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Embryonic suckling and maternal specializations in the live-bearing teleost Zoarces viviparus

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ABSTRACT

The European eelpout follows an aplacental viviparous reproductive strategy, in which gestation lasts 4–5 months. During the last months of development yolk reserves are depleted, and embryos depend on an external source of nutrients. Here we provide evidence for novel specialized physiological, morphological and behavioural adaptations, which we propose as the responsible mechanisms for the exchange of nutrients and gases between the maternal organism and her embryos. Ovarian follicles contain an internal glomerulus-like structure within the distal tip of each follicle. Ultrastructural examination indicated a capacity for steroid synthesis and secretion. Gel electrophoresis demonstrated a protein size distribution in the follicular fluid different from that of the maternal serum, and that ovarian fluid is devoid of protein. From vascular casts and histological sections the follicle was reconstructed. The glomerulus has a central canal that is exteriorized at the tip of the follicle, allowing passage of follicular fluid. Oxygen measurements across the ovary of near-term females showed a strongly hypoxic ovary lumen, yet ovarian fluid adjacent to follicles was oxygen saturated. As another novel observation, embryos were seen engaged in suckling on follicles. We hypothesize that embryos use the follicles on the ovarian wall as placental analogues and that they use their mobile jaw apparatus to attach themselves and apply suction.

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1. Introduction

The course of embryonic development in the European eelpout, Zoarces viviparus (Linnaeus, 1758), has been described in detail (Soin, 1968; Kristoffersson et al., 1973; Korsgaard and Andersen, 1985; Rasmussen et al., 2006). In brief, the female reproductive cycle is initiated by the simultaneous maturation of 25–400 oocytes in the ovarian wall (Soin, 1968; Rasmussen et al., 2006). Each oocyte has its own vascular supply in the form of follicles that grow during maturation and project perpendicularly into the ovary lumen. During ovulation, a single mature oocyte is released by each follicle into the lumen of the ovary, and follicles do not undergo atresia during the subsequent development of embryos.

Gestation in the eelpout lasts 18–22 weeks. Eggs hatch 3 weeks after fertilization and the developing embryos deplete their yolk sac reserves during the following 6 weeks (Rasmussen et al., 2006). For the remaining 9–13 weeks until parturition, embryos have a wet mass gain of more than 1500% (Kristoffersson et al., 1973; Korsgaard and Andersen, 1985), and must depend on an external nutritional source. As extra-vitelline sources must account for some 90% of the wet mass of near-term embryos, several hypothesis have been put forward as to how Z. viviparus provides nutrition for their young. Soin (1968) described the development of a mobile jaw apparatus early in the embryonic development, which for some time was considered a mechanism to facilitate ingestion of ovarian fluid, the uptake of which is facilitated by a hypertrophied hind gut (Kristoffersson et al., 1973). Nutritive substances in the ovarian fluid were thought to originate from a fatty mucous fluid secreted by the ovary wall (Stuhlmann, 1887). However, as no nutritional components have been found in the ovarian fluid (Kristoffersson et al., 1973; Korsgaard, 1983) this scenario appears unlikely.

In contrast, the fluid within the follicle has been shown to be rich in protein, free fatty acid and glucose, with levels identical to that of maternal serum (Korsgaard, 1983). Because the follicle is a closed structure, these nutritional substances are not readily available to developing embryos, and various proposals have been made as to which mechanisms might facilitate transepithelial passage (Skov et al., 2007).

Further to uncertainties pertaining embryonic nutrition, there is the question of oxygen delivery to the embryos. Embryos of Z. viviparus lie freely suspended in the ovarian fluid with no direct maternal connection or specialized structures to facilitate oxygen uptake. Oxygen content in the ovarian fluid of Z. viviparus is low, with a PO2 of less than 5 kPa (Hartvig and Weber, 1984). Despite a high

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2. Materials and methods

2.1. Animals

European eelpout, Z. viviparus were caught locally in set eel traps from October to November. Animals were kept in 4501 aerated holding tanks at the Marine Biological Laboratory, University of Copenhagen, according to Danish institutional guidelines for animal research. Holding tanks were continuously supplied (8 l min⁻¹) with 10 °C recirculating filtered seawater (30 ppt.) and kept under a 12 h light: 12 h dark regime. Fish were fed to apparent satiation twice weekly with chopped mussels (Mytilus edulis). All experiments were performed from mid December to late January.

