Immunohistochemical study of the nervous system of the tunicate *Thalia democratica* (Forsskal, 1775)

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Abstract

*Thalia democratica* is a cosmopolitan tunicate belonging to the Thaliacea class. To further investigate the anatomy of this species, immunohistochemical labelling was performed using anti-tubulin and anti-serotonin antibodies on specimens collected in the Mediterranean Sea. The anti-tubulin antibody stained the cilia of the endostyle, the pericoronal bands and of the gill bar, enabling a detailed description of these structures. Moreover, immunolabelling of the nervous system showed the presence of eight pairs of nerve fibres emerging from the neural ganglion. Serotonergic cells were observed in the distal tract of the intestine, along the pericoronal bands, and in the placenta of gravid blastozooids, as well as in the neural ganglion. The presence of serotonin in the central nervous system has also been reported in the larvae of ascidians and ascidian larvae. This work improves our knowledge of the anatomy of *T. democratica* and demonstrates the presence of a complex serotonergic system.

Introduction

Thaliaceans are tunicates known for their complex life cycles and their ecological relevance. Nevertheless, the morphology of members of this class is not as well studied as that of other tunicates, such as ascidians, which are among the favourite organisms of developmental biologists.1,2

*Thalia democratica* is a cosmopolitan thaliacean that is found in all oceans, with the exclusion of the Polar Regions. It swims by rhythmic contractions of the muscle bands, which normally project water from the anterior to the posterior opening, providing oxygen and food to the transverse gill bar. Like other thaliaceans, this species has a complex metagenetic life cycle that consists of an alternating succession of sexually produced forms, the oozooids, to blastogenetically produced forms, the blastozooids. The oozooid is a barrel-shaped solitary form, carrying five muscular bands. It generates a ventral stolon, producing a chain of 25–30 small blastozooids. A single blastozooid, endowed with four muscular bands, breaks off from the stolon and swims free. As in other salps, the blastozooids are protogynous hermaphrodites: a single oocyte develops from the simple ovary joined to the atrial wall by a solid rod called the fertilization duct,3 through which only selected sperm reaches the egg.4 The zygote gives rise to the oozooid, which develops very close to the layers of syncytial maternal tissues in order to build a placenta.5 The salps are the only tunicates that lack a real larval stage. After the oozooid is released into the seawater along with the placenta, the testis ripens and sperm cells are released.5

The embryonic development of the central nervous system of *T. democratica* has been described from electron microscope reconstructions.6 During development, the central nervous system passes through a neural tube stage, very similar to that of larvae of other tunicates. Briefly, an early dorsal mass of neurons with an open central canal (tube phase) becomes enriched with a thick mantle of neuroblasts. Afterwards, the neural tube shortens and the central canal disappears because it appears to be filled by the neurites originating from the surrounding neurons (ganglion phase). The nerves coming from the ganglion towards the periphery appear to originate from three paired clusters of cells with large cell bodies (C1, C2 and C3 from the anterior to the posterior). These clusters are located in the equatorial plane of the ganglion and three pairs of nerves directed anteriorly from C1 in addition to two nerves directed laterally and one posteriorly from C2 have been identified. The fibres emerging from the more posterior dorsal cluster C3 are directed posteriorly. Similar to other thaliaceans, the salps have cilia-rich organs that are the main components of the filter-feeding apparatus, such as the transverse gill bar and the endostyle. Bone6,7 accurately described the different regions of the *T. democratica* endostyle: at the bottom, ciliated zones and ciliary fences are located between glandular cells and produce an obliquely forward current; in a more external position, the columnar cilia beat upwards to the top of the endostyle.

In this work, the morphology of the nervous system and ciliary apparatus of *T. democratica* was further investigated by immunolabelling techniques using anti-tubulin and anti-serotonin antibodies to show the three-dimensional pathway of the peripheral nerves emerging from the cell clusters of the ganglion and to compare it with patterns of brain organization in other chordates.

Material and Methods

Animals

Blastozooids and oozoids of *T. democratica* were collected near the coast of Talamone (Italy) in September 2008 with a plankton net (mesh size of 300 μm) during the reproductive period of this species. The samples were sorted under a stereo microscope by means of a glass pipette, rinsed in Millipore-filtered seawater (MFSW), fixed in 4% paraformaldehyde in 0.1 M PBS pH 7.4 for 1 h at room temperature and stored in 70% ethanol at -20 °C.

