Stocking density effect on survival and growth of early life stages of maraena whitefish, Coregonus maraena (Actinopterygii: Salmoniformes: Salmonidae)

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Abstract

The maraena whitefish, Coregonus maraena (Bloch, 1779), is often considered a suitable candidate for intensive aquaculture diversification in the EU. However, only a few such farms in Europe are in operation. Rearing this species in recirculating aquaculture systems is a recent innovation, and optimisation is necessary to standardise aspects of larviculture. This 30-day study investigated the effect of stocking densities of 25/L, 50/L, 100/L, and 200/L on the survival and growth of maraena whitefish larvae in a recirculating aquaculture system. The four groups of larvae (initial weight = 7.4 ± 0.1 mg; initial total length = 13.0 ± 0.1 mm) in three repetitions were reared in a recirculating system. Larvae were fed fresh live brine shrimp metanauplii every 3 h at a rate converted to larval stocking density. After the experiment, 10 larvae from each tank (30 of each density group) were weighed on a digital microbalance (ABJ 220-4M KERN, Germany, readout = 0.1 mg) and measured manually on images taken with Leica MZ16 A stereomicroscope and a digital colour camera with 5-megapixel resolution for Leica DFC420 Image Analysis. No significant differences in final body weight, total length, size heterogeneity, condition factor, or survival were found among treatments (P > 0.05). The highest non-significant survival rate and growth parameters were observed in larvae reared at 25/L. On the contrary, it is possible to rear maraena whitefish larvae at high stocking density without any subsequent negative consequences for growth and survival. As no significant differences in any evaluated parameter were observed between groups of larvae at the highest and lowest stocking density, we conclude that it is possible to rear maraena whitefish larvae at high stocking density (and 200/L) without any subsequent negative consequences for growth and survival.

Keywords

coregonids, fry, growth metrics, larviculture, recirculation systems

Introduction

The maraena whitefish, Coregonus maraena (Bloch, 1779), is a promising species for inland freshwater aquaculture throughout east-central Europe (Mukhachev and Gunin 1999), and northern Europe, especially Finland (Jobling et al. 2010) and Norway (Siikavuopio et al. 2011). Several decades ago, due to predation by the great cormorant Phalacrocorax carbo, the population dramatically declined (Suter 1997). Eutrophication has also contributed to the decrease
Materials and methods

Eggs and larvae

Maraena whitefish were obtained from the Szczecin Lagoon (the River Odra estuary), north-western Poland. The broodstock comprised 120 fish at a 1:1 sex ratio. Gametes of three-year-old 60 females (mean weight, 800.4 ± 80.1 g, mean ± SEM; mean total length, 30.2 ± 1.1 cm) and three-year-old 60 males (650.5 ± 49.7 g, mean ± SEM; mean total length, 26.4 ± 0.9 cm) were stripped manually (no hormone stimulation) by commercial fishermen in December 2016 shortly after fish capture and transported to local hatcheries for fertilization and incubation. Eggs (100 mg) were fertilized with 0.5 mL of milt mixed with 50 mL of hatchery water and incubated at the ambient water temperature of the river (2–3°C) with initial water inflow 3 L/min, oxygen saturation to 90%, and pH near 7.0. In February 2017, the eggs were taken to the Department of Lake and River Fisheries (Olsztyn, Poland) where they were distributed among five 8-L Zug jars (n = ~150 000 eggs/jar) in a recirculating system and incubated at 3.0–3.5°C with water inflow 3 L/min, oxygen saturation to 90%, and pH near 7.0. In total, ~750 000 eggs were incubated. After 60 days, eggs were transferred to the second set of 8-L Zug jars and incubated at 8–9°C to accelerate development and hatching. After 5 days, the temperature was increased to 10°C for mass hatching. Hatching success was estimated at 90%, and about 675 000 larvae were available for the experiment. Hatched larvae swam across to a tank (total volume 1 m³) underlain with 0.2 mm mesh. After 24 h, larvae were transferred to tanks in the RAS.

