observed in the central nervous system in scattered patterns. Staining was inconsistent due to the low signal (not shown). This indicated that *lpPhox2* was expressed in brain structures, but at a low expression level. This observation agreed with a search in available transcriptomes of ganglia and brain lobe of *Loligo pealei* (http://ivory.idyll.org/blog/2014-loligo-transcriptome-data.html), showing the presence of lpPhox2 transcripts in the optical and buccal lobes, but not in the vertical lobe.

More intense signals of *lpPhox2* mRNA were obtained in the adult stellate ganglion (Fig. 3). We observed subregions of gene expression in the ganglion by *in situ* hybridization of β-tubulin (all neurons, and giant axon, Fig. 3G,1) and a transcript for a hypothetical protein homologous to mouse KIAA1109 (small giant fiber lobe neurons and giant axon, Fig. 3H). *lpPhox2* was expressed in small neurons in the giant fiber lobe (Figs 3J, 4A). Signals were far lower than those of β-tubulin (Fig 3H,1). Immunohistochemistry on isolated ganglia showed two types of lpPhox2 protein expression. One in nuclei of small neurons in the giant fiber lobe, and the other as non-nuclear staining in the central mass of the ganglion (Fig 4).
Relationship to Fmrf expression

We recently found that the Fmrf gene, coding for the neuropeptide FMRFamide, is highly expressed in the embryonic and adult stellate ganglion of Loligo pealei (Burbach et al., 2014). In the present study we noted similarities in the localization of lpPhox2 mRNA and Fmrf mRNA. Comparing the expression of lpPhox2 and Fmrf in the adult ganglion by in situ hybridization showed that both genes displayed a similar distribution of expression in the adult stellate ganglion. Both genes were expressed predominantly by the small neurons of the giant fiber lobe (Fig 5).

To investigate a potential involvement of lpPhox2 in the regulation of expression of the Fmrf gene, we determined the 5′ end of Fmrf transcripts by searching transcriptome databases of Loligo pealei nervous system lobes and a genome data dump of this squid (http://athyra.idyll.org/~t/blast/ceph/). In this way a short putative promoter region could be identified that contained two potential TATA boxes closely together (Fig 6). This sequence overlapped with a motif that resembles a Phox2a binding site in the human DBH gene promoter (Hwang et al., 2005), suggesting that Phox2a may regulate the expression of the Fmrf gene directly. Unfortunately, this suggestion could not be further tested functionally since lpPhox2 is not transcriptionally active in heterologous mammalian expression systems.

DISCUSSION

In many nervous systems homeodomain proteins play a developmental role in the differentiation and specification of neuronal systems. Moreover, they are engaged in regulatory processes when expression is sustained in the matured state. The latter function has not been explored extensively. In this study we identified seven paired-type homeobox genes homologous to mouse KIAA1109 in the giant fiber lobe of the stellate ganglion. (1) Magnifications of panel G showing neuronal expression of β-tubulin. (2) Neuronal expression of lpPhox2 in the adult giant fiber lobe. Abbreviations: sg, stellate ganglion; ga, giant axon; is, ink sac. Scale bars are 100 µm in panels A–E, 1 mm in G and H, and 10 µm in I and J.
Abbreviations: multiple tiles. In A and B sections from the same ganglion were used.

Individual axons from these neurons fuse to form the giant axons. (B) In situ hybridization of transcripts. Expression is highest in the small neurons of the giant fiber lobe. Phox2a/b ortholog in the adult stellate ganglion and lpPhox2 expression in the visceral neurons, both sensory and motoneurons, in all animal species, even in the ascidian Ciona intestinalis (D’Autréaux et al., 2011). These systems are particularly engaged in respiration. For Sepia officinalis it has been proposed that Phox2 serves the generation of stellate ganglionic neurons for a respiratory function (Nomaksteins et al., 2013).

