Commentary: How Does Oestradiol Influence the AVT/IT System in Female Round Gobies During Different Reproductive Phases?
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Teleostean neuropeptides such as arginine vasotocin (AVT) and isotocin (IT) and their homologues arginine vasopressin (AVP) and oxytocin (OT) are implicated in the regulation of reproduction and social behaviour in vertebrates\(^1,2,3\). In fish, those neurohormones have also been reported to influence sexual behaviour, including aggression\(^4,5,6\), courtship\(^7,8,9\), vocalization\(^10\), and spawning reflex\(^11\). In higher vertebrates, AVP/OT systems affect social behaviour by acting within a complex neuronal network defined as a ‘social behavioural neuronal network’ (SBNN)\(^12,13\). The SBNN is composed of groups of neurons, defined as nodes that are reciprocally connected, express gonadal steroid receptors, and constitute an important site of regulation or activation of multiple forms of social behaviour\(^14\). Furthermore, there is evidence that SBNN also exists in non-mammalian vertebrates, including fish\(^15\).

Oestrogens, synthesized in ovaries or locally in the brain, play well-recognized roles in the neuroendocrine control of reproductive physiology, secondary sex characteristics, and sexual behaviour across vertebrate females\(^16,17\). In female \textit{Lythrypnus dalli}, it was presented that the gonadal 17β-oestradiol (E\(_2\)) concentration was approximately 10 times higher than that in the brain\(^18\). In higher vertebrates and fish, there are indications of possible interactions between oestrogens and AVP/AVT- and OT/IT-ergic systems. However, studies were focused on the location of these hormones or their receptors in the brain and oestrogen replacement therapy after ovariectomy. So far, to the authors’ knowledge, nobody has attempted to check if there is a functional relationship between oestradiol and AVT and IT in fish. Our study presents, for the first time, a presumable mechanism for oestradiol action on the AVT/IT system in female round goby (\textit{Neogobius melanostomus}) during the spawning-capable phase and the regressing phase. We aimed to determine whether there is a functional relationship between circulating oestradiol and AVT and IT and to establish which pathway, genomic or non-genomic, is involved in this mechanism in different reproductive phases. For this purpose, the brain explants of female round goby were perfused in medium supplemented with E\(_2\) at doses mimicking the plasma levels of this hormone in nature during different phases of the reproductive cycle. In the perfusion of brain explants, we used E\(_2\) separately or in combination with Fulvestrant (ICI 182.780)—oestrogen receptors (ERs) antagonist or Actinomycin D (Act D)—transcription inhibitor.

How does oestradiol influence AVT and IT secretion and what pathway, genomic or non-genomic, is involved in this regulation?

Changes in oestradiol level and other hormones related to the phases
of the reproductive cycle or social interactions can affect AVP and OT levels by their ERs, which in turn could alter the social behaviour of the individual. Most of the research that has focused on the effect of oestradiol on the synthesis and secretion of nonapeptides has been performed in rodents. Usually, AVP and OT levels or mRNA were reduced following ovariectomy and restored with oestriadiol replacement. In contrast to higher vertebrates, in fish and ovulation at the hypothalamic and ovary level. Our results also suggest that in the regressing phase, the release of both nonapeptides is mediated through classical nuclear ERs via a genomic pathway. It should be mentioned that the mechanism presented here does not exclude the possibility of oestradiol action on the AVT/IT system via non-classical mERs, such as ER-X, GPER, and Gq-mER.

The old new story: from fish to humans

Various species of aquatic fish and invertebrates have long provided valuable models for the study of basic biological processes. Fish are also useful for pathological and toxicological studies because of the high fecundity and relatively brief generation times. Nowadays, the zebrafish (Danio rerio) is a versatile and robust model organism for the study of vertebrate biology, physiology, and human disease, suitable for both developmental and genetic analysis. Therefore, the zebrafish may bridge the gap between its vertebrate and invertebrate counterparts in genetics and developmental studies. Striking homologies have been found between the reproductive system of humans and zebrafish and many similarities from gene functions to reproductive physiology. Hence, fish may be a perfect model for studying the mechanisms whereby hormones modulate the sexual behaviours in both non-human vertebrates and humans. This group of vertebrates demonstrates an exceptional diversity of mating systems and reproductive behaviours. What is more, many fish species present a remarkable plasticity in their biology of reproduction, e.g. sex change, diversity of reproductive tactics that seems to be mediated by the hormone. Furthermore, because the fish neuroendocrine system is well conserved among vertebrates, the mechanisms of hormonal control of behaviour may show similarities with those of other vertebrates.

This kind of study requires a special method that allows monitoring the dynamic hormone secretion and registering even small and short-term fluctuations in their release. It should be noted that only the perfusion culture method allows detailed examination of changes in the release of hormones while ensuring optimal culture conditions. So far, organ perfusion methods have not often been used in fish for lack of suitable techniques. However, an innovative system for organ perfusion proposed by Minuth in the early 1990s gives more options for this kind of technique. In our study, we used the procedure for the gradient perfusion technique (3D) developed by Kalamarz-Kubiak et al., who for the first time applied the MINUCELS and MINUTISSUE tissue engineering technique for perfusion of fish brain tissues. In this study, brain explants were placed on the membrane between rings of tissue carriers inside the gradient container. A specific construction of this container facilitated the uniform supply of medium to the luminal and basal sides to avoid the dead space. It should be noted that although the procedure has been elaborated for studies of AVT and IT in fish explants, after only minor modification, if any, it can serve many other purposes.

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Competing interests

The authors declare no competing or financial interests.

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