2.2. Respirometry

Oxygen consumption studies were performed using computerized intermittent flow respirometry (Steffensen, 1989) using a Fibox oxygen meter and dipping probe (Loligo Systems, Denmark). Measurements were made on females during the last month of pregnancy (mean mass ± S.E. 81.9 ± 13.7 g, N = 10) and non-reproductive males (66.9 ± 5.5 g, N = 10) in a 1.448 l shielded respirometer. Animals were allowed to recover from handling overnight in the respirometer prior to measurements. To enable comparison of oxygen consumption in different sized respirometers, the working volume was corrected by the ratio of the volume of typical oxygen meter and dipping probe (Loligo Systems, Denmark) to the respective respirometers. In some viviparous poeciliid species, respiration rates of pregnant females are nearly submerised on a foam rubber mat, and ventilated with fully aerated seawater containing anaesthetic (0.03 g l⁻¹) using a Microx oxygen sensor with a needle-type fibre optic oxygen sensor (Precision Sensing GmbH, Regensburg, Germany). Fish were anaesthetized with Benzocaine (0.06 g l⁻¹), placed on their side, nearly submersed on a foam rubber mat, and ventilated with fully aerated seawater containing anaesthetic (0.03 g l⁻¹). A small lateral incision was made to access the ovary at its widest point, the ovary wall was punctured and the oxygen sensor advanced to the opposing ovary wall. Using a micromanipulator, the oxygen sensor was retracted 1 mm at a time and the partial pressure of oxygen recorded. Distances across the ovary varied from 32 to 38 mm. Measurement points were transformed to percentage of distance across the ovary and binned to the nearest 5% interval. Subsequently oxygen pressures were measured adjacent to the tip of 3 follicles from each individual. Animals were killed by spinal transection, the embryos were removed, counted, and weighed.

2.3. Ovarian oxygen availability

Profiles of the oxygen tension across the ovary lumen were made using a Microx oxygen sensor with a needle-type fibre optic oxygen sensor (Precision Sensing GmbH, Regensburg, Germany). Fish were anaesthetized with Benzocaine (0.06 g l⁻¹), placed on their side, nearly submersed on a foam rubber mat, and ventilated with fully aerated seawater containing anaesthetic (0.03 g l⁻¹). A small lateral incision was made to access the ovary at its widest point, the ovary wall was punctured and the oxygen sensor advanced to the opposing ovary wall. Using a micromanipulator, the oxygen sensor was retracted 1 mm at a time and the partial pressure of oxygen recorded. Distances across the ovary varied from 32 to 38 mm. Measurement points were transformed to percentage of distance across the ovary and binned to the nearest 5% interval. Subsequently oxygen pressures were measured adjacent to the tip of 3 follicles from each individual. Animals were killed by spinal transection, the embryos were removed, counted, and weighed.

2.4. Gel electrophoresis

The size distribution of proteins in female serum, ovarian fluid and follicular fluid from 2 late term pregnant females was analysed by SDS-PAGE using a NuPAGE system (Invitrogen, Denmark). Female serum was obtained from the caudal vein, while ovarian fluid was obtained by Pasteur pipettes after the ovaries were dissected out. Fluid from the follicles was obtained using 10 μl capillary tubes and pooled into 0.2 ml Eppendorf tubes. A 10% BIS–TRIS gel with MES running buffer was used (Invitrogen) with a Mark12 unstained standard (2.5–200 kDa) (Invitrogen). Female serum and follicular fluid were diluted 10-fold, while ovarian samples were diluted 2-fold prior to loading. A 15 μl sample was loaded onto individual lanes and the gel was run at fixed voltage (200 V) for 60 min before being stained in Coomassie Blue R-250 for 45 min and destained overnight in 10% acetic acid / 30% methanol.