Experimental system

Four groups of larvae in three replicates were transferred to the experimental aqua system consisting of twelve 2 L aquaria, 96 × 154 × 200 mm. The recirculating system (2300-L total water volume) included a series of filtration sections (total biofilter volume 1500-L), a settling tank (500-L water volume). Thirty fish were weighed and measured to obtain the initial values for weight and length. Maraeana whitefish larvae (initial weight, 7.4 ± 0.1 mg, mean ± SEM; initial total length, 13.0 ± 0.1 mm) were placed into each aquarium at stocking density of 25/L (S25), 50/L (S50), 100/L (S100), and 200/L (S200). A biomass by litre (g/L) was 0.185 (S25), 0.370 (S50), 0.740 (S100), 1.480 (S200). A total of 2250 larvae were used in the experiment.

Rearing conditions

The oxygen level, water temperature, and pH were checked daily at 0800 and 1600 h. The pH range was monitored using an OxyGuard H04PP Handy pH meter (OxyGuard International, Denmark). The initial tempera-
ture without supplemental heat was 10°C. Water temperature ~19°C was regulated by a HAILEA HC-1000A cooler (China). The temperature was gradually elevated from 10°C to 19°C (3°C/day). Oxygenation was maintained using two SICCE Syncra 5.0 pumps (5000 L/h) (Italy). Ammonia, nitrate, and nitrite concentrations were analysed using HACH, LCK 304, LCK 339, LCK 341 (Germany) with a HACH DR5000 spectrophotometer (Germany). Disinfection used a 30 W UV MCT Transformeren GmbH steriliser (Germany). NaCl was added at 1 g/L weekly to maintain a 16:1 chloride:nitrogen ratio. A constant inflow of 0.4 L/min was ensured. Dead larvae were removed and counted during daily cleaning. The level of organic matter remained low. A low CO₂ level was maintained via aeration and keeping alkalinity stable. During the 30-day trial, basic physico-chemical parameters were following: temperature = 19.1 ± 0.0°C, pH = 8.7 ± 0.0, O₂ saturation = 85.8 ± 0.9%, O₂ concentration = 7.9 ± 0.1 mg/L, NH₄ + = 0.1 ± 0.0 mg/L, NO₂⁻ = 0.8 ± 0.1 mg/L, NO₃⁻ = 21.2 ± 5.4 mg/L.

Feeding

Larvae were fed fresh live metanauplii of brine shrimp, Artemia salina (Ocean nutrition, HE > 230 000 NPG, Belgium) (20–24 h old, 0.4–0.5 mm) four times daily at 3 h intervals during the light phase (0830 to 1730 h). The feeding level was fixed to the range of 500–700 mg/L, NO₃ = 7.9 ± 0.1 mg/L, NH₄ + = 8.7 ± 0.0, O₂ saturation = 85.8 ± 0.9%, O₂ concentration = 7.9 ± 0.1 mg/L, NH₄ + = 0.1 ± 0.0 mg/L, NO₂⁻ = 0.8 ± 0.1 mg/L, NO₃⁻ = 21.2 ± 5.4 mg/L.

Sampling and measurements

After the experiment, 10 larvae from each tank (30 of each density group) were weighed on a digital microbalance (ABJ 220-4M KERN, Germany, readout = 0.1 mg) and measured manually from images taken with Leica MZ16 A stereomicroscope and a digital colour camera with 5-megapixel resolution for Leica DFC420 Image Analysis.

A sample size of ten larvae per tank, 30 larvae per treatment, was used as in a number of studies (Kaiser et al. 2003, Mahmood et al. 2004, Fletcher et al. 2007, Celda et al. 2008, Nowosad et al. 2013, Palińska-Zarska et al. 2014, Laczynska et al. 2016).

