

11-7-2015

The roles of individual variation and female choice in round goby (*Neogobius melanostomus*) acoustic signalling

Meghan Donovan
University of Windsor

Follow this and additional works at: <https://scholar.uwindsor.ca/etd>

Recommended Citation

Donovan, Meghan, "The roles of individual variation and female choice in round goby (*Neogobius melanostomus*) acoustic signalling" (2015). *Electronic Theses and Dissertations*. 5512.
<https://scholar.uwindsor.ca/etd/5512>

This online database contains the full-text of PhD dissertations and Masters' theses of University of Windsor students from 1954 forward. These documents are made available for personal study and research purposes only, in accordance with the Canadian Copyright Act and the Creative Commons license—CC BY-NC-ND (Attribution, Non-Commercial, No Derivative Works). Under this license, works must always be attributed to the copyright holder (original author), cannot be used for any commercial purposes, and may not be altered. Any other use would require the permission of the copyright holder. Students may inquire about withdrawing their dissertation and/or thesis from this database. For additional inquiries, please contact the repository administrator via email (scholarship@uwindsor.ca) or by telephone at 519-253-3000ext. 3208.

The roles of individual variation and female choice in round goby (*Neogobius melanostomus*) acoustic signalling

by
Meghan Donovan

A Thesis
Submitted to the Faculty of Graduate Studies
through Biological Sciences
in Partial Fulfillment of the Requirements for
the Degree of Master of Science at the
University of Windsor

Windsor, Ontario, Canada

© 2015 Meghan Donovan

The roles of individual variation and female choice in round goby (*Neogobius melanostomus*) acoustic signalling

by

Meghan Donovan

APPROVED BY:

A. Fisk
Department of Earth and Environmental Sciences

J. Ciborowski
Department of Biological Sciences

D. Higgs, Advisor
Department of Biological Sciences

August 17, 2015

DECLARATION OF ORIGINALITY

I hereby certify that I am the sole author of this thesis and that no part of this thesis has been published or submitted for publication.

I certify that, to the best of my knowledge, my thesis does not infringe upon anyone's copyright nor violate any proprietary rights and that any ideas, techniques, quotations, or any other material from the work of other people included in my thesis, published or otherwise, are fully acknowledged in accordance with the standard referencing practices. Furthermore, to the extent that I have included copyrighted material that surpasses the bounds of fair dealing within the meaning of the Canada Copyright Act, I certify that I have obtained a written permission from the copyright owner(s) to include such material(s) in my thesis and have included copies of such copyright clearances to my appendix.

I declare that this is a true copy of my thesis, including any final revisions, as approved by my thesis committee and the Graduate Studies office, and that this thesis has not been submitted for a higher degree to any other University or Institution.

ABSTRACT

Acoustic signalling in teleost fishes serves a variety of communicatory purposes, mostly centered around aggression and courtship. There is strong evidence that courtship calls are used by many fishes to locate mates, coordinate spawning, and for species recognition, but the possibility that these calls also act as honest signals is not yet highly explored. Calls produced by the male round goby, *Neogobius melanostomus*, were analyzed for dominant frequency, interpulse interval, duration and number of pulses in the call. Call characteristics were then analyzed for relationships to body morphometrics of total length, head width, total weight, and gonadosomatic index. Strong interactive relationships between male body traits and individual call characteristics were found. Females were shown to have a preference for longer interpulse interval, suggesting that calls are capable of honestly signalling male body size. These findings suggest that acoustic honest signalling as well as individual discrimination exists in this species.

DEDICATION

I dedicate this thesis to my family and friends for all of their love, support and understanding.

ACKNOWLEDGEMENTS

Firstly, I would like to thank my advisor, Dr. Dennis Higgs, for all of his help and guidance throughout this project. You really went above and beyond as a supervisor and I am incredibly happy that I joined your lab. I have to thank you and Kirsten both for making these two years really enjoyable, I will always remember our lab's humorous lunchtime conversations.

I am very grateful to Dr. Jan Ciborowski and Dr. Aaron Fisk for serving on my committee and for all of their suggestions throughout my project. I would also like to thank Dr. Oliver Love for serving as the chair on my defence.

I would also like to thank Dr. Christina Semeniuk for your help in interpreting interaction effects. Thank you to Drs. Clara Amorim, Stefano Malavasi and Gianluca Polgar for allowing me to use your oscillograms in this thesis. I also want to thank Dr. Mattias Johansson for your suggestions on the statistics of Chapter 3.

A huge thank you to everyone who came out to help me collect gobies in the field, especially Christian Carnevale, Bronson Goodfellow, Stephanie Cervi, Peter Cunha and Daniel Dubovan. Sarah Humphrey and Megan Mickle, thank both of you immensely for spending day after day fishing in the heat with me. It would not have been anywhere near as much fun without you ladies. Thank you to Shyane Wieggers for her help with video analysis. Thank you Mallory Wiper for all the advice and fantastic times I had in the lab with you. Thank you to all of the undergraduate students in the Higgs lab for making these past two years great.

To all the friends I have made during my time in Windsor, thank you all so, so much, you have all made my time here amazing and I am incredibly happy to have gotten to know you all.

I would like to thank all of the staff as well, especially Linda Sterling and Melissa Gabrieau down in animal care, Bob Hodge for all his technical support, the staff at the Technical Support Centre for all their help, Rodica Leu and Ingrid Churchill, and Nancy Barkley.

This research was funded by grants from NSERC awarded to Dennis Higgs and by funding from OFAH.

TABLE OF CONTENTS

AUTHOR’S DECLARATION OF ORIGINALITY	iii
ABSTRACT.....	iv
DEDICATION	v
ACKNOWLEDGEMENTS	vi
LIST OF TABLES	x
LIST OF FIGURES	xi
CHAPTER	
I. REVIEW ON SEXUALLY SELECTIVE BIOACOUSTICS IN THE Gobiidae	
Thesis Objectives.....	1
Sensory Systems in an Aquatic Environment.....	1
Underwater Sound Propagation	2
Uses of Sound in Fish	3
Sound Production in Fish.....	5
Sound Production in the Gobiidae	8
Phylogenetic Links Between Sound	8
Call structure	10
Seasonality and Nesting	11
Sound Producing Mechanisms.....	13
Bioacoustics as Sexually Selected Signals.....	15
Recommendations and Conclusions	15
References.....	17
II. VARIATION IN THE CALL STRUCTURE OF MALE ROUND GOBIES (<i>NEOGOBIUS MELANOSTOMUS</i>) IN RELATION TO BODY MORPHOMETRICS	
Introduction.....	21
Model Organism	23
Sampling Methods	23
Experimental Procedure	24
Call Analysis	25
Statistical Analysis.....	25
Results.....	27
Discussion	31
References.....	38
III. EXAMINING THE POTENTIAL FOR ACOUSTIC HONEST SIGNALLING IN THE ROUND GOBY (<i>NEOGOBIUS MELANOSTOMUS</i>) THROUGH FEMALE CHOICE	

Introduction.....	42
Sampling Methods	43
Effects of Hormone Injection.....	44
Male Call Playback	45
Statistical Analysis	47
Results.....	48
Discussion	52
References.....	58
IV. CONCLUSIONS AND RECOMMENDATIONS	
References.....	64
VITA AUCTORIS	65

LIST OF TABLES

Table 1.1. Sound production across families.	6
Table 1.2. Reproductive seasons of soniferous gobies.	12
Table 2.1. Inter- and intra-male variability of call frequency.	28
Table 2.2. Loadings of call and body measures.	29
Table 3.1. Statistical results of female choice GLMs.	53

LIST OF FIGURES

Figure 1.1. Phylogenetic tree of soniferous gobies.	9
Figure 2.1. The relationship between HWxTW, HWxTL and DF.	30
Figure 2.2. The relationship between HWxTW, HWxTL and IPI.	32
Figure 2.3. The relationship between HWxTLxTW and duration.	33
Figure 3.1. Diagram of the female choice experimental set-up.	46
Figure 3.2. Time spent in arena speaker across trials.	50
Figure 3.3. Number of females showing call preference.	51

Chapter I

REVIEW ON SEXUALLY SELECTIVE BIOACOUSTICS IN THE GOBIIDAE

Thesis Objectives

This thesis aims to determine if acoustic signalling in the round goby functions as an honest signal. For this purpose, I define honest signalling as a communicatory signal where some measurable aspect of male condition is related to the signals reliability and where females are able to accurately judge and have a preference for signal variability. The first portion of the study, presented in Chapter 2, examines the correlation between the male body characteristics of head width, total length, total weight and gonadosomatic index and the call characteristics of call duration, dominant frequency and average interpulse interval. If any of these call characteristics are strongly correlated to male body size, round goby courtship calls may act as reliable predictors of male body size. The second portion of this thesis, Chapter 3, focuses on female call preference, to determine if the relationship present between male body measures and call characteristics is a true honest signal. Confirming female preference is critical to defining these courtship calls as honest signals.

Sensory Systems in an Aquatic Environment

The underwater environment is a complex system, which heavily influences signal transmission and detection, therefore, communicating with conspecifics poses a challenge for inhabitants in such environments. One of the barriers that must be overcome is the challenge of communicating with conspecifics. To do this, many species will combine several types of sensory displays simultaneously (Saunders et al., 2010; Smith and van

Staadén, 2009; Thaker et al., 2006). While multimodal signals can be perceived as being redundant, the transference of the same information through several sensory systems assures that the receiver will perceive the sender's communication accurately (Partan and Marler, 2005). Redundant multimodal signals become increasingly important in habitats where the information communicated might be misinterpreted. For instance, visual displays are often limited to very short distances due to poor visibility, olfaction is complicated by highly variable water flow and auditory cues can be masked by the surrounding biotic and abiotic acoustic environment. Of these three common underwater signalling modalities, bioacoustic signals can often be transmitted the furthest, increasing the chances that the sender's information will reach a desired target. In many species, these acoustic displays are low in frequency (Ladich, 2014), allowing the sound to travel a great distance without attenuating in ideal conditions but causing propagation difficulties in shallow water (Rogers and Cox, 1988). Communicating with low frequency sounds can also create issues with sound reception and localization as the wavelength of the sound is larger than the distance between the ears of the fish (Popper and Fay, 1973) and yet the majority of fish communication calls are in the low frequency range (Ladich, 2014).

Underwater Sound Propagation

The basic physics of how sound travels underwater are important to understanding the reception and production of sound in the aquatic environment. Many differences exist between sound transmission through air compared to water. The substantial increase in speed from 332 m/s in air to 1484 m/s in water greatly increases the wavelengths of sounds underwater. Attenuation is the loss of energy as the sound propagates due to

effects like absorption and scattering. Low frequencies can travel great distances with very little attenuation in an unbounded medium, however higher frequency sounds tend to travel only very short distances. Absorption of sound also increases with increasing water temperature and salinity. This lack of attenuation means that the background noise caused by biotic and abiotic factors is immense, increasing the overall noise levels by up to 30 dB (Rogers and Cox, 1988). This increase in sound intensity becomes incredibly important when considering that communicatory sounds will potentially be masked, preventing the fish from hearing the sound of interest (Fay and Popper, 2012). As sound passes across a thermocline or halocline, the speed at which the sound wave is propagating will change, causing refraction of the sound wave. Similarly, when a sound wave encounters the surface it will both refract and reflect the sound. The substrate that the sound wave encounters will also affect how it travels; harder surfaces will reflect sound better than soft surfaces (Nedelec et al., 2015). As the angle of incidence increases, more sound will be reflected than refracted, until refraction becomes undetectable. This effect becomes important in shallow water where much of the energy of the sound wave is refracted instead of traveling to the potential receiver and can greatly complicate the use of sound as a communication signal in shallow waters.

Uses of Sound in Fish

Several types of calls are produced by fishes, most notably alarm, agonistic and courtship calls (Smith, 1992; Bass and McKibben, 2003; Van Staaden and Smith, 2011). Typically, fish produce alarm calls when being handled or when the fish encounters a predator (Smith, 1992). Alarm calls are those that elicit a response by conspecifics, while distress calls are meant to dissuade the predator from attacking (Smith, 1992). The

Atlantic cod (*Gadus morhua*) will produce grunting calls when chased by a novel object or by a predator and will continue to grunt when cornered and immobile (Brawn, 1961). In the cod, the cues may be used as a warning to nearby conspecifics or as a means of attracting a secondary predator, potentially allowing escape (Smith, 1992). A more clear alarm signal exists in the longspine squirrelfish (*Holocentrus rufus*) where an encounter with a predator will cause the squirrelfish to produce a staccato call, which either elicits retreat or mobbing behaviour by nearby conspecifics (Winn et al., 1964).

Many fish species are known to produce calls under agonistic contexts, especially when competing for food or mates. Competition for food stimulates the grey gurnard, *Eutrigla gurnardus*, to produce knocking sounds when grasping food and grunting sounds when doing frontal displays to competitors (Amorim et al., 2004). Growling and other sounds produced by gurnards may act as keep-away signals to nearby competitors (Amorim and Hawkins, 2000; Amorim et al., 2004). Acoustic signalling is also often used by territory holding males in both an agonistic context as well as in courtship. Parental male plainfin midshipmen, *Porichthys notatus*, have two call types, the grunt used in agonistic displays and the hum used in attracting mates (McKibben and Bass, 1998). Hums will elicit a phonotactic response from females, but grunts and modified calls that do not closely resemble a hum receive no response from females (Bass and McKibben, 2003).

Perhaps the most well documented uses of acoustic signalling in fish are courtship calls. These calls are used by monogamous pairs (Boyle and Tricas, 2011), group spawners (Rowe and Hutchings, 2003) and territory holders when attracting mates (Amorim et al., 2011). Courtship calls are used to locate mates (McKibben and Bass, 1998), ensure correct species recognition (Rollo and Higgs, 2008) and potentially even

act as honest signals of mate quality (Amorim et al., 2013a). In species like the burbot, *Lota lota*, which spawns under the cover of sea-ice, acoustic signalling seems to be critically important to the timing of gamete release (Cott et al., 2014). Calls are likely very important for species that nest under objects where the caller may be hard to locate (Lugli et al., 1997). Many fish calls may serve multiple purposes and have simply yet to be examined.

Sound Production in Fish

There is an exceptionally diverse variety of fish that are known to produce sounds as a means of communication. Fish and Mowbray (1970) provided extensive documentation on 36 families of Osteichthyes fish with sound producing members and one species of Chondrichthyes. Several more soniferous families have been discovered since the publication of this compendium. Over 700 species of fish are now known to produce sound (Kaatz, 2002; Luczkovich et al., 2008) and there are many bioacoustic sounds attributed to fish that have yet to be identified (Anderson et al., 2008). In the cases of the families examined by Fish and Mowbray (1970), all species documented were producing agonistic calls in response to electric or mechanical stimulus. Fish will also produce these calls under agonistic and aggressive interactions, such as when a predator or other intruder has entered their territory (Ladich, 1997). Many of these sound producing families also contain species that produce courtship sounds. Courtship sounds are typically only produced by males, whereas aggressive and agonistic calls are commonly produced by both sexes (Ladich, 1997). Several species have separate calls that they produce in agonistic and courtship contexts (Table 1.1). However, the characterization of these sound types may vary between researchers, leaving the simple

Table 1.1

Table 1.1. Sound production across families. The qualitative descriptions of fish sounds across 42 families of soniferous fishes. Most calls are described similarly and many families contain species that produce distinct calls for courtship and agonistic displays.