2.5. Vascular casting

Vascular casting was performed as described previously (Skov and Bennett, 2004). In brief, animals were euthanized in an overdose of Benzocaine (0.15 g L⁻¹), before the ventral aorta was cannulagrade via the heart. The animal was cleared of blood with 40–60 ml of 1.0% NaCl (pH 7.8) containing Heparin (20 IU ml⁻¹) and sodium nitroprusside (10 μM). A polyurethane resin (vasQtec, Zurich, Switzerland) diluted 4:3 with ethyl methyl ketone was injected into the vascular system and allowed to polymerize overnight before digesting the entire animal in several changes of 20% (w:v) KOH.

2.6. Scanning electron microscopy

Relevant pieces of vascular cast were dissected free and mounted on aluminium stubs using double-sided carbon tape, platinum coated and viewed in a field emission scanning electron microscope (JEOL JSM-6335F, Tokyo, Japan). Tissues for SEM were dehydrated in a graded series of ethanol, and then transferred to 100% hexamethyldisilazane (HMDS) (Braet et al., 1997) in a volume sufficient to cover the specimen. HMDS was allowed to evaporate in the fume-hood, mounted and coated as above.

2.7. Histology

Animals were cannulated as described above and flushed with saline, and perfusion fixed with 2% paraformaldehyde (PFA) in 0.1 M phosphate buffered saline (PBS). Ovaries were dissected out and post-fixed in 4% PFA in 0.1 M PBS at 4 °C overnight. Follicles were dissected out and dehydrated in a graded series of ethanol, infiltrated and embedded in blocks of Technovit 7100 (Heraeus Kulzer GmbH, Wehrheim, Germany) at 4 °C. Blocks were sectioned to distilled water at 3 μm and dried on a hotplate at 60 °C until well bonded, stained with H&E, rinsed in distilled water, air dried and cover slipped using DPX.
2.8. Transmission electron microscopy

Fish were perfusion fixed as described above, using 2.5% glutaraldehyde in 0.1 M PBS. Ovaries were removed and stored in 2.5% glutaraldehyde in 0.1 M PBS at 4 °C until processed for transmission electron microscopy (TEM). Suitable specimen samples were rinsed three times in 0.15 M sodium cacodylate buffer (pH 7.2) and subsequently post-fixed in 1% OsO4 in 0.12 M sodium cacodylate buffer (pH 7.2) for 2 h. The specimens were dehydrated in a graded series of ethanol, transferred to propylene oxide, and embedded in Epon according to standard procedures. Ultrathin sections were cut with a Reichert-Jung Ultracut E microtome and collected on 200-mesh copper grids with Formvar supporting membranes. The sections were stained with uranyl acetate and lead citrate and examined with a Philips CM 100 transmission electron microscope, operated at an accelerating voltage of 80 kV.

3. Results

3.1. Energetic costs of gestation and oxygen availability

The relationship between length and mass of 10 near-term females and 20 non-reproductive males was best described by non-linear regressions (Fig. 1A):

\[
\text{Female } BM = 7.01 \times 10^{-6} \times SL^{2.98} \quad (R^2 = 0.96) \quad (1)
\]

\[
\text{Male } BM = 3.59 \times 10^{-6} \times SL^{2.63} \quad (R^2 = 0.83) \quad (2)
\]

where BM is body mass in grams and SL is standard length in mm. Gestating females were significantly heavier than males of similar length (One way ANOVA, P<0.05). The difference between Eqs. (1) and (2) is the mass increase during gestation (GMI) of a near-term female at a given standard length (SL)

\[
GMI = 2.18 \times 10^{-8} \times SL^{3.80} \quad (3)
\]

Animal mass, intact ovarian mass, and embryonic mass was determined from 4 near-term females. The intact ovary accounted for 28.4±1.5% of the pregnant mass, of which 69.6±1.4% was embryos, approximately 10% was ovarian fluid and the remaining 20% was ovarian tissue.

