



PLASMA SEX STEROID PROFILES IN MEAGRE (*ARGYROSOMUS REGIUS*)

Vallés, R.¹, Bayarri, M.J.², Mañanós, E.² and Duncan, N.¹

¹ Institut de Recerca i Tecnologia Agroalimentàries (IRTA), Sant Carles de la Ràpita, Tarragona, Spain. Fax: +34-977744138. e-mail: neil.duncan@irta.cat

² Instituto de Acuicultura de Torre la Sal (CSIC), Torre la Sal, Castellón, Spain

Introduction:

Meagre is a new species for Mediterranean aquaculture and the reproduction is still relatively unknown. The aim of this work is to obtain the profile of sex steroids levels during one year and 8 months to gain knowledge on the meagre reproductive system.

Methods:

A meagre broodstock (first generation, G1, in captivity) of mean weight $6,79 \pm 0,19$ kg was held in IRTA, Sant Carles de la Ràpita, Spain, under natural photoperiod and temperature (min 12.8°C, max 23.7°C). Every month, from February 2009 to November 2010, fish were anesthetized with MS-222 (70 mg L^{-1}) and a blood sample of 1.5 mL collected. Plasma was obtained by centrifugation (3000g, 15min, 4°C) and stored at -80°C for steroid analysis. Levels of estradiol (E_2), testosterone (T) and 11-ketotestosterone (11-KT) were quantified by ELISA. Oocyte samples were obtained by canulation biopsy and sperm samples by abdominal pressure.

Results:

During the first year, female plasma levels of E_2 were elevated during February and increased reaching

the maximum in April ($0.54 \pm 0.09 \text{ ng ml}^{-1}$), which coincided with the presence of vitellogenic oocytes in the ovary. The E_2 plasma levels then returned to basal levels in June. In the second year, the E_2 started to increase in December and the highest levels were observed in March ($1.15 \pm 0.52 \text{ ng ml}^{-1}$). A similar pattern was observed in plasma levels of 11-KT in males, during the first year levels were significantly higher in February ($0.64 \pm 0.07 \text{ ng ml}^{-1}$) and decreased to a low in June. The highest percentage of spermiating males was encountered in April. In the second year, 11-KT began to increase in November with the maximum in March ($0.27 \pm 0.11 \text{ ng ml}^{-1}$) and decreased in May when 100% of males had flowing sperm. Plasma levels of T in males did not show significant differences over the sampling period.

Conclusion:

In conclusion, this study provides the profiles of three sex steroids that clearly identify the period of gametogenesis (December – March) before the spawning period (April-May).