In this respect it is of relevance to note the apparent coexpression of lpPhox2 and the Fmrf gene in small neurons of the giant fiber lobe of adult Loligo pealei. These neurons fuse their efferents into the giant axons that control mantle contraction, and enable physiological functions such as respiration and the escape response. The neurotransmitter of these neurons is not entirely certain; it has been hypothesized that glutamine and asparagine may be employed. We have shown that the Fmrf gene is expressed in these neurons, albeit the biologically active FMRF amide tetrapeptide is not produced due to differential precursor processing. Instead, the peptide is expressed in smaller fibers within the stellar nerves that innervate mantle muscles involved in respiration (Burbach et al., 2014). Several results from the present study suggest that the Fmrf gene is a target of lpPhox2. First, the onset of expression is timed similarly, around stage 27–28. Secondly, the location of expression overlaps for the two genes, and, thirdly, the putative promoter region of lpFmrf gene contains motifs that resemble Phox2a/b response elements of the hDBH promoter. However, this suggestion cannot be tested at present. Since lpPhox2 appeared to be functionally inactive in a mammalian cell line, we need to develop a homologous expression system, which is not available at present.

A noted feature of all invertebrate Phox2 proteins is that sequence conservation is exclusive to the homeodomain. Such restricted similarities have also been observed for other homeodomain orthologs. For example, species comparison of the Pitx family of paired-type homeodomain proteins displays a similar homology conservation is exclusive to the homeodomain. Such restricted similarities have also been observed for other homeodomain orthologs. For example, species comparison of the Pitx family of paired-type homeodomain proteins displays a similar homology.

The results suggest that transcriptional codes for the developmental specification and regulation of neuronal systems are at least in part conserved between mammals and squid. lpPhox2 may serve analogous functions in the development of the stellate ganglion.
neurons of Loligo pealei as Phox2a/b does in the regulation of neuronal development in mammals. Such conservation may render this ganglion and the giant fiber system as a useful model for addressing specific questions concerning neuronal development.

**MATERIALS AND METHODS**

**Animals**

Specimens of the North Atlantic long-finned squid Loligo pealei were caught in Vineyard Sound and obtained through the Marine Resources Centre of the Marine Biological Laboratory, Woods Hole, MA, USA. Catch, care and use of animal complied with all institutional and national rules. Egg mass deposits were obtained from squid kept in tanks with running seawater at 20°C and staged according to Arnold (1990). Embryos were harvested in seawater, fixed in 70% ethanol and stored at −20°C.

**Degenerate RT-PCR amplification of homeodomain cDNAs**

Total RNA was isolated from dissected squid stellate ganglia and subjected to RT-PCR using degenerate primers based on homologies within the homeodomain. Primer sequences, procedures and conditions were as described before (Asbreuk et al., 2002). PCR fragments were cloned and sequenced. PCR fragments encoding β-tubulin, a hypothetical protein homologous to mouse KIAA1109 and collagen were identified and cloned.

**cDNA libraries and cloning**

Stellate ganglia were dissected from adult squids and stored at −80°C. Poly(A+) RNA was isolated, size selected and multiple libraries were made in the lambda-ZAP Express vector by unidirectional cloning of cDNAs using the kits of Stratagene. Plague libraries were screened by filter hybridization with 32P-labeled homedomain fragments. Plasmids were excised from purified lambda clones and both strands were Sanger sequenced.

**In situ hybridization**

Whole mount in situ hybridizations were performed according to Harland (1991) with modifications (Wacker et al., 2004) on embryos fixed in MEM-buffered paraformaldehyde and stored in methanol. Samples were rehydrated, treated with protease K and acetylated in triethanolamine-buffered paraformaldehyde and kept in phosphate-buffered saline with 30% sucrose at 20°C. The lysate was used to assay luciferase activity.

**Competing interests**

The authors declare no competing or financial interests.

**Author contributions**

J.P.H.B. and H.C.P. conceived and designed the experiments, J.P.H.B. and A.J.C. performed the experiments, J.P.H.B. analysed the data, J.P.H.B. and P.G. wrote the paper.

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**Supplementary material**

Supplementary material available online at http://bio.biologists.orglookup/supplid=10.1242.bio.012476/-DC1

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