Metabolic rates increased with body mass for both males and females. Females had a significantly (t-test, P<0.001) higher SMR (31.5±1.6 mg O2 kg\(^{-1}\) h\(^{-1}\), N=10) than males (23.9±0.9 mg O2 kg\(^{-1}\) h\(^{-1}\), N=10). The correlation between oxygen consumption (mg O2 h\(^{-1}\)) and body mass was found to be

\[
\text{Female } MO_2 = 0.0313 \times BM^{1.01} \quad (R^2 = 0.95) \quad (4)
\]

\[
\text{Male } MO_2 = 0.0366 \times BM^{0.92} \quad (R^2 = 0.85) \quad (5)
\]

Transverse ovarian oxygen profiles showed that the centre of the ovary was nearly anoxic (\(PO_2<1\) kPa), while conditions close to the ovary wall were close to normoxic (\(PO_2\) between 8.6–16.8 kPa). Adjacent to the tip of the follicle ovarian fluid was fully oxygenated (\(PO_2 21.87±0.96\) kPa, N=4) (Fig. 1B).

3.2. Follicular anatomy

The gross organization of ovarian blood vessels and capillary organization of the follicles was described previously (Skov et al., 2007) using traditional acrylic casting agents. Acrylic resins cannot be dissected due to their brittle nature. The present study used polyurethane resin which allowed for micro-dissection of vascular casts. In combination with histological approaches, we were able to obtain novel information on the anatomy of the internal structure of the follicles. A semi-schematic representation of the internal follicular anatomy is given in Fig. 2. Follicles vary in length, between 8–12 mm. At the base, the follicle is supplied by a single central blood vessel from the ovary wall. Approximately one-third along the length of the follicle, the supplying vessel divides repeatedly into half a dozen smaller vessels which continue their course for a few millimeters before giving rise to a tuft of convoluted capillaries near the tip of the
follicle (Fig. 3C–D). This vascular structure resembles a glomerulus with some modifications; it is oval in shape and approximately 4–5 mm in length and 2–3 mm at its widest point. From histological sections and SEM images of dried tissue, it is apparent that the glomerulus is enclosed by an epithelial layer (Fig. 3E). The glomerulus is made up by a bi-layer of blood vessels on either side of a layer of connective tissue which is folded onto itself on the longitudinal axis, forming an elongate horseshoe shaped structure (Fig. 3A). The external base of this fold is fused to the inside of the follicular wall (Fig. 2, inset; Fig. 3A) and maintained in place by projections of connective tissue from the outer epithelial layer of the glomerulus to the endothelial lining of the follicle (Fig. 2). The fold created by these vessels forms a central lumen within the glomerulus, termed the glomerular space. Positioned in this manner, the tip of the follicle reaches the opening to the oesophagus (Fig. 6A). From sagittal sections of the head from an individual in which the follicle enters by the mouth, it is apparent that only the outer 3–5 mm are engulfed (Fig. 6B). This corresponds to the region of the follicle containing the glomerulus, and the mouth is closed down on the follicle proximal to this structure. Positioned in this manner, the tip of the follicle reaches the opening to the oesophagus (Fig. 6B).

3.3. Glomerular ultrastructure

The lumen of the blood vessels within the glomerulus ranged between 20–30 µm in diameter (Fig. 4A). The wall is composed of a layer of endothelial cells lined by a continuous layer of pericyte like cells (Fig. 4B,F,G). The endothelial cells are joined by tight junctions (Fig. 4G). In the luminal plasmalemma several invaginations, of which some are coated, could be observed, while the plasmalemma towards the pericytes often showed numerous caveolae (Fig. 4F). The endothelial cells contain mitochondria, Golgi complexes, rough endoplasmic reticulum and dense granula (Fig. 4C,D). Pericytes contained fewer organelles than the endothelial cells, but dense granula could also be observed in these cells (Fig. 4A,B). A thin layer of connective tissue with sparse collagen fibres separates the vessels from a mesangial-like core of the network (Fig. 4C), while the cells in the core are joined by desmosomes (Fig. 4H). The cells contain numerous mitochondria with tubular cristae, rough endoplasmic reticulum and Golgi complexes (Fig. 4G). In some of the cells an abundance of dense granula with a diameter of 3–400 nm could be observed (Fig. 4A,B). Interspersed between the cells smooth muscle cells could be seen (Fig. 4E).