Family	Agonistic call	Uncharacterized sound	Courtship call	Reference
Elopidae	thumps			Fish and Mowbray (1970)
Albulidae	thumps			Fish and Mowbray (1970)
Clupeidae	thumps and knocks			Fish and Mowbray (1970)
Arridae	sobs, yelps, grunts			Fish and Mowbray (1970)
Anguillidae	clucking and knocks			Fish and Mowbray (1970)
Gadidae	thumps		grunts and hums	Fish and Mowbray (1970); Cott et al. (2014)
Syngnathidae		snaps and clicks during feeding		Fish and Mowbray (1970)
Holocentridae	grunts			Fish and Mowbray (1970)
Centropomidae	thumps and knocks			Fish and Mowbray (1970)
Serranidae	thumps		pulses and a tonal sweep	Fish and Mowbray (1970); Lobel (1992)
Lutjanidae	thumps			Fish and Mowbray (1970)
Pomatomidae	thumps and knocks			Fish and Mowbray (1970)
Carangidae	thumps and knocks			Fish and Mowbray (1970)
Gerridae	weak knocks			Fish and Mowbray (1970)
Pomadasyidae	grunts			Fish and Mowbray (1970)
Sciaenidae	knocks and croaks		drumming	Fish and Mowbray (1970); Connaughton et al. (2002)
Mullidae	thumps and knocks			Fish and Mowbray (1970)
Sparidae	thumps and knocks			Fish and Mowbray (1970)
Kyphosidae	thumps and knocks			Fish and Mowbray (1970)
Ephippidae	thumps, knocks and grunts			Fish and Mowbray (1970)
Chaetodontidae	thumps and knocks; pulses			Fish and Mowbray (1970); Boyle and Tricas (2011)
Pomacentridae	thumps, knocks and clicks		clicks	Fish and Mowbray (1970); Parmentier et al. (2010)
Labridae	thumps and knocks		pulses	Fish and Mowbray (1970); Boyle and Cox (2009)
Scaridae	clicks and knocks		group spawning, individual sounds indiscernible	Fish and Mowbray (1970); Lobel (1992)

Acanthuridae	Knocks and grunts			Fish and Mowbray (1970)
Triglidae	growls, grunts and barks			Fish and Mowbray (1970); Amorim et al. (2004)
Cottidae	growls			Fish and Mowbray (1970)
Dactylopteridae	barks			Fish and Mowbray (1970)
Sphyrinaeidae		thumps and knocks with swift movement		Fish and Mowbray (1970)
Polynemidae	knocks			Fish and Mowbray (1970)
Balistidae	grunts, knocks, scrapes, thumps and clicks			Fish and Mowbray (1970)
Ostraciidae	groans, clicks and scrapes			Fish and Mowbray (1970)
Tetraodontidae		"erks" with inflation		Fish and Mowbray (1970)
Diodontidae		whines with inflation		Fish and Mowbray (1970)
Molidae	grunts	high-pitched groan when lifted from the water		Fish and Mowbray (1970)
Batrachoididae	grunts		boatwhistle	Fish and Mowbray (1970); Amorim et al. (2011)
Blenniidae			grunt	DeJong et al. (2007)
Cichlidae			pulses	Maruska et al. (2012)
Percidae			knocks, purrs and drums	Speares and Johnston (2011)
Mormyridae			grunts and moans	Crawford et al. (1997)
Doradidae	grunts and growls		pulse	Kaatz and Stewart (2012)
Auchenipteridae	grunts and growls		pulse	Kaatz and Stewart (2012)

description of the sound up for debate (Parmentier et al., 2006). While the majority of fish sounds are categorized as pulsed sounds resembling grunts, thumps or knocks, there is a huge amount of variation from species to species, and in many cases calls are species specific (Luczkovich et al., 2008).

Sound Production in the Gobiidae

The ability to produce sound has been documented across many taxa of fish, with one of the most intensively studied families being the Gobiidae. As these fish lack specialized structures that aid in hearing, they can be thought of as hearing generalists (Popper and Fay, 2011); the actual amount of sound detection abilities in these fish is debated as the group is exceptionally diverse. Currently, the family contains some 1950 species across 210 genera (Nelson, 2006) of which only 23 species are documented as sound producers (Lugli et al., 1997; Parmentier et al., 2013).

Phylogenetic Links Between Sound Production

Where the Gobiidae is such an expansive and widely distributed group, the taxonomic links between members are still under some consideration. Many studies use different names for some call producing species, most notably *Padogobius martensii* and *P. bonelli*, which appear to be the same species (Nocita and Vanni, 2001). Very detailed work has been done to attempt to organize this clade using morphological and genetic analyses (Thacker and Roje, 2011). While the phylogenetics of the Gobiidae is still unclear, there appears to be no evolutionary patterns in call structure with some species using pulse trains while other use simple grunts, regardless of phylogenetic position (Figure 1.1). There are also clear differences in call structure between species, allowing

Figure 1.1

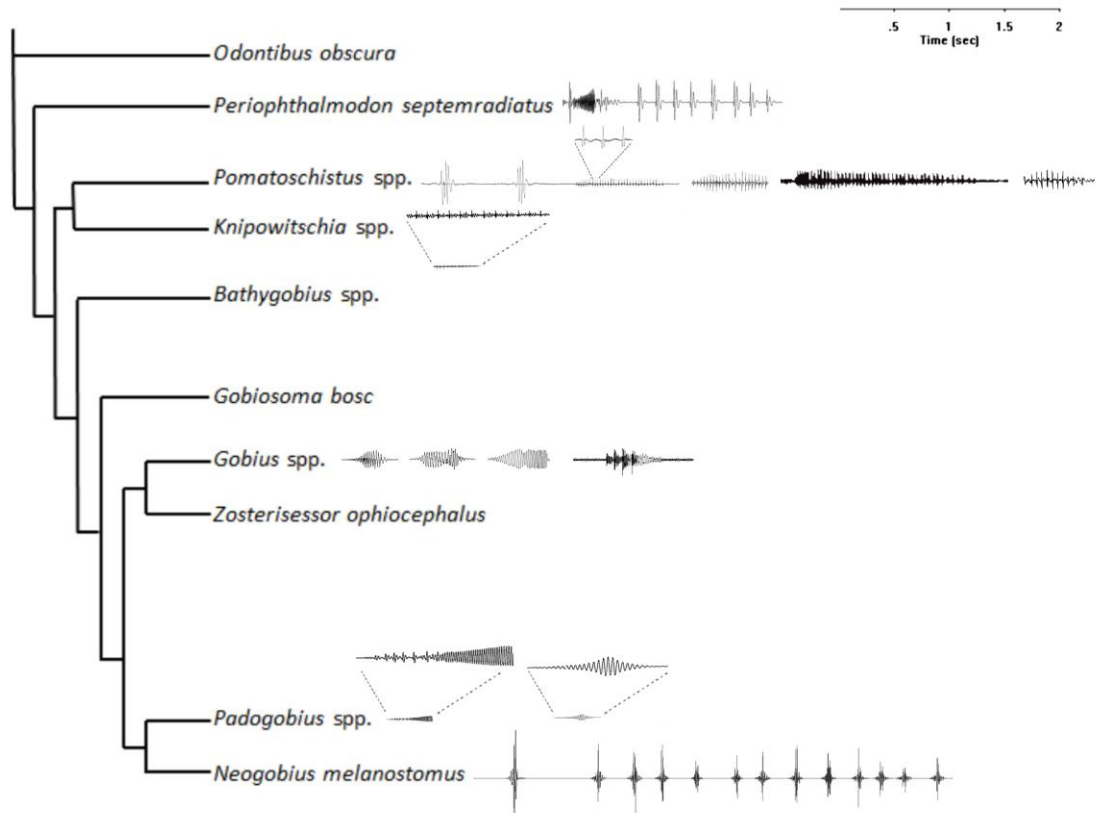


Figure 1.1. Phylogenetic tree of soniferous gobies. All oscillograms are scaled to the same time frame. Individual calls represent different species, from left to right in each genus where species are not labeled, calls belong to *Pomatoschistus*: *P. pictus*, *P. canestrinii*, *P. marmoratus*, *P. minutus*; *Knipowitschia*: *K. panizzae*; *Gobius*: *G. niger*, *G. cobitis*, *G. paganellus*, *G. cruentatus*; *Padogobius*: *P. martensii*, *P. nigricans*. This phylogeny was modified from Thacker and Roje (2011), to include soniferous species in the Gobiidae. Oscillograms were used with permission from Drs. S. Malavasi, C. Amorim, and G. Polgar.

for species recognition when examining the calls. The soniferous species in this family are widely distributed across the clade, and yet, more ancestral species like the mudskipper, *Periophthalmodon septemradiatus*, also produce sound (Polgar et al., 2011). This suggests that acoustic signalling may be far more common in this family than the few species already described.

Call structure

Several terms are used by different researchers seemingly interchangeably when referring to certain sounds, such as drums, grunts and pulses. A call is usually referred to as “drumming” when grunts are repeated in a grunt or pulse train (Lugli et al., 1997). A grunt is defined as a single, low frequency pulse and a pulse is a short, continuous sound burst. The grunt is possibly the most commonly produced call by gobiids. Grunts are generally low in frequency, typically between 100 and 200 Hz for most species (Malavasi et al., 2008). Grunts may be produced singly, in pulse trains resembling a drumming sound or as part of complex sound that includes a grunt train and a tonal sound (Lugli et al., 1995). The tonal sound is a continuous waveform with a low range of frequency modulation. It is possible that the tonal sound is produced by reducing the inter-pulse duration to the point that the pulse train becomes a single tone (Malavasi et al., 2008). Depending on the method of sound production, some species should be incapable of making tonal sounds as these sounds would likely depend on rapid stridulation, something not all species are capable of (Stadler, 2002). Similar to the pulse train is another call described as a stutter where the call is weaker than individual grunts, but otherwise similar in structure (Stadler 2002). The weakness of this call is potentially caused by the alternation of body sides when the fish is ejecting water through the gills

where single grunts would have water ejected simultaneously from both sides (Stadler, 2002). This method of calling has only been described in one species thus far, the notchtongue goby, *Bathygobius curacao*, though may be found in other species of this genus (Tavolga, 1958; Stadler, 2002). Differentiation of call components may be important for species recognition when several species of gobies live sympatrically to avoid hybridization (Blair, 1958).

Seasonality and Nesting

Sound production is generally linked to courtship behaviour in the Gobiidae and as such occurs during the spawning seasons (Table 1.2). Typically, acoustic signals are paired with visual (Lugli et al., 1995) and olfactory stimuli (Kasurak et al., 2012). Visual displays vary by species, but often the male will actively fan his pectoral fins (Meunier et al., 2013), swim in and out of the nest and perform tail beats (Lugli et al., 1995). Calls are produced solely by males (Parmentier et al., 2013) and occur during aggressive, courtship and spawning interactions (Stadler, 2002). During the spawning season, male gobies guard nests from which they produce these calls (Rollo et al., 2007). Generally, these nests are simple cavities under rocks or shells. However, this resource may be limiting, causing males to be territorial for the best nesting sites (Lugli et al., 1992). The distance between nesting freshwater goby (*Padogobius martensii*) increases with male size and when females are not a limiting factor, larger males will take over larger nesting sites (Lugli et al., 1992). When deciding on a nesting site, male sand gobies (*Pomatoschistus minutus*) preferentially choose the nest cavity that looks larger from the outside (Lindström, 1992). Larger males have a tendency to win more fights, thus gaining better

Table 1.2

Table 1.2. Reproductive seasons of soniferous gobies. The reproductive season of 17 of the 23 known soniferous gobies. The taxonomy of *Padogobius bonelli* and *Padogobius martensii* is unclear however and these may be the same species.

Species	Seasonality	Location	Reference
<i>Bathygobius fuscus</i>	June-September	Indo-Pacific	Zhang and Takemura 1989
<i>Gobius cobitis</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Gobius cruentatus</i>	March-May	Mediterranean Sea	Sebastianutto et al. 2008
<i>Gobius niger</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Gobius paganellus</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Knipowitschia panizzae</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Knipowitschia punctatissima</i>	March-May	Italy	Lugli et al. 1997
	March-June	Venice lagoon	Malavasi et al. 2008
<i>Neogobius melanostomus</i>	May-August	Detroit River	MacInnis and Corkum 2000
<i>Odontobutis obscura</i>	May-July	Japan	Takemura 1984
<i>Padogobius bonelli</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Padogobius martensii</i>	March-May	Italy	Lugli et al. 1997
<i>Padogobius nigricans</i>	March-May	Italy	Lugli et al. 1997
	March-June	Venice lagoon	Malavasi et al. 2008
<i>Pomatoschistus canestrinii</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Pomatoschistus marmoratus</i>	March-June	Venice lagoon	Malavasi et al. 2008
<i>Pomatoschistus minutus</i>	May-August	Baltic Sea	Lindström and Lugli 2008
	March-June	Venice lagoon	Malavasi et al. 2008
<i>Pomatoschistus pictus</i>	January-May	Portugal	Amorim et al. 2013
<i>Zosterisessor ophiocephalus</i>	March-June	Venice lagoon	Malavasi et al. 2008

nesting sites (Magnhagen and Kvarnemo, 1989). Larger nesting sites have a greater surface area and as such a greater number of eggs are deposited (Lugli et al., 1992). However, larger males are not necessarily preferred by females (Magnhagen and Kvarnemo, 1989). Likewise, females are not more likely to choose the winner of a fight (Forsgren, 1997). In the sand goby, preferred males receive more eggs and provide greater parental investment (Forsgren et al., 1996). These factors are somewhat confounding however, as males that provide more care to larger clutches may not inherently be better fathers (Forsgren, 1997). The importance of individual features of interest to females choosing mates is difficult to discern and may differ between species and across habitats.

Sound Producing Mechanisms

Sound production has been documented in at least 23 species in the Gobiidae (see Lugli et al., 1997 and Parmentier et al., 2013) and is likely displayed in other species as well. Several hypotheses exist on the mechanism behind sound production in this family and, due to the diversity of the group, more than one mechanism seems likely. Most recently, Parmentier et al. (2013) proposed that the levator pectoralis muscles are key to sound production, especially in species lacking swim bladders, though the exact mechanism remains unknown. Other possible mechanisms are by rubbing the pharyngeal plates together, coupled with an amplification through the swim bladder in species that possess one (Takemura, 1984; Lugli 1995) as well as squirting water through the opercular openings (Tavolga, 1958) though this latter method is debated (Stadler, 2002; Parmentier et al., 2013). Calls produced by *Odontobutis obscura* are amplified by the muscular connection between the pharyngeal teeth to the 2-4 vertebrae, which in turn are

in direct connection with the thin-walled swim bladder (Takemura, 1984). At least one vocal species lacks necessary structures for producing stridulatory calls (Stadler, 2002). However, the absence of pharyngeal teeth seems unlikely as other members of the *Bathygobius* have pharyngeal teeth (Miller and Smith, 1989). In vocal species of gobies, the large pectoral fins are often erected when fish are calling, potentially using the membrane as an amplifier for the sound (Parmentier et al., 2013) though this behaviour is usually attributed to being part of visual or olfactory displays (Meunier et al., 2013).

