Broodstock and Breeding

*Dicentrarchus labrax* and *Sparus aurata* egg diameters ~ 1mm
Broodstock management

- Broodstock collection
- Breeding programs
- Spawning control
  - Natural - Photoperiod/temperature control
  - Induced - hormones
- Broodstock nutrition
- Water quality and environment
- Hygiene aspects
- Disease aspects
- Egg quality
Broodstock management

Prophylaxis

- Fresh water / Formalin baths, Oxytetracycline

Nutrition 2% body weight per day

Substrate / Monitoring

- Aquamat®, Underwater Video, Artificial Cleaning Station

Long-term parasite control
Breeding Strategies

• Maximize survival of eggs and fry
  – natural habitat

• Ultimate Factors
  – food supply
  – absence of predators
  – water quality
Breeding Strategies

• Proximate Factors
  – environmental cues
    • Temperature
    • Day length
    • Rainfall
    • Salinity
    • Lunar cycle
  – anticipate future spawning
3 strategies for production of eggs and spawning

• Synchronous spawning
  – One crop of eggs and sperm are produced and spawned at one time after which the fish dies e.g. Salmon

• Group synchronous
  – Groups of eggs are produced and spawned at one time but several cycles of development of spawning can occur (annual or other interval e.g. common aquaculture species)

• Asynchronous spawning
  – Continual development and spawning of oocytes
Reproductive biology
- freshwater fish
Reproduction stages

- Proliferation of oogonial cells in the ovary of the female fish that will develop into oocytes (eggs) & initiation of meiosis (reduction of chromosome numbers in half)

- **Stage I** – Development of basic cellular structures
- **Stage II** – Vitellogenesis – synthesis and uptake of egg yolk proteins which provide nutrients for the developing embryo
- **Stage III** – Maturation caused by the steroid hormone progesterone
- **Stage IV** – Spawning - hydration and fertilization
- **Stage V** - Recovery
Stage I – Development of basic cellular structures

• Enlargement of the nucleus, appearance of multiple nucleoli and subcellular organelles including cortical aveoli (important in fertilization events)
• Endogenous vitellogenesis – protein synthesis
• Two cell layers (theca & granulosa cells) develop and surround the oocyte to form a follicle
• Theca and granulosa cells cells are responsible for production of reproductive steroid hormones
Stage I: Oocyte Development

- **Oogenesis**
  - Oogonia give rise to oocytes
  - Oogonia undergo meiosis
  - Surrounded by epithelial cells (follicle)
  - Meiosis stops and enters a long state of cytoplasmic growth

- **Gonadotropin independent and body size dependent**
Stage I: Yolk Vesicle Formation

• Response to Proximate Factors

• Surges in gonadotropin
  – Previtellogenic oocyte
  – Appearance of yolk vesicles
  – “endogenous vitellogenesis”
  – GtH triggers yolk vesicle formation
Stage II – Vitellogenesis

• Interaction of the pituitary gland in the brain, the follicle cells, the liver and the eggs to produce hormones (gonadotrophins)
• Gonadotrophin stimulates the theca and granulosa cells to produce estrogen which in turn stimulates the liver to produce vitellogenin the precursor to egg yolk proteins
• Vitellogenesis is the longest stage of oocyte development and requires nutrient input either directly from feed or body stores of muscle and fat
• If insufficient amounts of egg proteins are deposited in the oocytes then fry development form these eggs will not complete development and have high mortalities as eggs or sac-fry
Stage II: Vitellogenesis

• Vitellogenin (Vg)
  – Phospholipid produced in liver
  – Transported to oocyte via blood
  – Precursor to egg yolk protein

• High nutritional demand
• Longest phase
Stage II: Vitellogenesis

- Egg yolk production
  - Theca cells
    - Testosterone (T)
    - Induced by GtH
  - Granulosa cells
    - Estradiol (E$_2$)
    - E$_2$ stimulates liver to produce Vg