3.4. Gel electrophoresis

The electrophoretic gel showed that the ovarian fluid was practically devoid of any protein with only a few weak bands occurring around the 14.4 kDa marker (Fig. 5). Only 6–7 distinct bands were seen in the adult serum. Distinct protein bands were more numerous in the follicular fluid, several of which corresponded to molecular weights not seen in adult serum.

3.5. Behavioural adaptations

As mentioned in the Introduction, the present study was undertaken following the careful dissection of a single immersion fixed pregnant female which revealed for the first time that eelpout embryos show suckling behaviour, and several individuals were found with the tip of the follicle within their oral cavity. If animals are merely placed in an overdose of anaesthetic, the escape response from the mother will cause all embryos to detach, and subsequent dissection will not reveal any embryos attached to follicles. From several approaches it would appear that a slow and careful approach is best. Pregnant individuals can be isolated in a suitable thermostatted container and allowed to settle down following handling. Once settled, anaesthetic is gradually introduced until the fish lose equilibrium. Once fish lose equilibrium, they should be ventilated, but being careful when handling. Over the following hours the concentration of anaesthetic should be gradually increased, to reach a lethal dose within a few hours. With this approach it was always possible to observe 2–6 embryos (out of a maximum of 36) attached to follicles during dissection.

The follicle entered the oral cavity either via the operculum or directly through the mouth (Fig. 6A). From sagittal sections of the head from an individual in which the follicle enters by the mouth, it is apparent that only the outer 3–5 mm are engulfed (Fig. 6B). This corresponds to the region of the follicle containing the glomerulus, and the mouth is closed down on the follicle proximal to this structure. Positioned in this manner, the tip of the follicle reaches the opening to the oesophagus (Fig. 6B).
4. Discussion

The 32% increase in the resting metabolic rate of near-term females compared to non-reproductive males is in good agreement with previously published accounts on the metabolism of live-bearing fishes. In near-term females of rockfish *Sebastes schlegeli* oxygen consumption in near-term females was 68% higher than in non-reproductive males (Boehlert et al., 1991) and for *S. flavidus* this was 101% higher (Hopkins et al., 1995). Similar increases have been observed for the molly *Poecilia latipinna* (27%) (Timmerman and Chapman, 2003), and surfperches *Rhacochilus vacca* and *Embiotoca lateralis* (53%) (Webb and Brett, 1972). That mass specific metabolic rate scaled linearly with body mass is presumably an indicator that metabolic growth and organogenesis of the embryos are metabolically costly.

The metabolic oxygen requirement by a gestating female can be summarized as the sum of non-reproductive oxygen consumption, embryo oxygen consumption and oxygen requirement to maintain the litter (Timmerman and Chapman, 2003). Thus, for *Z. viviparus*, the metabolic cost associated with gestation can be approximated by subtracting the SMR of a non-reproductive individual; a 100 g near-term female with a SMR of 3.28 mg O₂ h⁻¹ (Eq. (4)), would weigh 71.6 g if she were non-reproductive and therefore have a SMR of 1.86 mg O₂ h⁻¹ (Eq. (5)). Therefore, the metabolic cost of litter maintenance and oxygen consumption by embryos, accounts for almost half (43%) of the SMR of a near-term female.

The present results showed that the ovarian environment is far more hypoxic than previously assumed. Hartvig and Weber (1984) reported oxygen tensions in the ovary of *Z. viviparus* to be ~5 kPa, but in the 4 individuals we examined, P<sub>O₂</sub> in the ovarian lumen rarely exceeded 0.5 kPa. A P<sub>O₂</sub> of 5 kPa was only observed in the peripheral 3–4 mm, although immediately adjacent to the follicles, the ovarian fluid was fully oxygenated. Korsgaard (1994) reported a pH of 7.5 in the ovarian fluid, which is likely to be somewhat lower than maternal plasma, and this may provide sufficient Root effect to unload oxygen from the maternal blood during passage of the follicular vascular network. The oxygen profile of the ovary would suggest that embryos must remain in close proximity to the follicles to obtain sufficient oxygen. Even given the higher oxygen affinity of embryonic blood (Hartvig and Weber, 1984) embryonic haemoglobin could achieve no more than 20% saturation in the ovary lumen. It might be argued that the low oxygen pressures measured in the current study were an...
artefact caused by reduced ovarian blood perfusion resulting from the anaesthesia. In our opinion, the high oxygen tensions measured adjacent to the follicles, including those located opposite to the entry site of the oxygen sensor, do not suggest this to be the case. Maintaining a juxtaposition of the follicle and the gills of the embryos also constitutes an energetic advantage in that the maternal organism does not have to expend large amounts of energy in maintaining high oxygen levels throughout the ovary. Embryos have a thick avascular epidermis making it unlikely that they lose oxygen to their environment across the body surface.