Gobies may use their nest cavities as a way of amplifying their calls, as nests tend to only have one opening, which alters the properties of the sound being emitted (Lugli, 2012). The amplificatory effect depends on the material of the nest as well as the frequency of the call. Nesting structures that have a density similar to water do not seem to have the capacity to act as resonance chambers (Lugli, 2012). Similarly, some structures may not interfere with the amplitude of calls, as toadfish boatwhistles were unaltered by terra cotta nesting structures (Barimo and Fine, 1998) due to the resonance frequency of the calls (Lugli, 2012). The form of these nesting structures is another factor to consider, as the opening to a cavity can often affect the radiation of the sound from its source (Fletcher, 2004). In ground crickets, the burrow from which the animal calls significantly increases the loudness of the calls (Bailey et al., 2001). The effectiveness of the nest cavity acting as a resonance chamber also depends on the coupling between the nest cavity and the substrate, as any discontinuity causes a large drop in the shelter's ability to act as an amplifier (Lugli, 2012). Sand gobies are known to move sand around the site of their nest, such that a sand pile forms over top of the nest and the entrance can become quite small (Lugli, 2013). This would prevent the goby's calls from escaping

from any openings other than the entrance, ensuring that the nest cavity could best act as a single-opening resonance chamber.

Bioacoustics as Sexually Selected Signals

In the Gobiidae, calls are produced both for aggressive purposes and in reproductive contexts (Lugli et al., 1995; Kasurak et al., 2012). In some species, sound is only produced in the presence of a gravid female, usually in combination with other courtship displays (Lindström and Lugli, 2000). In the freshwater goby complex calls are produced in combination with tail beats to encourage the female to enter the nest and sound production slowly decreases as spawning proceeds (Lugli et al., 1995). Sound may be produced as a means of encouraging females to approach or enter the nest (Malavasi et al., 2009) or once the female has entered as part of a multisensory display to show paternal reliability (Amorim et al., 2013b). The panzarolo goby (*Knipowitschia punctatissima*) calls only when a female is present inside the nest and calling ceases upon the female's departure (Lugli et al., 1995). Female round goby (*Neogobius melanostomus*) are known to be able to differentiate between calls of several species and show a preference for conspecific calls (Rollo and Higgs, 2008). This species is able to localize to sound sources (Rollo et al., 2007) and males may use calls to attract mates from a distance. Actively displaying males are more attractive mates ensuring proper parental care (Forsgren, 1997).

Recommendations and Conclusions

There is still much to discover about sound production and its uses in fish. In many cases, bioacoustics has only been explored in a few species within a family, leaving

gaps in knowledge about how these sounds are used. Since sounds produced by fish are species specific, a wide range of possibilities exist for human use. Bioacoustics has the capacity to help researchers better understand species interactions and aid in conservation and management without the often destructive consequences of capturing fish. However, in order to get to the point where bioacoustics could replace more traditional techniques, more research on a greater diversity of fish must be done. Some recommendations that could increase the success of future studies include 1) Researchers should share calls of known species through online sound libraries; 2) For management purposes, acoustically active fish should be characterized, enabling the use of passive acoustic monitoring of population health and invasion fronts; 3) The phylogenetic backgrounds of calling species should be analyzed in greater detail; and 4) The linkages between male quality and sound production should be further examined.

References

- Amorim, M. C. P., & Hawkins, A. D. (2000). Growling for food: acoustic emissions during competitive feeding of the streaked gurnard. *Journal of Fish Biology*, 57(4), 895-907.
- Amorim, M. C. P., Stratoudakis, Y., & Hawkins, A. D. (2004). Sound production during competitive feeding in the grey gurnard. *Journal of Fish Biology*, 65(1), 182-194.
- Amorim, M. C. P., Simões, J. M., Almada, V. C., & Fonseca, P. J. (2011). Stereotypy and variation of the mating call in the Lusitanian toadfish, *Halobatrachus didactylus*. *Behavioral Ecology and Sociobiology*, 65(4), 707-716.
- Amorim, M. C. P., Pedroso, S. S., Bolgan, M., Jordao, J. M., Caiano, M., & Fonseca, P. J. (2013a). Painted gobies sing their quality out loud: acoustic rather than visual signals advertise male quality and contribute to mating success. *Functional Ecology*, 27(2), 289-298.
- Amorim, M. C. P., da Ponte, A. N., Caiano, M., Pedroso, S. S., Pereira, R., & Fonseca, P. J. (2013b). Mate preference in the painted goby: the influence of visual and acoustic courtship signals. *The Journal of experimental biology*, 216(21), 3996-4004.
- Anderson, K. A., Rountree, R. A., & Juanes, F. (2008). Soniferous fishes in the Hudson River. *Transactions of the American Fisheries Society*, 137(2), 616-626.
- Bailey, W. J., Bennet-Clark, H. C., & Fletcher, N. H. (2001). Acoustics of a small australian burrowing cricket the control of low-frequency pure-tone songs. *The Journal of Experimental Biology*, 204(16), 2827-2841.
- Barimo, J. F., & Fine, M. L. (1998). Relationship of swim-bladder shape to the directionality pattern of underwater sound in the oyster toadfish. *Canadian Journal of Zoology*, 76(1), 134-143.
- Bass, A. H., & McKibben, J. R. (2003). Neural mechanisms and behaviors for acoustic communication in teleost fish. *Progress in Neurobiology*, 69(1), 1-26.
- Blair, W. F. (1958). Mating call in the speciation of anuran amphibians. *American Naturalist*, 27-51.
- Boyle, K. S., & Cox, T. E. (2009). Courtship and spawning sounds in bird wrasse *Gomphosus varius* and saddle wrasse *Thalassoma duperrey*. *Journal of fish biology*, 75(10), 2670-2681.
- Boyle, K. S., & Tricas, T. C. (2011). Sound production in the longnose butterflyfishes (genus *Forcipiger*): cranial kinematics, muscle activity and honest signals. *The Journal of experimental biology*, 214(22), 3829-3842.
- Brawn, V. M. (1961). Sound production by the cod (*Gadus callarias* L.). *Behaviour*, 18(4), 239-255.
- Connaughton, M. A., Fine, M. L., & Taylor, M. H. (2002). Weakfish sonic muscle: influence of size, temperature and season. *Journal of Experimental Biology*, 205(15), 2183-2188.
- Cott, P. A., Hawkins, A. D., Zeddies, D., Martin, B., Johnston, T. A., Reist, J. D., Gunn, J. M., & Higgs, D. M. (2014). Song of the burbot: Under-ice acoustic signalling by a freshwater gadoid fish. *Journal of Great Lakes Research*, 40(2), 435-440.
- Fay, R. R., & Popper, A. N. (2012). Fish hearing: new perspectives from two 'senior' bioacousticians. *Brain, Behavior and Evolution*, 79(4), 215-217.

- Fish, M. & Mowbray, W. (1970). *Sounds of western North Atlantic fishes; a reference file of biological underwater sounds*. Baltimore: Johns Hopkins Press.
- Fletcher, N. H. (2004). A simple frequency-scaling rule for animal communication. *The Journal of the Acoustical Society of America*, 115(5), 2334-2338.
- Forsgren, E. (1997). Female sand gobies prefer good fathers over dominant males. *Proceedings of the Royal Society of London. Series B: Biological Sciences*, 264(1386), 1283-1286.
- Kaatz, I. M. (2002). Multiple sound-producing mechanisms in teleost fishes and hypotheses regarding their behavioural significance. *Bioacoustics*, 12(2-3), 230-233.
- Kaatz, I. M., & Stewart, D. J. (2012). Bioacoustic variation of swimbladder disturbance sounds in Neotropical doradoid catfishes (Siluriformes: Doradidae, Auchenipteridae): Potential morphological correlates. *Current Zoology*, 58(1).
- Kasurak, A. V., Zielinski, B. S., & Higgs, D. M. (2012). Reproductive status influences multisensory integration responses in female round gobies, *Neogobius melanostomus*. *Animal Behaviour*, 83(5), 1179-1185.
- Ladich, F. (1997). Agonistic behaviour and significance of sounds in vocalizing fish. *Marine & Freshwater Behaviour & Physiology*, 29(1-4), 87-108.
- Ladich, F. (2014). Fish bioacoustics. *Current opinion in neurobiology*, 28, 121-127.
- Lindström, K. (1992). The effect of resource holding potential, nest size and information about resource quality on the outcome of intruder-owner conflicts in the sand goby. *Behavioral Ecology and Sociobiology*, 30(1), 53-58.
- Lindström, K., & Lugli, M. (2000). A quantitative analysis of the courtship acoustic behaviour and sound patterning in male sand goby, *Pomatoschistus minutus*. *Environmental biology of fishes*, 58(4), 411-424.
- Lobel, P. S. (1992). Sounds produced by spawning fishes. *Environmental Biology of Fishes*, 33(4), 351-358.
- Luczkovich, J. J., Mann, D. A., & Rountree, R. A. (2008). Passive acoustics as a tool in fisheries science. *Transactions of the American Fisheries Society*, 137(2), 533-541.
- Lugli, M., Bobbio, L., Torricelli, P., & Gandolfi, G. (1992). Breeding ecology and male spawning success in two hill-stream populations of the freshwater goby, *Padogobius martensi*. *Environmental biology of fishes*, 35(1), 37-48.
- Lugli, M., Pavan, G., Torricelli, P., & Bobbio, L. (1995). Spawning vocalizations in male freshwater gobiids (Pisces, Gobiidae). *Environmental Biology of Fishes*, 43(3), 219-231.
- Lugli, M., Torricelli, P., Pavan, G., & Mainardi, D. (1997). Sound production during courtship and spawning among freshwater gobiids (Pisces, Gobiidae). *Marine & Freshwater Behaviour & Physiology*, 29(1-4), 109-126.
- Lugli, M. (2012). Acoustics of fish shelters: Frequency response and gain properties. *The Journal of the Acoustical Society of America*, 132(5), 3512-3524.
- Lugli, M. (2013). Sand pile above the nest amplifies the sound emitted by the male sand goby. *Environmental biology of fishes*, 96(8), 1003-1012.
- MacInnis, A. J., & Corkum, L. D. (2000). Fecundity and reproductive season of the round goby *Neogobius melanostomus* in the upper Detroit River. *Transactions of the American Fisheries Society*, 129(1), 136-144.

- Magnhagen, C., & Kvarnemo, L. (1989). Big is better: the importance of size for reproductive success in male *Pomatoschistus minutus* (Pallas)(Pisces, Gobiidae). *Journal of Fish Biology*, 35(6), 755-763.
- Malavasi, S., Collatuzzo, S., & Torricelli, P. (2008). Interspecific variation of acoustic signals in Mediterranean gobies (Perciformes, Gobiidae): comparative analysis and evolutionary outlook. *Biological Journal of the Linnean Society*, 93(4), 763-778.
- Malavasi, S., Valerio, C., & Torricelli, P. (2009). Courtship sounds and associated behaviours in the Canestrini's goby *Pomatoschistus canestrinii*. *Journal of fish biology*, 75(7), 1883-1887.
- Maruska, K. P., Ung, U. S., & Fernald, R. D. (2012). The African cichlid fish *Astatotilapia burtoni* uses acoustic communication for reproduction: sound production, hearing, and behavioral significance. *PLoS One*, 7(5).
- McKibben, J. R., & Bass, A. H. (1998). Behavioral assessment of acoustic parameters relevant to signal recognition and preference in a vocal fish. *The Journal of the Acoustical Society of America*, 104(6), 3520-3533.
- Meunier, B., White, B., & Corkum, L. D. (2013). The role of fanning behavior in water exchange by a nest-guarding benthic fish before spawning. *Limnology and Oceanography: Fluids and Environments*, 3(1), 198-209.
- Miller, P. J., & Smith, R. (1989). The West African species of *Bathygobius* (Teleostei: Gobiidae) and their affinities. *Journal of Zoology*, 218(2), 277-318.
- Nedelec, S. L., Simpson, S. D., Holderied, M., Radford, A. N., Lecellier, G., Radford, C., & Lecchini, D. (2015). Soundscapes and living communities in coral reefs: temporal and spatial variation. *Marine Ecology Progress Series*, 524, 125-135.
- Nelson, J. (2006). *Fishes of the world*. Hoboken, N.J: John Wiley.
- Parmentier, E., Vandewalle, P., Frederich, B., & Fine, M. L. (2006). Sound production in two species of damselfishes (Pomacentridae): *Plectroglyphidodon lacrymatus* and *Dascyllus aruanus*. *Journal of fish biology*, 69(2), 491-503.
- Parmentier, E., Kéver, L., Casadevall, M., & Lecchini, D. (2010). Diversity and complexity in the acoustic behaviour of *Dacyllus flavicaudus* (Pomacentridae). *Marine biology*, 157(10), 2317-2327.
- Parmentier, E., Kéver, L., Boyle, K., Corbisier, Y. E., Sawelew, L., & Malavasi, S. (2013). Sound production mechanism in *Gobius paganellus* (Gobiidae). *The Journal of Experimental Biology*, 216(17), 3189-3199.
- Partan, S. R., & Marler, P. (2005). Issues in the classification of multimodal communication signals. *The American Naturalist*, 166(2), 231-245.
- Polgar, G., Malavasi, S., Cipolato, G., Georgalas, V., Clack, J. A., & Torricelli, P. (2011). Acoustic communication at the water's edge: evolutionary insights from a mudskipper. *PloS one*, 6(6).
- Popper, A. N., & Fay, R. R. (1973). Sound detection and processing by teleost fishes: a critical review. *The Journal of the Acoustical Society of America*, 53(6), 1515-1529.
- Popper, A. N., & Fay, R. R. (2011). Rethinking sound detection by fishes. *Hearing Research*, 273(1), 25-36.
- Rogers, P. H. & Cox, M. (1988). Underwater sound as a biological stimulus. In *Sensory Biology of Aquatic Animals* (Atema, J., Fay, R. R., Popper, A. N. & Tavolga, W. N., eds), pp. 131-149. New York, NY: Springer-Verlag.
- Rollo, A., Andraso, G., Janssen, J., & Higgs, D. (2007). Attraction and localization of round goby (*Neogobius melanostomus*) to conspecific calls. *Behaviour*, 1-21.

- Rollo, A., & Higgs, D. (2008). Differential acoustic response specificity and directionality in the round goby, *Neogobius melanostomus*. *Animal Behaviour*, 75(6), 1903-1912.
- Rowe, S., & Hutchings, J. A. (2003). Mating systems and the conservation of commercially exploited marine fish. *Trends in Ecology & Evolution*, 18(11), 567-572.
- Saunders, K. M., Brockmann, H. J., Watson, W. H., & Jury, S. H. (2010). Male horseshoe crabs *Limulus polyphemus* use multiple sensory cues to locate mates. *Current Zoology*, 56(5).
- Schuijf, A. (1975). Directional hearing of cod (*Gadus morhua*) under approximate free field conditions. *Journal of comparative physiology*, 98(4), 307-332.
- Smith, R. J. F. (1992). Alarm signals in fishes. *Reviews in Fish Biology and Fisheries*, 2(1), 33-63.
- Smith, A. R., & van Staaden, M. J. (2009). The association of visual and acoustic courtship behaviors in African cichlid fishes. *Marine and Freshwater Behaviour and Physiology*, 42(3), 211-216.
- Speares, P., & Johnston, C. (2011). Sound production in *Etheostoma oophylax* (Percidae) and call characteristics correlated to body size. *Environmental biology of fishes*, 92(4), 461-468.
- Stadler, J. H. (2002). Evidence for a hydrodynamic mechanism of sound production by courting males of the notchtongue goby, *Bathygobius curacao* (Metzelaar). *Bioacoustics*, 13(2), 145-152.
- Takemura A (1984) Acoustical behaviour of the freshwater goby *Odontobutis obscura*. *Bulletin of the Japanese Society of Scientific Fisheries*, 59: 561–564.
- Tavolga, W. N. (1958). The significance of underwater sounds produced by males of the gobiid fish, *Bathygobius soporator*. *Physiological Zoology*, 259-271.
- Thaker, M., Gabor, C. R., & Fries, J. N. (2006). Sensory cues for conspecific associations in aquatic San Marcos salamanders. *Herpetologica*, 62(2), 151-155.
- Van Staaden, M. J., & Smith, A. R. (2011). Cutting the Gordian knot: complex signalling in African cichlids is more than multimodal. *Current Zoology*, 57, 237-252.
- Winn, H. E., Marshall, J. A., & Hazlett, B. (1964). Behavior, diel activities, and stimuli that elicit sound production and reactions to sounds in the longspine squirrelfish. *Copeia*, 413-425.
- Zhang, G., & Takemura, A. (1989). Acoustical behavior of brown goby, *Bathygobius fuscus*. *Bulletin of the Faculty of Fisheries-Nagasaki University (Japan)*.

