

# Oral treatments for monogenean parasites of farmed yellowtails, *Seriola* spp. (Carangidae)

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---Title page images from L – R: *Seriola quinqueradiata* (Carangidae) sea-cage, Kyushu, Japan; *Benedenia seriolae* (Capsalidae) on the eye of a *Seriola lalandi* (Carangidae); *Heteraxine heterocerca* (Heteraxinidae). Images: R. E. Williams. This work contains no material which has been accepted for the award of any other degree or diploma in any university or other tertiary institution to R. E. Williams and, to the best of my knowledge and belief, contains no material previously published or written by another person, except where due reference has been made in the text.

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Rissa Williams 30 November 2009 This page has been left intentionally blank.

#### DEDICATION

To my parents, Daisy and Terry Williams

You taught me how to watch, listen and learn. You gave me the freedom to grow and be independent and a loving home to come back to. Thank you for believing in me.

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### PUBLICATIONS ARISING FROM THIS PHD

**Williams, R.E.**, Ernst, I., Chambers, C.B., Whittington, I.D., 2007. Efficacy of orally administered praziquantel against *Zeuxapta seriolae* and *Benedenia seriolae* (Monogenea) in yellowtail kingfish *Seriola lalandi*. Diseases of Aquatic Organisms 77, 199-205. doi: 10.3354/dao01824

#### ABSTRACT

Japanese yellowtail *Seriola quinqueradiata* has been commercially farmed in Japan since the 1940s. In comparison, sea-cage farming of yellowtail kingfish *Seriola lalandi* in Australia is still developing, with commercial production commencing in 1998. In Australia, *S. lalandi* is parasitised by *Zeuxapta seriolae* and *Benedenia seriolae*. In Japan, *S. quinqueradiata* is parasitised by *Heteraxine heterocerca* and *B. seriolae*. These monogeneans affect industries in both countries and management of these parasites is required to prevent impacts on fish health and commercial losses.

I investigated efficacy (% reduction of mean parasite abundance) for orally administered praziquantel, fenbendzole and oxfendazole against *Z. seriolae* and *B. seriolae* on *S. lalandi* and the efficacy of orally administered praziquantel and febantel against *H. heterocerca* and *B. seriolae* on *S. quinqueradiata*. Medications were administered to fish by surface coating feed pellets or via direct intubation of the stomach. *Seriola lalandi* administered fenbendazole and oxfendazole by surface coating of feed had lower abundance of the gill parasite *Z. seriolae*. *Seriola quinqueradiata* intubated with febantel had lower abundance of the gill parasite *H. heterocerca*. Neither fenbendazole nor oxfendazole administered to *S. lalandi* in Australia, nor febantel administered to *S. quinqueradiata* in Japan resulted in a lower abundance of the skin parasite *B. seriolae*.

Praziquantel was first administered to *S. lalandi* by surface coating of feed. Fish rejected medicated feed, suggesting praziquantel affected its palatability. Fish treated with feed medicated with praziquantel had fewer *Z. seriolae* and *B. seriolae* than untreated fish. Praziquantel administered to *S. lalandi* by intubation allowed a more accurate dose to be tested without differential feeding or reduced palatability obstructing results, and resulted in fewer *Z. seriolae* (99.5-100 % reduction) and *B. seriolae* (91 – 97.7 % reduction). Intubated praziquantel also led to fewer recruitment life stages of *Z. seriolae* and *B. seriolae*, even at low doses, but did not completely eliminate them from *S. lalandi*. Praziquantel administered to *S. lalandi* alone and combined with cimetidine had high efficacy (>99%) against *Z. seriolae*. In comparison, praziquantel administered alone resulted in fewer *B. seriolae* (68.3 – 69.7 % reduction) than the same doses of praziquantel combined with cimetidine (36.9 - 40.9 % reduction). A 90.4 -100 % reduction in *H. heterocerca* was achieved when praziquantel was administered by intubation to *S. quinqueradiata* in Japan but there was only a 22-77.8 % reduction in *B. seriolae*. The dose of PZQ (150 mg kg<sup>-1</sup> body weight day<sup>-1</sup> for 3 days) on the label of a commercially available product used to treat *B. seriolae* in Japanese aquaculture resulted in a 50.9% reduction against *B. seriolae*, but completely eliminated *H. heterocerca*.

In trials against *Z. seriolae* and *B. seriolae* on *S. lalandi* in South Australia, I also screened 27 other anthelmintics and antiparasitics from the chemical groups: amprolium derivatives, benzimidazoles, benzyl ureas, diphosphate salts, imidazothiazoles, macrocyclic lactones, nitromidazoles, organophosphates, piperazines, salicylanilides, substituted phenols and tetrahydropyrimidines. Of these, only the benzimidazole, albendazole, was effective against *Z. seriolae* and none appeared to have an effect against *B. seriolae*.

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