Viviparity has evolved on 132 independent occasions in the lower vertebrates; the vast majority of these being placental reptiles (72%) or cartilaginous fishes (8%) (Blackburn, 1992). Aplacental matro-trophic viviparity has evolved on 12 occasions in teleost fishes (Blackburn, 1992). To date, nutrients have been considered derived from the maternal circulation in all species, except the European eelpout. The present work shows that embryos of the eelpout obtain nutrients by suckling on ovarian follicles, and that these nutrients are also derived from the maternal circulation. The profiles from the gel electrophoresis support the suggestion that nutrients are passed directly from the follicle to the embryos without intermediary mixing in the ovarian fluid. The role of the mobile jaw apparatus is most likely to facilitate swallowing of fluid (Soin, 1968). However, we also believe it constitutes an anatomical requirement for suckling, both for the purpose of attachment to the follicle and for obtaining follicular fluid. The developmental stage at which the initial attachment of embryos

Fig. 4. Transmission electron micrographs of the elaborate network of vessels of the glomerulus-like structure within the outermost 4–5 mm of the follicle. (A) The lumen (L) of the vessels is 20–30 μm in diameter. (B) The vessel wall is comprised of a layer of endothelial cells (E) surrounded by a layer of pericytes (P). A thin layer of connective tissue with sparse collagen fibres (C) separate the pericytes from the cells within the core of the glomerulus network which could be heavily loaded with dense granules (arrows). (C) Dense granules (arrows) were observed in the core, as well as mitochondria (M). Occasionally smooth muscle (SM) cells could be observed. (D) In some regions of the core mitochondria (M) were densely packed with the appearance of tubular cristae. (E) Smooth muscle cells could be found interspersed between dense granules. (F) Invaginations in the plasmalemma were found both on the luminal side of the endothelial cells as well as towards the pericyte layer (arrows). (G) Endothelial cells also contained Golgi complexes, rough endoplasmic reticulum and dense granules (arrows). (H) Cells within the network core were connected by tight junctions with prominent desmosomes (D). Scale bars: A, 10 μm; B, 2 μm; C–E, 5 μm; D, F, H, 500 nm; G, 1 μm.
Suckling behaviour has never previously been observed in *Z. viviparus*. The reason for this is probably that the attachment between embryo and follicle is not static. Embryos are easily disturbed when handling the female and detach from the follicles. Even with the utmost care, the dissection of fresh specimens has never allowed observations of embryos engaged in suckling. Admittedly it is difficult to ensure that embryos remain attached to the follicles during fixation, and when successful, it is only a few that do. There is nothing to suggest that embryos are permanently attached to the follicles; most probably they are able to detach and relocate at will. Following handling, pregnant females can subsequently be brought to spawn, implying that embryos are able to locate a vacant follicle following disturbances, yet the homing mechanisms responsible are unknown. Suckling is an efficient strategy for several reasons; one being that embryos receive concentrated nutrients rather than having to forage within the ovary. The presence of one follicle per embryo presumably also ensures an equal distribution of nutrients, and would account for the small variation in individual size at term (Rasmussen et al., 2006).

We have demonstrated a strong dependency between the eelpout embryo and the ovarian follicles, in which the latter serves a functional analogy to a combined placenta and teat during embryonic development. This bears some resemblance to the ovarian leaf-shaped projections in *Jenynsia lineata* described by Turner (1940). The morphological and functional specializations of the follicles lend evidence to a very complex matrotrophic relationship in *Z. viviparus*, with numerous physiological and behavioural adaptations.
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