The survival rate (SR), size heterogeneity (SH), and condition factor (K) and specific growth rate (SGR) were assessed as follows:

\[ SR(\%) = 100 \times \left( \frac{N_f}{N_i} \right) \]

in which \( N_i \) and \( N_f \) = initial and final number of larvae, respectively;

\[ SH(\%) = 100 \times \left( \frac{SD}{W_m} \right) \]

in which \( SH = \) size heterogeneity; \( SD = \) mean standard deviation of weight of 10 randomly selected larvae/tank; \( W_m = \) mean weight [mg] of 10 larvae/tank.

\[ K = 100 000 \times W \times (TL^3)^{-1} \]

in which \( W = \) mean weight [g] of 10 larvae/tank; \( TL = \) mean total length [mm] of 10 larvae/tank

\[ SGR(\%) = 100 \times \left[ \frac{\ln W_f - \ln W_i}{d} \right] \]

in which \( W_f \) and \( W_i \) are final and initial weight of larvae, respectively [g]; \( d = \) duration of the experiment [days].

Statistical analysis

Statistical analyses were performed using STATISTICA 12.0 (StatSoft, Praha, Czech Republic). Data are presented as mean ± SEM. The effects of stocking density on \( W, TL, SR, K, SH, \) and SGR were analysed by one-way ANOVA with stocking density as a fixed variable. Differences were considered significant at \( P < 0.05 \). Prior to ANOVA, SR, K, SH, and SGR were arcsine-transformed. All data were tested for homogeneity of variance using the Cochran, Hartley, and Bartlett test, and for normality with the Shapiro–Wilk normality test. The parametric Tukey test was used for assessing differences among groups in \( W, TL, SR, SH, K, \) and SGR (Table 2).

Results

At the conclusion of the trial, no significant \( (P>0.05) \) differences among treatments were observed in SR, \( W, TL, SH, K, \) or SGR (Table 2). The highest SR (92.7% ± 2.4%),
Table 2. One-way ANOVA results for the factor stocking density on total length (TL), body weight (W), size heterogeneity (SH), condition factor (K), survival rate (SR), and specific growth rate (SGR) of larvae of maraena whitefish, Coregonus maraena (Bloch, 1779).

| Parameter | Source of variation | SS  | DF  | F   | MS  | P  |
|-----------|---------------------|-----|-----|-----|-----|----|
| TL        | SD                  | 0.9 | 3.0 | 0.3 | 2.3 | 0.2|
| W         | SD                  | 466.3 | 3.0 | 155.4 | 2.7 | 0.1|
| SH        | SD                  | 22.7 | 3.0 | 7.6  | 0.2 | 0.9|
| K         | SD                  | 0.0  | 3.0 | 0.0  | 2.2 | 0.2|
| SR        | SD                  | 3.6  | 3.0 | 1.2  | 0.2 | 0.9|
| SGR       | SD                  | 0.0001 | 3.0 | 2.1  | 0.00005 | 0.2|

SD = stocking density; SS = sum of square; DF = degrees of freedom; F = distribution fitting; MS = mean square; P = probability.

Table 3. Effect of stocking density on growth and survival of larvae of maraena whitefish, Coregonus maraena (Bloch, 1779), in a 30-day growing trial.

| Group   | SR [%] | TL [mm] | W [mg] | SH [%] | K     | SGR [%] |
|---------|--------|---------|--------|--------|-------|---------|
| S25     | 92.7 ± 2.4 | 30.7 ± 0.3 | 147.9 ± 5.8 | 22.5 ± 4.3 | 0.51 ± 0.01 | 0.50 ± 0.003 |
| S50     | 91.3 ± 1.5 | 30.4 ± 0.2 | 135.7 ± 1.6 | 20.3 ± 3.6 | 0.48 ± 0.01 | 0.49 ± 0.001 |
| S100    | 91.3 ± 1.1 | 30.4 ± 0.1 | 135.1 ± 3.5 | 21.1 ± 4.9 | 0.48 ± 0.00 | 0.49 ± 0.003 |
| S200    | 91.8 ± 1.0 | 30.0 ± 0.2 | 131.3 ± 5.2 | 18.7 ± 2.3 | 0.49 ± 0.01 | 0.49 ± 0.004 |

Groups represent stocking densities of 25, 50, 100, and 200 larvae/L, respectively; SR = survival rate, TL = total length, W = body weight, SH = size heterogeneity, K = condition factor, SGR = specific growth rate.