Chapter II

VARIATION IN THE CALL STRUCTURE OF MALE ROUND GOBIES (*NEOGOBIUS MELANOSTOMUS*) IN RELATION TO BODY MORPHOMETRICS

Introduction

The sounds produced by animals can often be used by listeners and eavesdroppers as a means of determining information about the call producer (reviewed in Mock et al., 2011; Kaplan, 2014). Many terrestrial animals perform acoustic displays to attract potential mates (Peters and Peters, 2010), act as a mode of species recognition (Rivero *et al.* 2000), and advertise their quality (Vannoni and McElligott, 2007; Voituron *et al.*, 2012). Inherent differences in sound production mechanisms can create signals that are variable, both within and between individuals (Ryan and Guerra, 2014) and a great deal of the variability in an acoustic display of an individual is based on the individual's "condition". For example, in chacma baboons (*Papio hamadryas ursinus*), a calling bout will contain signals of varying intensity, where intensity decreases as the individuals exhaust themselves (Kitchen et al., 2003). In cases where these signals are costly to produce, it is expected that females can use the display as a measure of the potential mate's condition (Zahavi, 1975). Honest signals should be those that accurately advertise a desirable trait, whether it is male size and resource holding capabilities (Wells, 1977), lipid reserves to protect against filial cannibalism (Manica, 2004) or possession of desired heritable genes (Bentsen et al., 2006). Acoustic honest signalling has been found in many terrestrial species across a wide range of taxa (birds: Spencer et al., 2003; amphibians: Gingras *et al.*, 2013; mammals: Vannoni and McElligott, 2008, Kitchen *et al.*, 2013) and

is suspected to be in many others (arachnids: Rivero et al., 2000; tortoises: Galeotti *et al.*, 2005).

In fish, the role of acoustic signalling as an honest signal is much less clear. It is often difficult to break down acoustic cues to determine the information being carried with them and what may be true for some groups does not necessarily apply to others. Dominant frequency, an acoustic feature that is often negatively correlated with male size in some species (e.g. Myrberg et al. 1986; McKibben and Bass, 1998; Amorim et al. 2003; De Jong *et al.*, 2007), appears to have no relationship to male size in others (Boyle and Tricas, 2011). In the painted goby (*Pomatoschistus pictus*), acoustic activity is related to lipid stores, with fatter males displaying more and with a greater sound pressure level (Amorim *et al.*, 2013). Female midshipman (*Porichthys notatus*) prefer calling bouts that are longer and uninterrupted (McKibben and Bass, 1998) and both sound amplitude and calling activity are indicators of condition in sand goby, *Pomatoschistus minutus* (Pedroso et al. 2013). There is also limited evidence that acoustic parameters influence mate choice in some fish species (Myrberg et al. 1986; Rowe & Hutchings 2008), suggesting that sound can be a true honest signal in fish, but the data remain sparse and more investigation is needed.

The purpose of the current study was to determine if the sounds produced by male round goby (*Neogobius melanostomus*) have the potential to act as size indicators. Previous work shows that round goby males vocalize and females are able to localize to calling males (Rollo et al., 2007), but whether or not these calls also function as honest signals is not yet known. I examined the correlations between male body characteristics and call variability in several call components of the round goby as a means of determining the potential use of calls as a measure of male condition. I test the prediction

that acoustic signals will correlate to at least one measure of body size, thus creating a link to body condition. Determining a linkage between calls and body condition is the first step in determining if acoustic honest signalling has the potential to exist in this species and to explain observed variability in call characteristics.

Methods

Model Organism

The round goby is a benthic fish, originally from the Ponto-Caspian region that has established populations throughout the Great Lakes (Jude et al., 1992). This fish is a brood guarder (Meunier et al. 2009), with the males keeping and defending nests.

Territory holding males, hereafter referred to as parental males, are very readily identified by their nuptial characteristics of black colour, swollen cheeks, a slimy coating and an enlarged urogenital papilla (Marentette et al., 2010). Parental males are known to call to attract females (Rollo et al. 2007; Rollo & Higgs 2008) much like other species of gobies (Lugli et al. 1996). The acoustic component of courtship is accompanied by visual and olfactory cues, whereby the male will fan with his pectoral fins (Meunier et al., 2013). It has been hypothesized (Marentette et al., 2009) that these parental males must defend their nests from sneaker males, reproductive males that do not display nuptial characteristics.

Sampling Methods

Round gobies were caught by angling at the Erieau Marina (N 42.2572945 W 81.9084025), on the north shore of Lake Erie and in the Detroit River, Ontario, in the summers of 2013 and 2014. Fish were then transported to the Animal Care Facility at the

University of Windsor where they were kept on a 12L:12D light cycle at $18\pm 1^{\circ}\text{C}$ and fed daily (Hikari Cichlid Gold; Kyorin Corporation, Japan). Males were visually and acoustically isolated from each other in a Z-Hab System (Aquatic Habitats Incorporated; Apopka, Florida, USA) and kept for no longer than one month. A length of PVC tube was supplied in each tank to act as a shelter. Females were kept in communal tanks of up to ten females.

Experimental Procedure

Only reproductive round gobies were used in this study. Male reproductivity was assessed by the enlargement of the urogenital papilla (Marentette et al., 2010). Both parental and “sneaker” males may have been used as parental males lost their nuptial colouration after transport and could not be differentiated by size. Reproductive females were identified by the enlargement and colour of the urogenital papilla; the papilla appears round and swollen in reproductive females, with the colour shifting from white to yellow-orange (Marentette et al., 2010).

Fish were tested at night as round gobies are more vocal between the hours of dawn and dusk than during daylight hours (Higgs and Humphrey, 2014). During testing all filters, lights and air pumps were turned off to reduce abiotic noise and a plastic, grid barrier was placed into the tank with the male. The PVC shelter was removed to remove any chance of noise being produced by banging against the shelter and a hydrophone (RESON TC4013-4, $-210.9\text{ dB re } 1\mu\text{Pa/V}$ at 1m) was inserted into the male’s side of the tank. Recordings were made using a solid state recorder (Marantz PMD670) after running through a preamplifier (RESON VP2000 Voltage Preamplifier EC6081) with a high pass setting of 1 Hz and a low pass setting of 15 kHz. The hydrophone remained in close

proximity to the male throughout the trial. After five minutes of acclimation, the recording was started for a 30 minute control recording and then a reproductive female was added to the unoccupied side of the tank. The male could see and smell the female, but could not access her. After each half hour trial, the female was placed back into the communal female tank to be used for other trials.

After the experiment, males were euthanized using clove oil (>20 ppm after anaesthesia with 10 ppm) and measured for head width above the eyes (HW), total length (TL), total wet weight (TW) and gonad weight (GW). Total weight and gonad weight were used to calculate gonadosomatic index (GSI) using the equation

$$GSI = \left(\frac{\text{gonad mass}}{\text{total body mass}} \right) \times 100\%.$$

Call Analysis

Male calls were analyzed using Adobe Audition® 3.0 software (Adobe Systems Incorporated, USA). Sound files were cleaned of background noise by using the software's Noise Reduction feature and each call was measured for the dominant frequency (DF, characterized as the frequency with the greatest energy determined by the power spectra, FFT size 32768, Blackmann-Harris window), number of pulses, total duration of the call and the interpulse interval (IPI). Interpulse interval was defined as the time in between two peaks of consecutive pulses when viewed using the waveform setting of Adobe Audition® 3.0.

Statistical Analysis

Data was Log_{10} transformed to obtain a normal distribution. Call and body characteristics were associated against each other using principal components analysis (PCA) on JMP 10 software (SAS Institute Incorporated, North Carolina, USA) to determine general correlative relationships between body quality and call characteristics. Redundant and non-correlative variables identified by the PCA were removed from all subsequent tests. Generalized Linear Models (GLMs) were performed using call characteristics of IPI, DF and duration as dependent factors with body morphometrics of total length, head width and total weight acting as explanatory variables. For these analyses, only calls of more than one pulse were included. In cases where a three-way or two-way interaction between these body characteristics was non-significant, the interaction term was removed and the model run again. Effects of interaction terms on call characteristics were examined using procedures provided by Dr. Jeremy Dawson (<http://www.jeremydawson.co.uk/slopes.htm>). The residuals of all of the GLMs were normally distributed.

The variation of the calls' dominant frequency component was analyzed by examining the coefficients of variation between and within male calls. Coefficient of Variation (CV) is defined with the function $= \frac{SD}{\bar{x}} \times 100\%$. For this purpose, only the first pulse of each call was examined and any males with at least 3 calls were included in the within male variability (CV_w) analysis. As in Bee et al. (2001), between-male variability (CV_b) uses the mean of all of the first and single pulses across all males recorded. Variation is considered greater between males than among males when the ratio of $CV_b/CV_w > 1$.

Results

A total of 36 males were tested, of which 24 produced calls, one of which only called during the control trial. Sixteen actively calling males produced at least one call of more than one grunt, with two of these males producing more than a single call of one pulse. The total number of calls was 37 with 20 of these being calls of more than one pulse. Only 4 males produced at least three calls during a single recording, these calls were analysed for CV_w , with 3 of the four having much lower variation in frequency within than between males (Table 2.1).

The principal components analysis identified three main components (Table 2.2), of which, the first principal component accounted for nearly 41% of the variability in the model. Three body characteristics, total length, head width and total weight were strongly associated with scores of PC1, as was a single call characteristic, interpulse interval. The second component accounted for 26% of the variability. Call characteristics of frequency, duration and number of pulses were all strongly associated with this component. Total width and total size were weakly positively associated with this component. Gonadosomatic index is most strongly associated with scores of PC4 and is not tightly correlated with any feature of the call or other body measure so GSI was discounted from further analyses.

Generalized Linear Models of call characteristics against body morphometrics identify several significant interactions between body features affecting the call. Frequency is affected by two two-way interactions of HWxTW and HWxTL. The HWxTW cross showed heavier males having lower frequency calls than their smaller counterparts (X^2 ; $df=1$, $N = 20$ $p = 0.0293$; Figure 2.1a). On the other hand, the HWxTL cross showed longer males had higher frequencies than smaller males (X^2 ; $df=1$, $N = 20$

Table 2.1

Table 2.1. Inter- and intra-male variability of call frequency. The coefficients of variation in dominant frequency for four male round goby that each called at least 3 times compared to the variation in the calls of all recorded males. Only the dominant frequency of the first pulse of each call was used in the analysis. CV_w within male variability, CV_b between male variability.

Male	CV_w	CV_b	CV_b / CV_w
5	13.6	36.0	2.7
31	17.3		2.1
32	7.0		5.1
36	50.0		0.7

Table 2.2

Table 2.2. Loadings of call and body measures. Principal components loadings

showing interpulse interval, total length, head width and total weight were most highly associated with the first component, frequency, duration and number of pulses grouped in the second component and GSI most strongly in the fourth component by itself.

Characteristic	PC1	PC2	PC3	PC4	PC5	PC6	PC7	PC8
Log#pulses	0.191	0.783	0.524	0.152	-0.091	-0.158	0.131	0.001
LogDuration	0.471	0.772	0.280	0.224	0.019	0.170	-0.151	-0.002
LogFreq	-0.180	-0.572	0.506	0.554	0.275	-0.005	0.007	0.001
LogTW	0.916	-0.378	0.082	0.045	-0.058	0.023	0.054	-0.020
logGSI	-0.450	0.102	-0.498	0.702	-0.198	0.056	0.044	-0.001
logTL	0.820	-0.464	0.214	0.020	-0.209	0.137	0.049	0.014
logHW	0.877	-0.125	-0.297	0.235	-0.016	-0.238	-0.117	0.004
logIPI	0.700	0.380	-0.508	0.022	0.296	0.070	0.113	0.004
Total variability	40.839	25.847	15.738	11.660	3.243	1.733	0.932	0.009

Figure 2.1

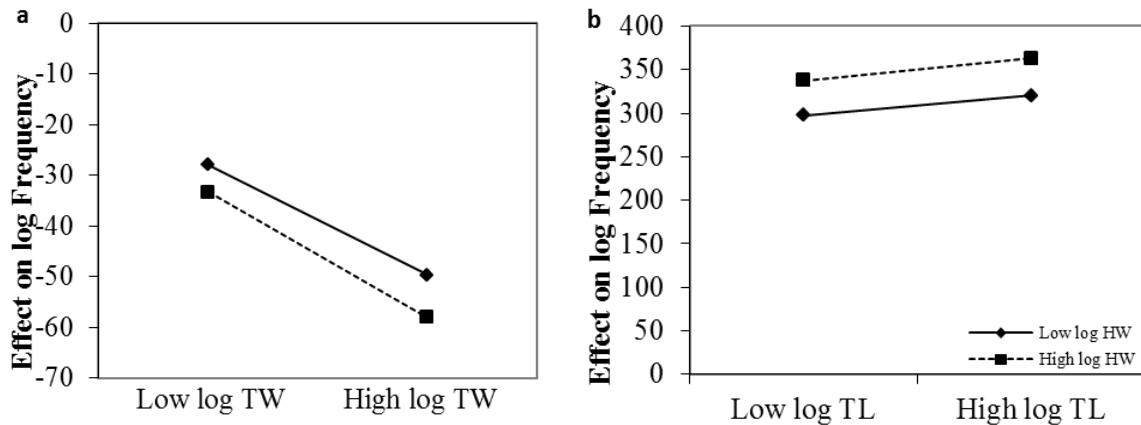


Figure 2.1. The relationship between HWxTW, HWxTL and DF. a.) The negative relationship between the log₁₀ transformed dominant frequency of calls and the two-way interaction effect of head width and total weight. This interaction suggests larger males have lower dominant frequencies of their calls. b.) The positive relationship between the log₁₀ transformed dominant frequency of calls and the two-way interaction effect of head width and total length. This interaction suggests larger males have higher dominant frequencies of their calls.

$p = 0.0213$; Figure 2.1b. These interactions work in opposite directions with large headed, heavier males having lower frequency calls, but large headed, long males having higher frequency calls.