Immunohistochemistry

After rehydration, fixed samples were rinsed with 0.1 M PBS and were processed for immunolocalization experiments according to the method described by Pennati et al.22 with a few modifications. Whole-mount individuals were washed in PBT (PBS plus 0.25% Triton -X and 0.1% Tween 20) twice for 10 min and incubated for 2 h with 50% heat-inactivated normal goat serum (NGS) in PBT. Then, they were incubated at 4 °C overnight with rabbit anti-s-5-hydroxytryptamine antibody (Medak), diluted 1:400 in PBS/NGS (1:1), with monoclonal anti-β-tubulin antibody (clone 2-28-33; Sigma, Milano, Italy) diluted 1:200 in PBS/NGS.
(1:1) or with monoclonal anti-acetylated α-tubulin antibody (clone 6-11B-1, Sigma). The samples were rinsed several times in PBT (PBS plus 0.2% Tween-20), incubated for 2 h in 1% BSA in PBT and then incubated overnight with a solution containing the secondary antibody diluted at 1:500 in PBT and TRITC-conjugated Phalloidin. The anti-5-HT antibody was detected by Alexa Fluor 488 anti-rabbit IgG secondary antibody, and anti-β-tubulin and anti-acetylated tubulin antibodies were detected by the Alexa Fluor 488 anti-mouse IgG secondary antibody. Specimens were mounted in 1,4-diazabicyclo [2.2.2] octane (DABCO, Sigma) plus MOWIOL (Sigma) on microscope slides. The negative control samples were processed without incubation in primary antibodies. No detectable fluorescence was exhibited by the specimens in the control experiments. The samples were examined using a Leica TCS-NT confocal laser scanning microscope (Leica Microsystems, Heidelberg, Germany) equipped with an argon/krypton (75 mW) multiline laser.

Results

Morphology

Both oozooids and blastozooids of T. democrita were present in the plankton collected. This species is easily recognizable by the violet colour of the tunic and the visceral nucleus, which is characteristic of living animals and disappears after fixation. Barrel-like oozoid specimens presented a ventral stolon with small immature blastozooids (Figure 1A). Actin staining by TRITC-conjugated phalloidin revealed five muscular bands (Figure 1B). Free blastozooids were identifiable by four muscular bands (Figure 1 C,D). They carried embryos at different developmental stages (Figure 1 E,F).

Tubulin immunolocalization

Whole-mount specimens of T. democratica were labelled with an anti-β-tubulin antibody; subsequent observations by confocal laser microscopy showed a strong signal present in the cilia in many structures of the filter feeding apparatus (Figure 2 A,B). In particular, the gill bar, endostyle and pericoronal bands, which are two lateral structures that run anteriorly from the ventral endostyle to the dorsal gill bar, were intensely marked cilia-rich organs (Figure 2 B-E). The anti-acetylated tubulin antibody marked the microtubules present in long projections more specifically, which started from the epidermis and passed through the tunic. These projections were very abundant, regularly distributed along the entire body and were up to 500 μm long.
Acetylated tubulin staining appeared discontinuous, probably indicating that microtubules were in course of stabilization when the samples were fixed. In fact, acetylation is generally considered to occur on stable microtubule assemblies. The cerebral ganglion was intensely marked by the anti-β-tubulin antibody. It is located in the anterior part of the body, dorsal to the pericoronal arches (Figure 2H). This pear-shaped organ is characterized by a narrowing in its anterior portion that divides it into two unequal regions: a small anterior region carrying the pigmented spots and a larger posterior region (Figure 2I,J).

Eight pairs of fibres and several minor ones were identified (Figure 2K). The two most anterior fibres connected the visceral ganglion to the ciliated funnel, which is a cilia-rich organ with an as yet unknown function (Figure 2I,K).

Serotonin (5-HT) immunolocalization

In the whole mount oozooids incubated with an anti-serotonin antibody, a positive signal was present in the cells located in the pericoronal bands, in the proximal tract of the intestine and in the cerebral ganglion (Figure 3A). Serotonin-positive cells were numerous (more than 50) in the pericoronal bands and were regularly positioned to form two almost uninterrupted rows (Figure 3B,C).

In addition, 5-HT immunoreactivity was localized in the first tract of the intestine in several scattered cells with an irregular shape (Figure 3D,E). Serotonin was present in numerous pericarya localized in the posterior region of the cerebral ganglion (Figure 3F-I). Optical sections taken by confocal laser microscopy indicated that serotonin-positive cellular bodies were particularly abundant in the deeper layer of the ganglion. Moreover, a 5-HT signal was present in the placenta of blastozooids that were carrying embryos (Figure 3J-M).

Discussion

An antibody against acetylated α-tubulin was used to stain the ciliary apparatus of oozooids of *T. democratica*. This type of antibody has been widely used in ascidian larvae, mainly to stain the cilia. We observed very long and numerous projections crossing the tunic of the animals all around the body. Although TEM observations are necessary to define the structure of these projections, they were reminiscent of the long cilia of epidermal sensory cells of the trunk and tail of ascidian larvae.

In *Ciona intestinalis* larvae, cell bodies of epidermal sensory neurons are found in pairs...
The presence of serotonin in the ganglion, the ciliated funnel and the intestinal metamorphosis, and serotonin is also known to immunolocalize in the dorsal ganglia during the signalling cascade that triggers thaliacean ed that serotonin is essential in ascidian larvae. In adult thaliaceans and ascidian larvae, the ascidian neural ganglion is similar to that in ascidian larvae. Interestingly, both adult thaliaceans and ascidian larvae have a common pelagic life. T. democratica ooozoids can be considered as directly developing juveniles that lead a planktonic life without the need to choose a substratum on which to adhere and begin metamorphosis. Serotonin is known to modulate circadian rhythms in different organisms. In particular, many planktonic species accomplish vertical and circadian migrations that are directly controlled by light stimuli. Therefore, the presence of serotonin in the central nervous system may be required for orientation and muscle coordination in directional locomotion. It will be of interest to investigate whether or not the contraction of circular muscle bands in thaliaceans is controlled by serotoninergic neurons. In fact, serotoninergic neurons project towards motor neurons and have functions in the coordination and modulation of locomotion in a variety of taxa.