**W** (147.9 ± 6.3 mg), TL (30.7 ± 0.4 mm), SH (22.5% ± 1.1%), K (0.51 ± 0.01), and SGR (0.50 ± 0.003%) was observed at S25 (Table 3).

**Discussion**

The fact that growth–weight parameters did not differ significantly means that maraena whitefish growth was not influenced by stocking density at the tested levels. Slightly lower (non-significant) growth was found with increasing stocking density. It is important to sustain uniformity of fish size in aquaculture (Biswas et al. 2010). The effect of stocking density on larva size heterogeneity may be species-dependant. For instance, the relation of stocking density to size heterogeneity has been reported to be positive in red tilapia Oreochromis niloticus (Linnaeus, 1758) × Oreochromis mossambicus (Peters, 1852), when stocking density was 0.1, 0.2, 0.4, 0.8, 1.6, and 3.2 fry per litre (Huang and Chiu 1997), but negative in Arctic charr, Salvelinus alpinus (Linnaeus, 1758), with stocking density 10, 20, 28, 40, 60, 80, and 100 fry per litre (Wallace et al. 1988). We found size variation with respect to stocking density at the levels tested to be negligible with the only non-significant more uniform size in the S200 group and the least uniform in the S25 group. North et al. (2006) observed the same trend, with the highest size heterogeneity observed in rainbow trout, Oncorhynchus mykiss (Walbaum, 1792), reared in low stocking density and vice versa.

Stocking density can influence mortality rate, with survival often negatively correlated with stocking density as shown for silver perch, Bidyanus bidyanus (Mitchell, 1838) (see Rowland et al. 2006). Fish species can be classified as density-independent or density-dependent. Tilapia larvae (Huang and Chiu 1997) were reported to be density-dependent. Survival was high and not significantly affected by stocking density in the presently reported study, thus maraena whitefish seem to be density-independent, and stocking density is not likely a limiting factor in their survival in intensive rearing. High survival in all groups indicates that high-density aquaculture may be suitable for the production of this species. This phenomenon was also seen in Kupren et al. (2011) for asp, Leuciscus aspius (Linnaeus, 1758); ide, Leuciscus idus (Linnaeus, 1758); and chub, Squalius cephalus (Linnaeus, 1758).

Stocking density has been reported to be an important factor in fish growth (Saoud et al. 2008) and is of particular concern in the welfare of intensively farmed fish (Ashley 2007, Wocher et al. 2011). Mortality (Ellis et al. 2012), as well as susceptibility to pathogen infections and fin damage (Turnbull et al. 1998, Jones et al. 2011), in farmed fish, are generally considered important indicators of welfare. Ashley (2007) suggests that unsuitable stocking density can result in damage or death of fish. Negative effects of high stocking density on fish growth and survival can be attributed to impaired water quality associated with accumulation of fish metabolites and carbon dioxide, with accompanying decline in pH level (Hosfeld et al. 2009). As no technical problems or disease occurred during the course of our study, we can conclude that water quality and stocking density effects were accurately evaluated. The high survival rate at all density levels and lack of observable damage to fins are evidence of appropriate rearing conditions with respect to fish welfare.

**Conclusions**

No significant differences in any evaluated parameter were observed between groups of larvae at the highest and lowest stocking density. It is possible to rear maraena whitefish larvae at high stocking density with no subsequent negative consequences for growth and survival. This study examined fry and early-stage larvae, but a further study, focusing on juvenile and adult maraena whitefish, is warranted. The effects of stocking density on stress hormone response, body composition, and haematological and biochemical parameters of maraena whitefish should be studied.

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