The IPI is also affected by two two-way interactions of HWxTW and HWxTL. The HWxTW cross showed heavier, large-headed males having longer IPIs compared to thinner, small-headed males (X^2 ; $df=1$, $N = 20$ $p = 0.045$; Figure 2.2a). This interaction is contradicted by the HWxTL cross, which shows shorter, small-headed males having longer IPIs (X^2 ; $df=1$, $N = 20$ $p = 0.0383$; Figure 2.2b).

The GLM indicated that duration was significantly influenced by a three way interaction of HWxTLxTW X^2 ($df=1$, $N = 20$ $p = 0.0005$; Figure 2.3). This interaction suggests that long duration calls are produced by short, heavy males and short duration calls are produced by thin, long males. In this case, males with a greater TW did not change duration as considerably as males with a lesser TW. Males producing the shortest duration calls were those with long, thin bodies and a large head. Long, thin males with a small head produced calls with durations comparable to short, thin males with large heads.

Discussion

Honest signalling requires that a signal accurately and consistently displays the trait of interest (Zahavi, 1975; Johnstone, 1995). In the current study, the accuracy of call features predictive ability for body measures is quite high, suggesting that these acoustic signals could be used by conspecifics to determine the “quality” of the caller. There is also a greater inter-male variability in calls than intra-male variability,

Figure 2.2

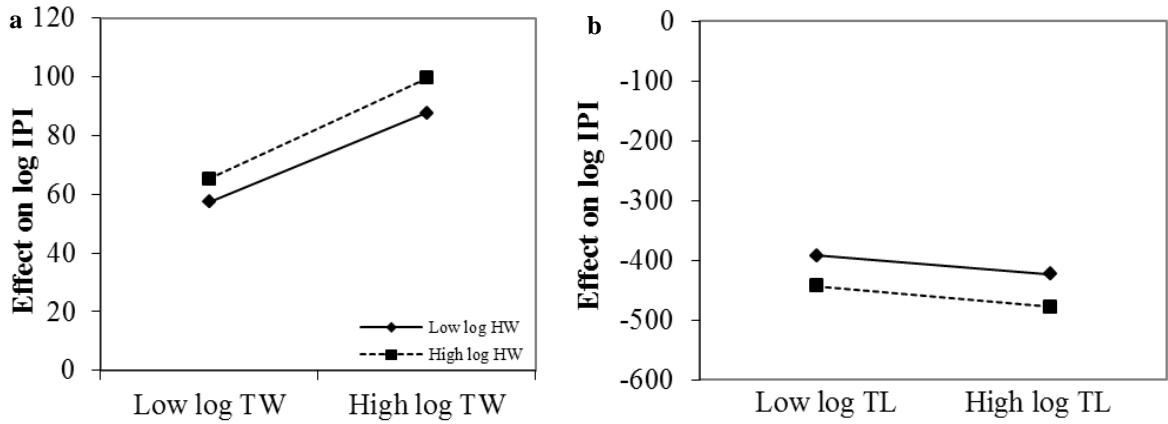


Figure 2.2. The relationship between HWxTW, HWxTL and IPI. a.) The positive relationship between the log₁₀ transformed interpulse interval of calls and the two-way interaction effect of head width and total weight. This interaction suggests larger males have greater interpulse intervals in their calls. b.) The negative relationship between the log₁₀ transformed interpulse interval of calls and the two-way interaction effect of head width and total length. This interaction suggests larger males have shorter interpulse intervals in their calls.

Figure 2.3

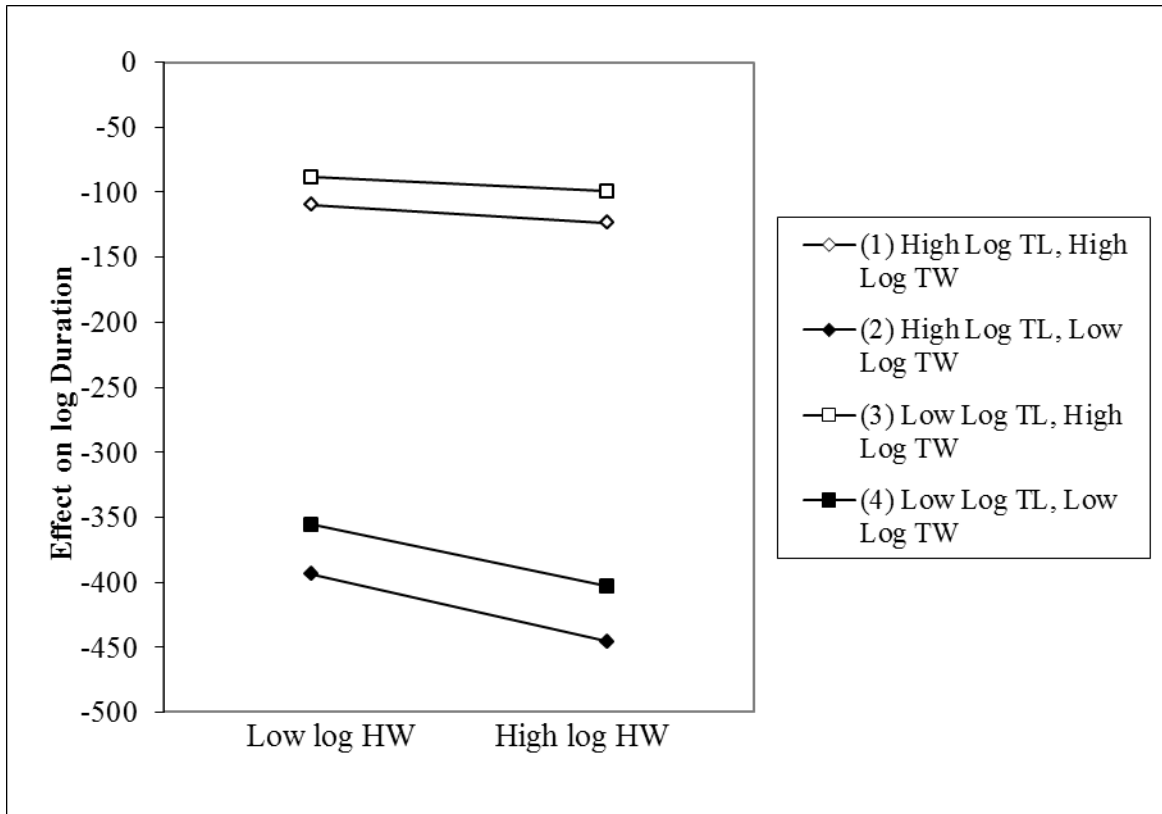


Figure 2.3. The relationship between HWxTLxTW and duration. The negative relationship between the log10 transformed duration of calls and the three-way interaction effect of head width, total length and total weight. Males with a greater weight show a considerably longer call duration regardless of head size and length. The shortest call durations are produced by long, thin males with large heads. Call produced by short, thin, big-headed males and long, thin, small-headed males are of a comparable length.

another factor necessary for honest signalling to exist. The calls produced by individual males were consistent over the short time frame of this study, showing a high reliability in signalling body condition. While the mechanism for sound production is not known in this species, calling may not be overtly costly (Amorim et al 2013) as honest signals are usually thought to be (Gintis et al.,2001; Számadó, 2011), however, the physical mechanism of sound production is likely limited by the body traits measured. In species, such as the weakfish (*Cynoscion regalis*), which use sonic muscles, larger fish produce sounds of lower frequencies due to the increased time it takes for the longer muscle fibres to contract and relax. Similarly, these larger fish produce sounds of greater intensity because they have greater muscle mass (Connaughton et al., 2002). A size-restricted sound production mechanism can be assumed to be the case where head size and body length of the fish affect characteristics of the call, as was seen in the current study. In other species of gobies where the mechanism of sound production has been examined, the call invariably originates from the head region of the fish. Some suspected modes of call production in gobies include projection of water through the operculum (Tavolga, 1958), grinding pharyngeal teeth together (Takemura, 1984; Lugli, 1995) and vibration of the levator pectoralis muscle which originates on the skull and inserts on the pectoral girdle (Parmentier et al., 2013). Should one of these modes of call production prove to be true for the round goby, the physical properties of the fish's head would alter the call, consistent with my results.

In the current study, long males with big heads produced short, high frequency calls with short IPI, while heavy, big headed males produce calls with lower frequencies and longer IPI. Depending on female preference for male traits, the female may be able to tease apart the various levels of a call to determine which nests to visit. In other species of

gobies, there is very little evidence supporting female preference for male size (Magnhagen and Kvarnemo, 1989; Forsgren, 1997; Amorim et al., 2013). Conversely, larger males are more capable fighters and tend to win larger territories, for which females do display a preference (Magnhagen and Kvarnemo, 1989). This contradiction may in part reflect the effort put into courtship. In Pacific blue-eye fish (*Pseudomogil signifier*) females do not mate more frequently with larger, more competitive males, instead choosing males that put greater effort into courtship (Wong, 2004). Bicolor damselfish (*Stegastes partitus*) females are less likely to mate with males that are not actively courting, perhaps because these males have lower energy reserves (Knapp and Kovach, 1991). Female gobiids are thought to prefer males with greater lipid stores (Amorim et al., 2013) and in the current study total weight was a significant predictor of IPI, DF and call duration. Choosing males with greater lipid stores would be adaptive because fattier males may be less likely to cannibalize their eggs (Neff, 2003), larger testes allowing for greater sperm production (Taborsky, 1998), and larger males may have greater brood survival rates (Sabat, 1994).

Call duration provides some counterintuitive results, wherein the longest call durations were performed by short, heavy males with small heads. There is not as great a difference between the groups of heavy males with HW and TL varying as there is between the groups of light males. This could suggest that males must have attained a minimum weight to be able to call for any substantial length of time, perhaps resorting to alternate, less costly modes of communication. While frequency and IPI might be related to morphological restrictions of the sound production mechanism, duration of the call may be more heavily influenced by the male's condition. Plainfin midshipman males that are in better condition and have greater mass, also have a greater number of young at the

end of the season (Sisneros et al., 2009). These males likely use their greater energy reserves for increased courtship effort, to produce calls of longer duration, to which females are more attracted (McKibben and Bass, 1998). Greater fat stores would allow the male to increase courtship, gain more mating opportunities and receive more eggs. Male gobies that care for more eggs exhibit lower incidence of filial cannibalism and more fanning behaviour than those that care for fewer eggs (Forsgren et al., 1996).

For both IPI and dominant frequency I saw opposite relationships for total length or total weight. Longer fish had a shorter IPI and higher frequency and heavier fish had a longer IPI and lower frequency than their counterparts. This disparity may be explained by the different information portrayed to the female by these two metrics of size. Total body weight is a more immediate measure of condition in fish (Cargnelli and Gross, 1997), with immediacy due to ability to thrive under current conditions while total length may be a sum total of growth capability over longer time frames (Nate and Bremigan, 2005). A heavier fish will be more capable of putting in the reproductive effort in the current season rather than waiting until it is larger, but possibly in worse condition. While I do not know which metric is more preferred by female round gobies, it is possible that the two metrics convey different information, or conversely that the call characteristics are influenced more by weight than length. This possibility seems unlikely however, as these two characteristics are very tightly correlated in the reproductive males used in this study (Pearson's $r = 0.959$). Until we can ascertain how calls are produced in round goby however I cannot assess what is driving these differences.

Although acoustic recordings are greatly affected by small tank dynamics (Akamatsu et al. 2002), the recordings in the current study were taken with the hydrophone very close to the fish and all experiments were done in the same conditions

so individual comparisons would not be affected. The removal of the male's shelter for the duration of the recording may also have negatively impacted call production. This does not seem to be the case however, as male call rate is similar to that found in nature (unpublished data), as are the call characteristics (Rollo et al., 2007). There also remains the possibility that male mate choice affected the results of this study. However, females were size matched and always checked for status of the urogenital papilla before being placed in the experimental setup. Regardless, male courtship effort may have been altered by the use of several females across trials. The production of calls by males thought to have been "sneakers" may suggest that this reproductive stage could be an alternate behavioural tactic when more dominant males are present.

In conclusion, male round goby have call parameters that are strongly linked to several body characteristics. These acoustic traits may act as an honest signal to females during courtship of the male's condition or parental capabilities. Male courtship calls may carry a large amount of information not only of male lipid stores, but also of overall size. These traits, if the female is able to detect them, would be beneficial for determining risks of filial cannibalism, effort put into rearing, as well as fighting capabilities. Future studies should aim to determine the relative importance of acoustic cues in relation to other male courtship signalling modalities and test the response of females to these courtship signals as a true test of honest signalling in this species.

References

- Akamatsu, T., Okumura, T., Novarini, N., & Yan, H. Y. (2002). Empirical refinements applicable to the recording of fish sounds in small tanks. *The Journal of the Acoustical Society of America*, *112*(6), 3073-3082.
- Amorim, M. C. P., Pedroso, S. S., Bolgan, M., Jordao, J. M., Caiano, M., & Fonseca, P. J. (2013). Painted gobies sing their quality out loud: acoustic rather than visual signals advertise male quality and contribute to mating success. *Functional Ecology*, *27*(2), 289-298.
- Bee, M. A., Kozich, C. E., Blackwell, K. J., & Gerhardt, H. C. (2001). Individual variation in advertisement calls of territorial male green frogs, *Rana clamitans*: implications for individual discrimination. *Ethology*, *107*(1), 65-84.
- Bentsen, C. L., Hunt, J., Jennions, M. D., & Brooks, R. (2006). Complex multivariate sexual selection on male acoustic signalling in a wild population of *Teleogryllus commodus*. *The American Naturalist*, *167*(4), E102-E116.
- Boyle, K. S., & Tricas, T. C. (2011). Sound production in the longnose butterflyfishes (genus *Forcipiger*): cranial kinematics, muscle activity and honest signals. *The Journal of experimental biology*, *214*(22), 3829-3842.
- Cargnelli, L. M., & Gross, M. R. (1997). Notes: Fish energetics: Larger individuals emerge from winter in better condition. *Transactions of the American Fisheries Society*, *126*(1), 153-156.
- De Jong, K., Bouton, N., & Slabbekoorn, H. (2007). Azorean rock-pool blennies produce size-dependent calls in a courtship context. *Animal Behaviour*, *74*(5), 1285-1292.
- Forsgren, E., Karlsson, A., & Kvarnemo, C. (1996). Female sand gobies gain direct benefits by choosing males with eggs in their nests. *Behavioral Ecology and Sociobiology*, *39*(2), 91-96.
- Forsgren, E. (1997). Female sand gobies prefer good fathers over dominant males. *Proceedings of the Royal Society of London. Series B: Biological Sciences*, *264*(1386), 1283-1286.
- Galeotti, P., Sacchi, R., Fasola, M., Rosa, D. P., Marchesi, M., & Ballasina, D. (2005). Courtship displays and mounting calls are honest, condition-dependent signals that influence mounting success in Hermann's tortoises. *Canadian journal of zoology*, *83*(10), 1306-1313.
- Gingras, B., Boeckle, M., Herbst, C. T., & Fitch, W. T. (2013). Call acoustics reflect body size across four clades of anurans. *Journal of Zoology*, *289*(2), 143-150.
- Gintis, H., Smith, E. A., & Bowles, S. (2001). Costly signalling and cooperation. *Journal of theoretical biology*, *213*(1), 103-119.
- Green, B. S., & McCormick, M. I. (2005). O₂ replenishment to fish nests: males adjust brood care to ambient conditions and brood development. *Behavioral Ecology*, *16*(2), 389-397.
- Higgs and Humphrey (August 2014) *Passive monitoring of round goby acoustic ecology and the influence of anthropogenic noise*. Presented at the International Congress on the Biology of Fish, Edinburgh, Scotland.
- Johnstone, R. A. (1995). Honest advertisement of multiple qualities using multiple signals. *Journal of theoretical Biology*, *177*(1), 87-94.