- End of Vitellogenesis
  - GtH II produced in higher levels than GtH I
  - Follicle produces maturation inducing substance (MIS)
Stage III – Maturation

• Final stage of oocyte development and lasts 24-72 hrs
• Nucleus of the egg migrates from the centre of the egg to the periphery and meiosis resumes again to pause before completion
• The membrane surrounding the nucleus disappears – geminal vesicle breakdown
• Uptake of water occurs in some species at this point
• When maturation is complete the oocytes are ovulated from the follicle due to influence of protaglandin’s
• Depending on the species
  – Some oocytes retained in the ovary
  – Eggs released into the peritoneal cavity until spawning
Stage III: Oocyte Maturation

- Maturation occurs in less than 24 hrs
  - Germinal vesicle migration
  - Germinal vesicle breakdown
  - Resumption of meiosis
  - Water uptake
Stage IV – Spawning - hydration and fertilization

- Occurs shortly after maturation of the eggs or the eggs will become overripe
- Spawning behaviour depends on the species
- Spawning can be induced by injection of gonadotrophin hormones (stage III) – carp pituitary extract or human chorionic gonadotrophin
- At fertilization activation of the oocytes involves the release of the contents of the cortical alveoli into the space between the egg membrane and the chorion which forms the egg shell. There is a hole in the chorion called the micropyle which allows the sperm to pass through and fertilize the egg.
- On activation and the release of the cortical alveoli the chorion lifts off the egg and the micropyle begins to close
Stage IV: Spawning

• Optimum environment
  – Mating behaviors

• Ovulation controlled by F-prostaglandin (PG)
  – PG release stimulated by MIS
  – GtH II is required for this process

• Common stage for hormone therapy
Stage V Recovery

- Restore body condition
- New oocyte development
Overview of the stripping process

Cover the head to prevent stress
Towel to have good grip
Anesthesia → Draining off water → Stripping of ova

Sperm collected in immobilizing solution (IS)
(170-200 mM NaCl, 30 mM TRIS, pH 7)

Possibility of sperm storage up to 72 hours at ±2°C

100 g of eggs, 2 ml of sperm with IS and 50 ml of AS

Egg storage, max 4 hours at 15-18°C

Activation solution (AS)
(17 mM NaCl 5 mM TRIS, pH 7)

Stirring solution with eggs (~ 10 s)
Standing 1-2 min

Slowly stirring

Adding 25 ml of AS

Elimination of stickiness:
3 min after activation by adding diluted enzyme alcalase
* (100 ml per 100 g eggs)

Stirring enzyme with eggs to 2 min

Eggs are fast rinsed with water and transferred into Wels jars (7-10 l)
Spawning

- Spawning behaviour
- Egg release
- Pheramone release
- Release of milt
  - Fertilization

Methods

- Simulate natural environment
- Dry stripping
- Hormone induction
Spawning

- Simulate natural environment
  - Nesting basins
  - Floating plants
  - Floating nests
  - Fiber mats
Spawning

• Dry Stripping
  – Strip eggs and milt
  – Mix together
  – Add water
    • Activate eggs and milt
closure of micropyle
  – Separation of sticky eggs
    eg Silting

MS222 is toxic to eggs and milt!
Spawning

- Induced Spawning
  - Stage 3 Oocyte maturation
  - Egg Sampling
    - Egg size
    - Yolk clarity
    - Oil globules

Ripe eggs at ovulation
Spawning

- Oral
- Hormone Injection
  - Intra muscular
  - Intra peritoneal
    - Carp Pituitary Extract
    - Human Chorionic Gonadotropin (HCG)
    - GnRHa
- Single or multiple injection
  - Stage 3 complete in 12 - 48 h
- Slow release capsules
- Natural Environment or Dry Stripping
Induced spawning

• Induced spawning
• Natural spawning
Regulation of spawning

From Broodstock management and selective breeding Yonathan Zohar
Methods of inducing spawning

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Development of antibodies

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Use of analogues

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Injecting implants or microspheres

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Induced sperm production using GnRH implants

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Commerically available

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