Figure 3. Immunolocalization of serotonin revealed by the FITC-conjugated antibody. A-E and H-I samples were counterstained with TRITC-conjugated phalloidin; A) whole-mount specimen observed by fluorescent microscopy. Serotonin is present in the cerebral ganglion, pericoronal bands and visceral nucleus; B-D) details of the serotonin-positive cells in the pericoronal bands; C,D) Confocal laser microscope (CLM) images; E) magnification of a detail in A showing the arrangement of serotonin-positive cells in the first tract of the digestive apparatus; F-I) serotonin localization in the cerebral ganglion; G) CLM image obtained by the sum of 54 optical sections, step size of 1 µm; F) superimposition of G with a transmission microscope image. H,I) fluorescent microscope images at different focal depths showing the presence of serotonin-positive cells mainly in the core region of the cerebral ganglion; J) CLM image of an early embryo obtained by the sum of 38 optical sections (step size of 2 µm) showing an intense signal in the region adjacent to the placenta; K) superimposition of J with a transmission microscope image. L) CLM image of a late embryo obtained by the sum of 44 optical sections, step size of 2 µm. m, superimposition of L with a transmission microscope image. Serotonin is present in the region contacting the placenta (asterisk) and in the placenta itself (arrowhead). cg, cerebral ganglion; el, cleoblast; es, endostyle; gb, gill bar; m1-m5, muscle bands; pb, pericoronal bands; pl, placenta; sc, serotoninergic cells; vn, visceral nucleus.
as mollusks31,32 and hirudinean annelids.33

Notably, strong 5-HT immunoreactivity was shown in T. democratica by the placental cells. The mature placenta of the pelagic tunicate Salpa fusiformis is formed by two syncytial layers, intimately connected by interdigitating microvilli, that separate maternal and embryonic circulations.14 The placenta shows two distinct regions of serotonin localization in late embryos of T. democratica: one area is close to the embryo whilst the other is closer to the maternal tissue, and these two regions are separated by a central area in which no signal has been found. The presence of serotonin and serotonin receptors has been reported in the human placenta, in which they are thought to play a role not only in placental development and pregnancy maintenance, but also in regulating foetal development.35 It is well known that among diverse physiological functions, serotonin can act as a regulator of cell growth in a variety of cell types, including mammalian placental cells.36 We suggest that 5-HT has a conserved role in the development of chordates.

It is interesting to compare the organization of the cerebral ganglion of the salps with the central nervous system (CNS) of other chordates. The common ground plan of the vertebrate embryonic CNS is characterized by a tripartite organization, consisting of the anterior forebrain, central midbrain and posterior hindbrain expressing the Otx, Pax2/5/8 and Hox genes, respectively. Comparative studies in ascidian larvae showed a homologue tripartite organization. In the larvae of Ciona intestinalis, one of the most studied ascidian species, the anterior part of the CNS, where the main sensory organs are located, consists of a sensory vesicle expressing Otx. This region is considered to be homologous to the forebrain of vertebrates. Posterior to this region, the visceral ganglion expressing Hox and hosting the motor neurons that innervate the tail is considered to be homologous to the vertebrate hindbrain. These two regions are separated by an intervening gap, in which Pax2/5/8 is expressed.17 This is the so-called neck region, which is considered the mid-hindbrain boundary equivalent.38

Based on our results, we developed a hypothetrical structural relationship of the T. democratica ganglion in comparison to that of the ascidian larva. The anterior zone containing the photoreceptor organ is comparable to the sensory vesicle of ascidian larvae in which an ocellus is usually present and, in turn, to the vertebrate forebrain. There is no evidence of a neck region in T. democratica, but the middle zone containing the two motor neuron clusters C1 and C2 could be equivalent to the visceral ganglion of the ascidian larvae in which the motor neurons are located, which, in turn, has the vertebrate hindbrain as a counterpart. This homology is further supported by the presence of serotonergic neurons in this region. In ascidian larvae, the serotonergic neurons are located in the ventral part of the anterior visceral ganglion, as evidenced by the expression of the gene Ci-Tph, the rate limiting synthetic enzyme of serotonin.37 Finding the presence of serotonin in the middle zone of the T. democratica cerebral ganglion is one of the main results of this paper, while it is known that the main serotonergic neurons in the vertebrate CNS are the raphe nuclei of the hindbrain.

Finally, the posterior region around the C3 cluster of the T. democratica cerebral ganglion that contains fibres directed towards the posterior muscular bands, could tentatively be considered equivalent to the ascidian posterior nerve cord. Future gene expression analysis would be required to corroborate these homologies.

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