- Jude, D. J., Reider, R. H., & Smith, G. R. (1992). Establishment of Gobiidae in the Great Lakes basin. *Canadian Journal of Fisheries and Aquatic Sciences*, 49(2), 416-421.
- Kaplan, G. (2014). Animal communication. *Wiley Interdisciplinary Reviews: Cognitive Science*, 5(6), 661-677.
- Kitchen, D. M., Seyfarth, R. M., Fischer, J., & Cheney, D. L. (2003). Loud calls as indicators of dominance in male baboons (*Papio cynocephalus ursinus*). *Behavioral ecology and sociobiology*, 53(6), 374-384.
- Kitchen, D. M., Cheney, D. L., Engh, A. L., Fischer, J., Moscovice, L. R., & Seyfarth, R. M. (2013). Male baboon responses to experimental manipulations of loud “wahoo calls”: testing an honest signal of fighting ability. *Behavioral Ecology and Sociobiology*, 67(11), 1825-1835.
- Knapp, R. A., & Kovach, J. T. (1991). Courtship as an honest indicator of male parental quality in the bicolor damselfish, *Stegastes partitus*. *Behavioral Ecology*, 2(4), 295-300.
- Lugli, M., Pavan, G., Torricelli, P., & Bobbio, L. (1995). Spawning vocalizations in male freshwater gobiids (Pisces, Gobiidae). *Environmental Biology of Fishes*, 43(3), 219-231.
- Lugli, M., Pavan, G., & Torricelli, P. (1996). The importance of breeding vocalizations for mate attraction in a freshwater goby with a composite sound repertoire. *Ethology Ecology & Evolution*, 8(4), 343-351.
- Magnhagen, C., & Kvarnemo, L. (1989). Big is better: the importance of size for reproductive success in male *Pomatoschistus minutus* (Pallas)(Pisces, Gobiidae). *Journal of Fish Biology*, 35(6), 755-763.
- Manica, A. (2004). Parental fish change their cannibalistic behaviour in response to the cost-to-benefit ratio of parental care. *Animal Behaviour*, 67(6), 1015-1021.
- Marentette, J. R., Fitzpatrick, J. L., Berger, R. G., & Balshine, S. (2009). Multiple male reproductive morphs in the invasive round goby (*Apollonia melanostoma*). *Journal of Great Lakes Research*, 35(2), 302-308.
- Marentette, J. R., Gooderham, K. L., McMaster, M. E., Ng, T., Parrott, J. L., Wilson, J. Y., Wood, C. M., & Balshine, S. (2010). Signatures of contamination in invasive round gobies (*Neogobius melanostomus*) A double strike for ecosystem health?. *Ecotoxicology and Environmental Safety*, 73(7), 1755-1764.
- McKibben, J. R., & Bass, A. H. (1998). Behavioral assessment of acoustic parameters relevant to signal recognition and preference in a vocal fish. *The Journal of the Acoustical Society of America*, 104, 3520-3533.
- Meunier, B., Yavno, S., Ahmed, S., & Corkum, L. D. (2009). First documentation of spawning and nest guarding in the laboratory by the invasive fish, the round goby (*Neogobius melanostomus*). *Journal of Great Lakes Research*, 35(4), 608-612.
- Meunier, B., White, B., & Corkum, L. D. (2013). The role of fanning behavior in water exchange by a nest-guarding benthic fish before spawning. *Limnology and Oceanography: Fluids and Environments*, 3(1), 198-209.
- Mock, D. W., Dugas, M. B., & Strickler, S. A. (2011). Honest begging: expanding from signal of need. *Behavioral Ecology*, 22, 909-917.
- Nate, N. A., & Bremigan, M. T. (2005). Comparison of mean length at age and growth parameters of bluegills, largemouth bass, and yellow perch from length-stratified subsamples and samples in Michigan Lakes. *North American Journal of Fisheries Management*, 25(4), 1486-1492.

- Neff, B. D. (2003). Paternity and condition affect cannibalistic behavior in nest-tending bluegill sunfish. *Behavioral Ecology and Sociobiology*, 54(4), 377-384.
- Parmentier, E., Kéver, L., Boyle, K., Corbisier, Y. E., Sawelew, L., & Malavasi, S. (2013). Sound production mechanism in *Gobius paganellus* (Gobiidae). *The Journal of Experimental Biology*, 216(17), 3189-3199.
- Partan, S. R., & Marler, P. (2005). Issues in the classification of multimodal communication signals. *The American Naturalist*, 166(2), 231-245.
- Pedroso, S. S., Barber, I., Svensson, O., Fonseca, P. J., & Amorim, M. C. P. (2013). Courtship sounds advertise species identity and male quality in sympatric *Pomatoschistus* spp. Gobies. *PLoS ONE* 8(6)
- Peters, G., & Peters, M. K. (2010). Long-distance call evolution in the Felidae: effects of body weight, habitat, and phylogeny. *Biological Journal of the Linnean Society*, 101(2), 487-500.
- Rivero, A., Alatalo, R. V., Kotiaho, J. S., Mappes, J., & Parri, S. (2000). Acoustic signalling in a wolf spider: can signal characteristics predict male quality?. *Animal Behaviour*, 60(2), 187-194.
- Rollo, A., Andraso, G., Janssen, J., & Higgs, D. (2007). Attraction and localization of round goby (*Neogobius melanostomus*) to conspecific calls. *Behaviour*, 144(1), 1-21.
- Rollo, A., & Higgs, D. (2008). Differential acoustic response specificity and directionality in the round goby, *Neogobius melanostomus*. *Animal Behaviour*, 75(6), 1903-1912.
- Ryan, M. J., & Guerra, M. A. (2014). The mechanism of sound production in túngara frogs and its role in sexual selection and speciation. *Current opinion in neurobiology*, 28, 54-59.
- Sabat, A. M. (1994). Costs and benefits of parental effort in a brood-guarding fish (*Ambloplites rupestris*, Centrarchidae). *Behavioral Ecology*, 5(2), 195-201.
- Sisneros, J. A., Alderks, P. W., Leon, K., & Sniffen, B. (2009). Morphometric changes associated with the reproductive cycle and behaviour of the intertidal-nesting, male plainfin midshipman *Porichthys notatus*. *Journal of fish biology*, 74(1), 18-36.
- Spencer, K. A., Buchanan, K. L., Goldsmith, A. R., & Catchpole, C. K. (2003). Song as an honest signal of developmental stress in the zebra finch (*Taeniopygia guttata*). *Hormones and Behavior*, 44(2), 132-139.
- Számádó, S. (2011). The cost of honesty and the fallacy of the handicap principle. *Animal Behaviour*, 81(1), 3-10.
- Taborsky, M. (1998). Sperm competition in fish: bourgeois' males and parasitic spawning. *Trends in Ecology & Evolution*, 13(6), 222-227.
- Takemura A (1984) Acoustical behaviour of the freshwater goby *Odontobutis obscura*. *Bulletin of the Japanese Society of Scientific Fisheries*, 59: 561–564.
- Tavolga, W. N. (1958). The significance of underwater sounds produced by males of the gobiid fish, *Bathygobius soporator*. *Physiological Zoology*, 259-271.
- Vannoni, E., & McElligott, A. G. (2008). Low frequency groans indicate larger and more dominant fallow deer (*Dama dama*) males. *PloS one*, 3(9).
- Voituron, Y., Brepson, L., Richardson, C., Joly, P., & Lengagne, T. (2012). Energetics of calling in the male treefrog *Hyla arborea*: when being large means being sexy at low cost. *Behaviour*, 149, 775-793.

- Wells, K. D. (1977). Territoriality and male mating success in the green frog (*Rana clamitans*). *Ecology*, 750-762.
- Wong, B. B. (2004). Superior fighters make mediocre fathers in the Pacific blue-eye fish. *Animal Behaviour*, 67(3), 583-590.
- Zahavi, A. (1975). Mate selection—a selection for a handicap. *Journal of theoretical Biology*, 53(1), 205-214.

Chapter III

EXAMINING THE POTENTIAL FOR ACOUSTIC HONEST SIGNALLING IN THE ROUND GOBY (*NEOGOBIUS MELANOSTOMUS*) THROUGH FEMALE CHOICE

Introduction

Acoustically-mediated honest signalling is dependent upon two factors: the first being that the acoustic signal correlates with male condition and the second being that females can discern this relationship through the sound. Both of these aspects are well studied in terrestrial systems. Male condition is usually examined as a function of male size (Schulte-Hostedde et al., 2005), though some studies are beginning to look into condition as a factor of social ranking (Kitchen et al., 2003, Vannoni and McElligott, 2008). Larger males are often more capable combatants, winning themselves a greater social rank (Le Boeuf , 1974; Bowyer, 1986). From the aspect of size, sound production should be a reliable signal as sound tends to scale linearly with the size of the larynx or syrinx and diameter of the mouth, or in the case of insects, the size of the vibrating organ producing the sound (Fletcher, 2004). When presented with acoustic signals produced by conspecific males, females are often able to judge male size accurately, often by the frequency of the call (Ryan et al, 1990). Female crickets (*Teleogryllus commodus*) prefer males that present greater calling effort, a trait that is condition dependent and suspected of being heritable (Bentson et al., 2006). Having offspring that in turn produce more attractive calls fits into the “sexy son” hypothesis, increasing overall fitness as the female would ultimately have more descendants (Weatherhead and Robertson, 1979). Males in better condition put greater effort into raising young in species that do so (Iwasa and Pomiankowski, 1999).

In fish, evidence for female choice for male acoustic signals is limited. While male acoustic signals often correlate strongly with male body size (Amorim et al., 2010; Chapter 2), few studies examine honest signalling from the female choice perspective. In cases where female preference for male calls is examined, females tend to prefer calls where the temporal aspects of the call correlate with male body characteristics. In the plainfin midshipman (*Porichthys notatus*), male calls that are uninterrupted and longer in duration are preferred by females (McKibben and Bass, 1998). Greater size and muscle mass may be attributed to the longer call duration (Connaughton et al., 2002) and if larger males are preferred, duration may act as an honest signal of size. Greater courtship effort is also linked to greater male energy reserves, reducing the chances that males will cannibalize their brood (Knapp and Kovach, 1991), thus increasing female fitness.

The current study focuses on the potential for honest signalling in the round goby (*Neogobius melanostomus*). Previous work has shown that acoustic characteristics do correlate strongly with male body conditions (Chapter 2). Where male calls function as indicators of body size, the potential exists for females to exploit this trait. The preference for larger males and the ability to tell a male's "condition" from his call would make the acoustic signal a true honest signal. The objective of this chapter is to determine if female round goby show some level of preference for male call characteristics, providing evidence for true honest signalling of male size in this species.

Methods

Sampling Methods

Round gobies were caught by angling at the Eriean Marina (N 42.2572945 W 81.9084025), on the north shore of Lake Erie and in the Detroit River, Ontario in the

summers of 2013 and 2014. Fish were then transported to the Animal Care Facility at the University of Windsor where they were kept on a 12L:12D light cycle at $18\pm 1^{\circ}\text{C}$ and fed daily (Tetramin Fish Flakes; Tetramin Inc, Blacksburg, VA, USA). Fish were housed in communal tanks of up to 10 females. Lengths of PVC tubes and clay pots were provided as artificial shelters.

Effects of Hormone Injection

Once recordings of reproductive males had concluded, out-of-season female round gobies were tested for male call preference. Initial tests were run at night during the months of October and November, 2013. Previously reproductive females were given an intraperitoneal injection of $0.5\ \mu\text{L/g}$ fish body mass of Ovaprim (Syndel Laboratories Ltd., Vancouver, B.C.) 5 days before testing, and then again after testing. The second test was done 48 hours after the second injection. This timing was decided as the fish were no longer in reproductive condition, thus the longer timeline for injection with a booster was chosen. This method has been used previously for inducing maturation in juvenile fish (Hansen and Routledge). Two trials were done in order to see if female choice was consistent across time within the same female. A total of 10 fish were used for this test. Hormone injection was decided upon in order to enhance female response rates, since the fish used in this time frame were no longer in reproductive condition. The experimental procedure for the hormone injection study was the same procedure used in subsequent tests. After the second trial females were euthanized with clove oil and measured for total length (mm), total weight (g) and gonad weight (g). When females were in reproductive condition during the summer of 2014, Ovaprim injection did not occur, and an additional 34 females were tested for call preference.

Male Call Playback

I had recordings from several reproductive male round gobies from previous work (Chapter 2) that I used in female call preference experiments. Both parental and “sneaker” males were used in this portion of the study, as parental male nuptial colouration returned to normal colouration and the two morphs were not distinguishable by size. These calls were recorded in individual tanks in a Z-Hab System (Aquatic Habitats Incorporated; Apopka, Florida, USA), at night during the summer months of 2013 and 2014 while in the presence of a reproductive female. All calls used were produced by males while in the presence of a reproductive female. The longest call train from each male was sectioned out of a 30 minute recording using Adobe Audition 3.0 software (Adobe Systems Incorporated, USA) so that the file used for playback was of a single, clear fish call. Calls were saved with a 15 decibel gain to allow for easy playback.

A 1020-L rectangular ($243.8 \times 91.4 \times 91.4$ cm) fibreglass tank with a holding area in the center and speakers (UW-30, Lubell Labs, Columbus, OH, USA) at either end was used for all female choice trials (Figure 3.1). The barriers had four 10 cm wide exits located approximately 20 cm apart so that fish must navigate through the barrier to gain access to the speaker of choice. Speakers were on acoustic foam to reduce vibration and were surrounded by a nylon mesh barrier so that fish could not use the speaker as a shelter. A GoPro Hero 2 camera (Woodman Labs Inc., USA) was situated above the tank and recorded 960p definition video of each thirty minute trial. The tank was filled with dechlorinated tap water to a depth of 30 cm and was held at 13 ± 1 °C. To start a trial, a reproductive female was placed in the centre of the holding area using a net. In order to reduce the chances that a female would seek shelter and refuse to move when calls began,

Figure 3.1

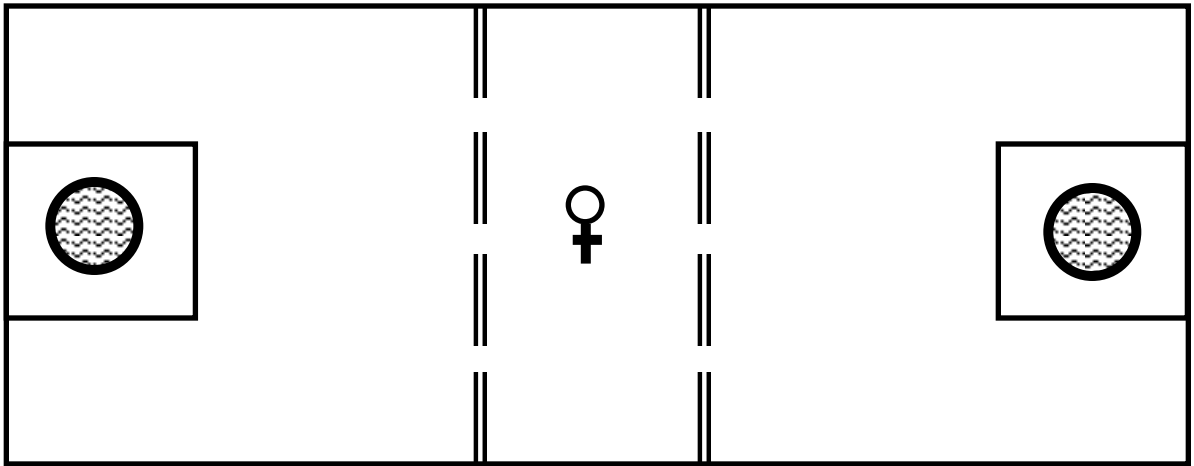


Figure 3.1. Diagram of the female choice experimental set-up. The tank in which female call preference was tested consisted of three “zones”, the two outer zones in which a speaker played a call from and the centre or starting zone. Dashed lines represent the barrier from which females had to navigate out of to reach a speaker. Speakers are represented by circles and the nylon mesh cage around the speakers are represented by the boxes.

there was no acclimation period. Two calls were chosen at random from a total of 12 male calls, and the speakers never played the same call during the same trial. A call was then played from each speaker simultaneously and each at an intensity of 140 dB re 1 μ Pa as measured before each trial by a precalibrated hydrophone (Interocean Inc., San Diego, CA, U.S.A.). Calls were left playing on a continuous loop for the duration of the 30 minute trial. The fish was left without any other outside stimulus for the duration of the trial. After the trial, females were euthanized with clove oil and measured for total length (mm), total weight (g) and gonad weight (g).

Female behavioural responses to male playback were quantified from video to determine speaker of choice for male preference, latency of response before approach began, and time spent near speaker. A responsive female was any female that oriented suddenly to a call and navigated through the barriers to approach one of the speaker zones. Females that oriented towards a call but stopped at the barriers were counted as non-responders as the motivation for moving from the starting zone could not be judged. A female was considered to have chosen a speaker when she was within 10 cm of the cage in which the speaker was placed. "Time spent near a speaker" was determined by recording the period of time during which a female was situated within 10 cm of a speaker cage.

Statistical Analysis

All statistical analyses were performed using JMP 10 statistical software (SAS Institute Incorporated, North Carolina, USA). A matched pairs t-test was performed to compare the mean amount of time spent within each speaker region between trials of hormone injected females that responded in both trials. The responses of injected and

non-injected fish were tested and found not to be significantly different. Therefore, all females were analyzed as a single group.

Data from trial 1 of each responsive fish was \log_{10} transformed to obtain a normal distribution. A Generalized Linear Model (GLM) was run using latency to respond as the dependent variable and the dominant frequency (DF) and interpulse interval (IPI) of the first male chosen as explanatory variables. A separate GLM was run with time spent in the speaker area as the explanatory variable and the dominant frequency and interpulse interval as explanatory variables. The GLM of time spent at the speaker of choice was then repeated with the initially chosen speaker's DF and IPI.

To determine if females exhibited a preference for particular male call characteristics, males' calls were ranked as attractive or unattractive based on the amount of time a female spent in their respective speaker arena. An attractive male was ranked a 1 if that male had a lower frequency than the unattractive male and a 2 if that male had a higher frequency than the unattractive male. For IPI, attractive males were given a rank of 1 if that male had a longer IPI than the unattractive male and a 2 if that male had a shorter IPI than the unattractive male. The binomial probability of each outcome was calculated using the equation $P(X) = \frac{n!}{(n-x)!x!} \cdot (p)^x \cdot (q)^{(n-x)}$ with a 50% chance of occurrence in either direction.

Results

Of the total 44 female fish used, 24 responded to the male call stimulus. Of these 24 fish, 7 were from the hormone injection trials. Of the 7 responsive Ovaprim-injected females, 3 females were only responsive during one of the trials. From the matched pairs

t-test, females consistently spent the same amount of time in each speaker region across both trials ($t(df=7) = -0.47, p = 0.651$; Figure 3.2). The 4 females that responded in both trials spent more time with the same male across each of their trials. Injected and non-injected fish were not significantly different in their responses ($F\text{-Ratio}(df=1,38) = 3.3907, p = 0.0738$).

In trial 1, 21 females responded to male call stimulus. Winning calls were those where the female spent more time in the speaker's arena. One of the males (male 31) was consistently rejected by females in all 5 of the trials where females were responsive; this male was atypical in sound, with a call that was "tinny" in quality and probabilities were calculated both including and excluding data where this male was an option. Females did spend time at the speaker playing this call and would occasionally choose this speaker first; however, females consistently spent less time at this speaker. Of the responding females, 1 had the option of choosing between males of the same frequency and was discounted from the probability calculation. The remaining 20 females showed an even distribution of choice between high and low frequency males ($P(x=10) = 0.176$). When data from male 31 was removed, the total number of trials dropped to 17 and choosing the low frequency male had a probability of random occurrence of ($P(x=10) = 0.148$) (Figure 3.3a).

Examination of the IPI chosen by responsive females shows that females had a strong preference for male calls with a longer average IPI. When including data from male 31, females chose the male with the longer interpulse interval in 15 of the 21 trials ($P(x=15) = 0.0259$). Without the data using male 31, the number of trials was 17 and the probability of random occurrence of 14 "wins" dropped to ($P(x=14) = 0.00519$) (Figure 3.3b).

Figure 3.2

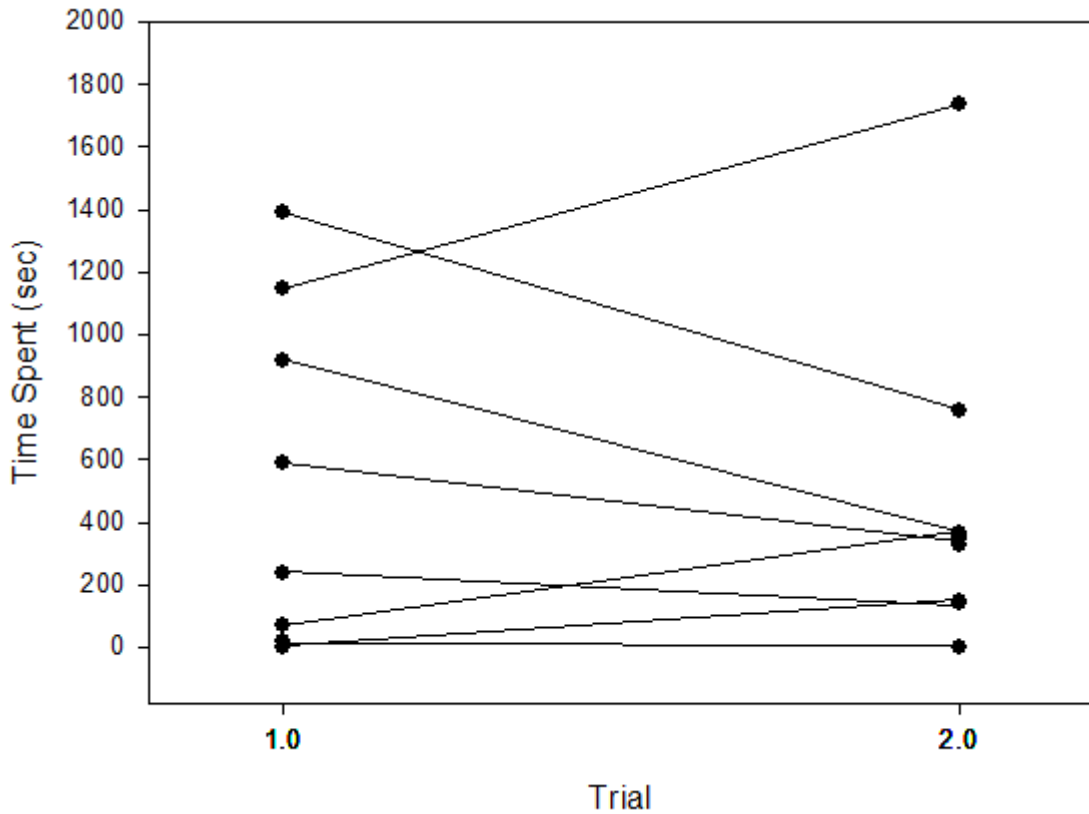


Figure 3.2. Time spent in arena speaker across trials. The 4 hormone injected females that responded in both trials spent similar amounts of time in the same speaker arenas during both trials. These 4 females spent time in both arenas, for a total of 8 choices, during the two trials and time spent in the arena was consistent between trials. Each line represents one chosen speaker, where the time spent at that speaker between trials is not significantly different. The hormone injected females consistently chose the same males across trials, and also spent similar amounts of time in the chosen speaker arena ($t(df=7) = -0.47, p = 0.651$).

Figure 3.3

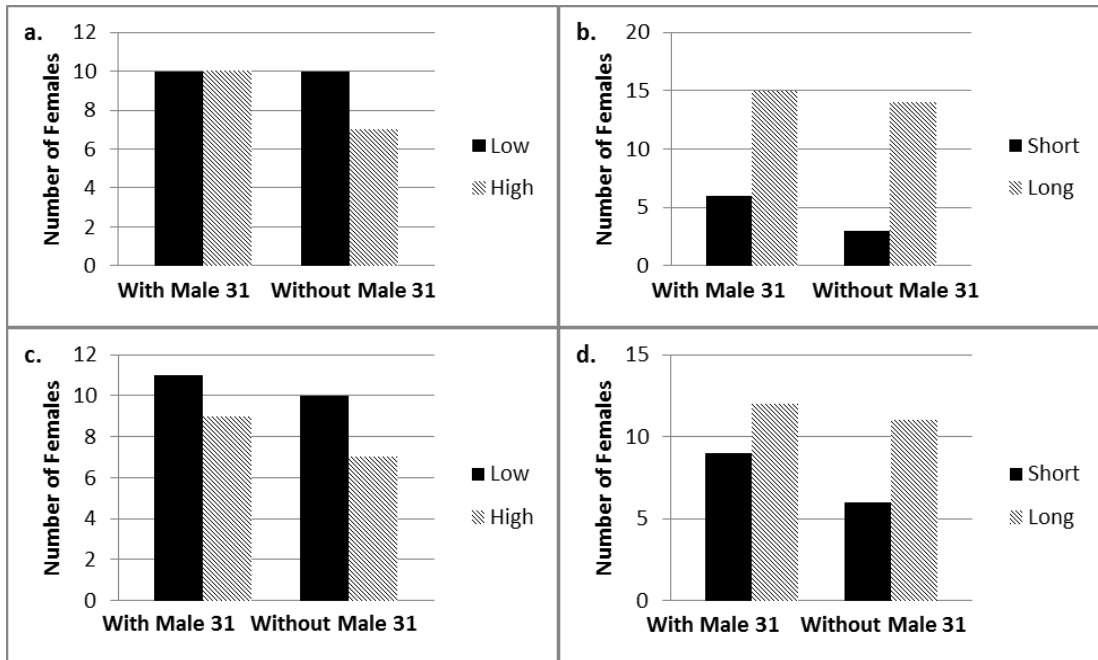


Figure 3.3. Number of females showing call preference. The call with which a female spent the greater amount of time or chose first was classed as the attractive male of the two calls and tallied as the preferred call. a. shows the frequency of the call where attractive males were decided by time spent in the speaker arena by the female; b. shows the interpulse interval of the call where attractive males were decided by time spent in the speaker arena by the female; c. shows the frequency of the call where attractive males were decided by the initially chosen call; and d. shows the interpulse interval of the call where attractive males were decided by the initially chosen call.

Initial call of choice showed a similar pattern, where frequency had a probability of $(P(x=11) = 0.160)$ with male 31 and $(P(x=10) = 0.0944)$ without male 31 (Figure 3.3c). The initial IPI chosen loses some of the distinction, with females only slightly preferring longer IPIs ($(P(x=12) = 0.140)$ with male 31 and $(P(x=11) = 0.0944)$ without male 31) (Figure 3.3d).

None of the GLMs performed showed any clear pattern (Table 3.1). Time spent near speaker was independent of call DF or IPI ($p > 0.05$), nor did it depend on the DF or IPI of the initially chosen call ($p > 0.05$). Removing interaction terms from these models created no change in the results. Latency of the response was also not determined by the initially chosen call's DF or IPI ($p > 0.05$). All residuals were normally distributed.

Discussion

The possibility for female choice is evidenced by the very low probability (2.6% chance of random occurrence) of females choosing males with longer IPI over males with shorter IPI, showing that this was a choice rather than a random occurrence. When choices involving male 31 were removed from the analyses, the probability drops even lower (0.5% chance of random occurrence). The atypical call of male 31 may be due to the hydrophone resting against the wall of the tank, producing a call with a tinny quality. In other fish species, the temporal aspect of the call seems to have a greater impact on female preference (McKibben and Bass, 1998). In the midshipman, females are much more discriminatory over pulse duration than over modulations in frequency, where even short breaks of a few hundred milliseconds in a call would decrease female responsiveness (McKibben and Bass, 2001). Another fish species where temporal

Table 3.1

Table 3.1. Statistical results of female choice GLMs. The results of the Generalized Linear Models with a normal distribution performed on time spent with the chosen speaker and latency of the females' response. No significant results were found.

		DF	ChiSquare	Prob>ChiSq
Time spent	Initial DF chosen	1,38	0.3544	0.5516
	Initial IPI chosen	1,38	0.114	0.7356
	Initial DFxIPI chosen	1,38	0.0202	0.8868
Time spent	DF chosen	1,38	0.1558	0.693
	IPI chosen	1,38	0.7538	0.3853
	DFxIPI chosen	1,38	0.563	0.453
Latency	Initial DF chosen	1,21	0.5266	0.468
	Initial IPI chosen	1,21	1.5234	0.2171
	Initial DFxIPI chosen	1,21	0.2837	0.5943

discrimination has been discovered, the bicour damselfish, *Eupomacentrus partitus*, increases courtship effort when calls with longer pulse intervals are played (Myrberg and Spires, 1972). The enhancement in courtship effort is evident with as little as a 10 msec increase in pulse interval; however, increasing the length of the pulse itself does not affect the courtship effort (Myrberg and Spires, 1972). The fact that female choice was stronger on the basis of time spent rather than speaker of first choice may be suggestive that females will visit multiple male nests before choosing a mate. This seems to be a distinct possibility as round goby will lay their eggs in several nests (Corkum et al., 1998). Acoustic signalling is part of male courtship effort, and so females may assess males on several courtship tactics before mating (Malavasi et al., 2009).

The lack of obvious choice by females based on frequency is not entirely unexpected. Round goby do not have a strong ability to differentiate between low range frequencies (Belanger et al., 2010), however this is condition dependent, as reproductive females are capable of distinguishing differences in frequencies (Zeyl et al., 2013). Even in the midshipman, a species that does display a preference for frequency, the preference is for a call frequency that is expected at a certain temperature and not for a frequency that displays information about the male (McKibben and Bass, 1998). In species that produce sound through vibratory muscles, larger males produce lower frequency calls, as larger muscles take longer to contract (Crawford et al., 1997; Connaughton et al., 2002). Male call frequency may indicate male size accurately, but larger males are not necessarily more capable fathers (Forsgren, 1997). Thus frequency may not be an important factor for females when selecting a mate.

Between trial 1 and trial 2 of the Ovaprim-injected females, all 4 of the hormone treated females that were responsive in both trials were consistent in their responses. The

precision of the amount of time spent between trials at each speaker for the Ovaprim-injected females suggests that female round goby can differentiate between calls and will consistently choose the same male over time (Figure 3.2). Honest signalling likely plays a large part of the reasoning behind choosing the same male consistently. Another possibility may be that females associated certain territories with the call, as each call was played from the same speaker across both trials. Since male gobies have quite high site fidelity (Marentette et al., 2011) and inter-male variability is greater than intra-male variability (Chapter 2), females may have made decisions based on previous experience from trial 1.

The GLMs performed did not show evidence of female choice based on latency of response or time spent in the arena of the speaker of choice. This may be due to the female only having the choice of two males that were randomly selected from the group of 12 male calls. The random playback of two of twelve male calls was deemed necessary so as to avoid pseudoreplication that is so common in sound playback experiments (Slabbekoorn and Bouton, 2008). The limitation of using this method is that comparisons across the entire data set are made. Trends that are clear from individual choices become impossible to discern when examined as a whole. While the individual trials show the female spent more time with the males with longer IPIs, the individual variation of these female choices was great enough to mask patterns present in the overall data set. Females did not spend the most time with the speakers producing the largest IPIs, instead spending roughly equal amounts of time with whichever male of the pair presented to them had the longer IPI. The discrepancy caused by comparing randomly chosen calls could perhaps have been alleviated by having groups of calls from which females could randomly choose, so that a long IPI call would always be compared against a short IPI call. Using

this method would prevent pseudoreplication of only offering the female two choices, while simultaneously presenting two very clear choices that would be easier to analyze.

Certain factors that may be important to female choice were not examined in the current study. The sound pressure level of many species is also a key factor in female choice (Searcy, 1996; Nandi and Balakrishnan, 2013), however all calls were played back at a standard of 140 dB re 1 μ Pa, preventing me from examining this as a potential factor. Likewise, where calls were played continuously to encourage female responsiveness, duration of the call could not be examined as a measure of female choice. Duration is a significant predictor of male size in this species (Chapter 2) as well as in some others (McKibben and Bass, 1998). Future studies delving into female acoustic preference in this species could attempt to modulate these two factors to determine their importance to female decision making.

The current study is one of only a few studies that have examined and found acoustic honest signalling in fish from the female choice perspective. Female choice is critical to the distinction of true honest signals from physiological by-products. This is especially true for acoustic signals, where the characteristics of a call will be highly dependent on the mechanism of production. From an allometric stand point, one would expect call characteristics to scale with the size of the call producing mechanism. While the sound producing mechanism is not known in the round goby, other goby species create sound using the head region (see Chapter 1) and Chapter 2 of this thesis shows that body characteristics interact in complex ways with male head size to alter call characteristics. Future studies examining bioacoustics as honest signals should be sure to incorporate female choice experiments before acoustic signals are deemed honest. Further

research needs to be done to determine what specific benefits female round goby receive by choosing larger males.

References

- Amorim, M. C. P., Simões, J. M., Mendonça, N., Bandarra, N. M., Almada, V. C., & Fonseca, P. J. (2010). Lusitanian toadfish song reflects male quality. *The Journal of experimental biology*, 213(17), 2997-3004.
- Bentsen, C. L., Hunt, J., Jennions, M. D., & Brooks, R. (2006). Complex multivariate sexual selection on male acoustic signalling in a wild population of *Teleogryllus commodus*. *The American Naturalist*, 167(4), 102-116.
- Bowyer, R. T. (1986). Antler characteristics as related to social status of male southern mule deer. *The Southwestern Naturalist*, 289-298.
- Connaughton, M. A., Fine, M. L., & Taylor, M. H. (2002). Weakfish sonic muscle: influence of size, temperature and season. *Journal of Experimental Biology*, 205(15), 2183-2188.
- Corkum, L. D., MacInnis, A. J., & Wickett, R. G. (1998). Reproductive habits of round gobies. *Great Lakes Research Review*, 3(2), 13-20.
- Crawford, J. D., Cook, A. P., & Heberlein, A. S. (1997). Bioacoustic behavior of African fishes (Mormyridae): potential cues for species and individual recognition in *Pollimyrus*. *The Journal of the Acoustical Society of America*, 102(2), 1200-1212.
- Fletcher, N. H. (2004). A simple frequency-scaling rule for animal communication. *The Journal of the Acoustical Society of America*, 115(5), 2334-2338.
- Forsgren, E. (1997). Female sand gobies prefer good fathers over dominant males. *Proceedings of the Royal Society of London B: Biological Sciences*, 264(1386), 1283-1286.
- Hansen, E., & Routledge, D. Evaluation of Induced Spawning Techniques and Requirements in Captive June sucker (*Chasmistes liorus*). (http://wildlife.utah.gov/fes/pdf/induction_of_spawning.pdf)
- Iwasa, Y., & Pomiankowski, A. (1999). Good parent and good genes models of handicap evolution. *Journal of Theoretical Biology*, 200(1), 97-109.
- Kitchen, D. M., Seyfarth, R. M., Fischer, J., & Cheney, D. L. (2003). Loud calls as indicators of dominance in male baboons (*Papio cynocephalus ursinus*). *Behavioral ecology and sociobiology*, 53(6), 374-384.
- Knapp, R. A., & Kovach, J. T. (1991). Courtship as an honest indicator of male parental quality in the bicolor damselfish, *Stegastes partitus*. *Behavioral Ecology*, 2(4), 295-300.
- Le Boeuf, B. J. (1974). Male-male competition and reproductive success in elephant seals. *American Zoologist*, 14(1), 163-176.
- Malavasi, S., Valerio, C., & Torricelli, P. (2009). Courtship sounds and associated behaviours in the Canestrini's goby *Pomatoschistus canestrinii*. *Journal of fish biology*, 75(7), 1883-1887.
- Marentette, J. R., Wang, G., Tong, S., Sopinka, N. M., Taves, M. D., Koops, M. A., & Balshine, S. (2011). Laboratory and field evidence of sex-biased movement in the invasive round goby. *Behavioral Ecology and Sociobiology*, 65(12), 2239-2249.
- McKibben, J. R., & Bass, A. H. (1998). Behavioral assessment of acoustic parameters relevant to signal recognition and preference in a vocal fish. *The Journal of the Acoustical Society of America*, 104, 3520-3533.

- McKibben, J. R., & Bass, A. H. (2001). Effects of temporal envelope modulation on acoustic signal recognition in a vocal fish, the plainfin midshipman. *The Journal of the Acoustical Society of America*, 109(6), 2934-2943.
- Myrberg, A. A., & Spires, J. Y. (1972). Sound discrimination by the bicolor damselfish, *Eupomacentrus partitus*. *Journal of experimental biology*, 57(3), 727-735.
- Nandi, D., & Balakrishnan, R. (2013). Call intensity is a repeatable and dominant acoustic feature determining male call attractiveness in a field cricket. *Animal Behaviour*, 86(5), 1003-1012.
- Ryan, M. J., Fox, J. H., Wilczynski, W., & Rand, A. S. (1990). Sexual selection for sensory exploitation in the frog *Physalaemus pustulosus*. *Nature*, 343, 66-67.
- Schulte-Hostedde, A. I., Zinner, B., Millar, J. S., & Hickling, G. J. (2005). Restitution of mass-size residuals: validating body condition indices. *Ecology*, 86(1), 155-163.
- Searcy, W. A. (1996). Sound-pressure Levels and Song Preferences in Female Red-winged Blackbirds (*Agelaius phoeniceus*)(Aves, Emberizidae). *Ethology*, 102(2), 187-196.
- Slabbekoorn, H., & Bouton, N. (2008). Soundscape orientation: a new field in need of sound investigation. *Animal behaviour*, 76(4), e5-e8.
- Vannoni, E., & McElligott, A. G. (2008). Low frequency groans indicate larger and more dominant fallow deer (*Dama dama*) males. *PloS one*, 3(9).
- Weatherhead, P. J., & Robertson, R. J. (1979). Offspring quality and the polygyny threshold: "the sexy son hypothesis". *American Naturalist*, 201-208.
- Zeyl, J. N., Love, O. P., & Higgs, D. M. (2013). Condition-dependent auditory processing in the round goby (*Neogobius melanostomus*): links to sex, reproductive condition and female estrogen levels. *The Journal of experimental biology*, 216(6), 1075-1084.

Chapter IV

CONCLUSIONS AND RECOMMENDATIONS

This thesis expands our understanding of the roles of bioacoustics in fish, particularly in terms of honest signals and male call variation. Previous work from our lab showed that the round goby is capable of localizing to a sound source (Rollo et al., 2007), distinguishing conspecifics from heterospecifics (Rollo and Higgs, 2008) and that auditory responses are dependent on the sexual condition of the fish (Zeyl et al., 2013). We have now also added that calls correlate strongly with male body characteristics and that females are capable of detecting these differences.

In Chapter 2, I collected calls from round goby at night in the presence of a reproductive female and analyzed several characteristics of these calls. Principal Components Analysis showed a strong association among Interpulse Interval (IPI), Total Length (TL), Total Weight (TW) and Head Width (HW) as well as links between Dominant Frequency (DF), Call Duration and Number of Pulses. Gonadosomatic Index was not highly related to any call or body features and was discounted from further analyses. Generalized Linear Models used individual call characteristics as dependent variables and incorporated the remaining body characteristics as explanatory variables. Interaction terms in the models were significant for all three call variables. In the case of frequency and IPI, the interactions between HWxTL and HWxTW worked in opposite directions. The three-way interaction of HWxTLxTW that affected call duration showed that a decreased TW greatly decreased call duration. The shortest calls were produced by long, thin males with big heads.

These results suggest that there is a very complex interplay between male body morphometrics and the calls that are produced. The fact that calls do correlate to male

body condition also provides evidence of the potential that these calls function as true honest signals of male body size. The next Chapter focused on validating whether or not females actually could differentiate between male calls and would have a preference for certain call features.

Chapter 3 used 12 of the calls recorded from the male portion of the study to determine if female round goby would display a preference for call characteristics when presented with 2 randomized calls. Calls were played continuously with no breaks for the duration of each trial and female responsiveness in the form of latency to respond, initial speaker of choice and time spent in each speaker zone was quantified. While there was no relationship with latency to respond, there was a pattern observed for time spent in the speaker zone with regards to IPI of the calls being played. The pattern was similar when considering the IPI and the initial speaker of choice, albeit not as strong. The chances of females randomly choosing the longer IPI in 15 out of 21 trials was extremely low ($P(x=15) = 0.0259$), suggesting that female round goby prefer male calls with a longer interpulse interval.

Taken together, the results of both chapters suggest that females prefer males that have large heads and high body mass. Male length affects call parameters in an unexpected way, making the effects of individual body characteristics much harder to discern. While the frequency of a male call was also strongly correlated to an individual's body morphometrics, females did not show any preference for this trait. Male size is known to alter call frequency in several species of fish (Crawford et al., 1997; Connaughton et al., 2002); however, females may be more interested in calling effort typically presented by better condition males than simply the largest males (Amorim et al., 2010). Males that put forth greater effort during courtship may be less likely to

cannibalize their brood, perhaps because of greater energy reserves (Forsgren, 1997). If interpulse interval is directly related to male energy reserves, the female would benefit from increased survival of her young. Further studies into honest signalling in round goby could assess male body lipid content and how this body characteristic affects courtship and paternal care.

Our understanding of the ecology of this invasive species is slowly increasing. Greater knowledge of how the round goby breeds is important to help prevent the spread of this species and other future invaders from this family. Incorporating passive acoustic monitoring into management practices could allow for more specific targeting of the larger more desirable males, and aid in early detection of a moving invasion front. Round gobies at the edges of an established area that disperse to establish new populations are often larger (Brandner et al., 2013) or much more aggressive (Groen et al., 2012). Using acoustic monitoring to determine which invasion fronts are more likely to be successful will more effectively slow the spread of this invasive species. While I realize that the round goby will likely never be successfully eradicated from the regions in which it has now become established, it is still important to prevent the movement of this species into new bodies of water.

This study is also very important in terms of the overall understanding of fish bioacoustics and understanding the diverse uses of underwater acoustic signalling. There are important benefits of determining in which fish species acoustic cues reliably signal mate condition, especially as anthropogenic impacts on aquatic ecosystems increases. Increasing our understanding of fish reproduction could aid us in reducing the detrimental effects of human activities. Where the Gobiidae is such a large family, there is still much to learn about the individual species within this group. It is also very likely that a great

many more species than the ones highlighted in Chapter 1 are soniferous. Ideally, field studies will also be done on this topic as acoustic laboratory studies usually cannot be used to represent the natural environment. The restrictions of the tank affect the sound being produced by the animals, preventing a direct comparison between lab and field-based studies.

True acoustic honest signalling in fish is not widely examined. This thesis highlights the importance of confirming the occurrence of female choice before signals can be labeled as honest indicators of male “quality”. The confirmation of female preference for certain male call variables in the round goby is one of the few studies to take this approach to honest signalling in fish. This study also highlights the abilities for sound detection in a fish with no swim bladder or auditory accessory organs.

References

- Amorim, M. C. P., Simões, J. M., Mendonça, N., Bandarra, N. M., Almada, V. C., & Fonseca, P. J. (2010). Lusitanian toadfish song reflects male quality. *The Journal of Experimental Biology*, *213*(17), 2997-3004.
- Brandner, J., Cerwenka, A. F., Schliewen, U. K., & Geist, J. (2013). Bigger is better: characteristics of round gobies forming an invasion front in the Danube River. *PLoS One*, *8*(9).
- Connaughton, M. A., Fine, M. L., & Taylor, M. H. (2002). Weakfish sonic muscle: influence of size, temperature and season. *Journal of Experimental Biology*, *205*(15), 2183-2188.
- Crawford, J. D., Cook, A. P., & Heberlein, A. S. (1997). Bioacoustic behavior of African fishes (Mormyridae): potential cues for species and individual recognition in *Pollimyrus*. *The Journal of the Acoustical Society of America*, *102*(2), 1200-1212.
- Forsgren, E. (1997). Female sand gobies prefer good fathers over dominant males. *Proceedings of the Royal Society of London B: Biological Sciences*, *264*(1386), 1283-1286.
- Groen, M., Sopinka, N. M., Marentette, J. R., Reddon, A. R., Brownscombe, J. W., Fox, M. G., Marsh-Rollo, S. E., & Balshine, S. (2012). Is there a role for aggression in round goby invasion fronts?. *Behaviour*, *149*(7), 685-703.
- Rollo, A., Andraso, G., Janssen, J., & Higgs, D. (2007). Attraction and localization of round goby (*Neogobius melanostomus*) to conspecific calls. *Behaviour*, *144*(1), 1-21.
- Rollo, A., & Higgs, D. (2008). Differential acoustic response specificity and directionality in the round goby, *Neogobius melanostomus*. *Animal Behaviour*, *75*(6), 1903-1912.
- Zeyl, J. N., Love, O. P., & Higgs, D. M. (2013). Condition-dependent auditory processing in the round goby (*Neogobius melanostomus*): links to sex, reproductive condition and female estrogen levels. *The Journal of Experimental Biology*, *216*(6), 1075-1084.

VITA AUCTORIS

NAME: Meghan Donovan

PLACE OF BIRTH: Ottawa, Ontario

YEAR OF BIRTH: 1991

EDUCATION: Glebe Collegiate, Ottawa, ON
2005-2009

Memorial University, St. John's, NL
2009-2012 B.Sc. Honours

University of Windsor, Windsor, ON
2013-2015 M.Sc.