

1 *In situ* source levels of mullo way (*Argyrosomus japonicus*) calls.

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27 **ABSTRACT**

28

29 Mullet ( *Argyrosomus japonicus* ) in Mosman Bay, Western Australia produce three  
30 call categories associated with spawning behavior. The determination of call source  
31 levels and their contribution to overall recorded sound pressure levels is a significant  
32 step towards estimating numbers of calling fish within the detection range of a  
33 hydrophone. The source levels and ambient noise also provide significant  
34 information on the impacts anthropogenic activity may have on the detection of *A.*  
35 *japonicus* calls. An array of four hydrophones was deployed to record and locate  
36 individual fish from call arrival-time differences. Successive *A. japonicus* calls  
37 produced samples at various ranges between 1 and 100 m from one of the array  
38 hydrophones. The three-dimensional localization of calls, together with removal of  
39 ambient noise, allowed the determination of source levels for each call category using  
40 observed trends in propagation losses and interference. Mean source levels (at 1 m  
41 from the hydrophone) of the three call categories were calculated as:  $163 \pm 16$  dB re  
42  $1 \mu\text{Pa}$  for Category 1 calls (short call of 2-5 pulses);  $172 \pm 4$  dB re  $1 \mu\text{Pa}$  for Category 2  
43 calls (long calls of 11-32 pulses); and  $157 \pm 5$  dB re  $1 \mu\text{Pa}$  for Category 3 calls (series  
44 of successive calls of 1-4 pulses, increasing in call rate).

45

46 Keywords: source level, soniferous, range, localization, passive acoustics, fish  
47 calls

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51 **I INTRODUCTION**

52

53 Many species of fish use vocalizations as social cues for spawning behavior (Winn,  
54 1964, Myrberg and Spires, 1972, Fine *et al.*, 1977) and often aggregate in large  
55 numbers to call *en masse* (Luczkovich *et al.*, 1999, Parsons *et al.*, 2006). Such  
56 choruses are often used to delineate areas used by spawning aggregations (Luczkovich  
57 *et al.*, 1999, Parsons *et al.*, 2006). However, the ultimate goal of fisheries passive  
58 acoustics is the unbiased estimation of abundance. The determination of absolute  
59 numbers of mobile animals such as aggregating fish requires the ability to  
60 simultaneously census large volumes of water and to eliminate bias from multiple  
61 detections of the same fish. The sensitivity of hydrophones in areas of low ambient  
62 noise allows the discrimination of fish calls emitted at large ranges from the receiver  
63 and in some cases the ability to monitor individual fish over time, by listening and  
64 localizing their calls (Parsons *et al.*, 2009).

65 One method of estimating the number of fish vocalizing within the detection range of  
66 a hydrophone is to measure the contribution of individual callers to the overall sound  
67 pressure levels (SPLs) received (McCauley, 2001, Sprague and Luczkovich, 2002).  
68 To achieve these caller density estimates *a priori* knowledge of species call source  
69 level (SL) is required (Sprague and Luczkovich, 2004). In addition to SL, the  
70 transmission characteristics of a call in its local environment require quantification to  
71 assess the proportion of emitted energy that is received by the hydrophone  
72 (McCauley, 2001).

73

74 To quantify a call SL it is necessary to isolate energy associated with the signal.  
75 Mulloway (*Argyrosomus japonicus*) often spawn in groups (Ueng *et al.*, 2007) and

76 can produce high densities of overlapping calls in the wild (Parsons *et al.*, 2006),  
77 complicating the discrimination of separate calls. However, during the  
78 commencement of an evening spawning cycle in the Swan River, Perth, Western  
79 Australia, calls are of sufficiently low density to offer the opportunity to monitor  
80 individual fish (Parsons *et al.*, 2009).

81

82 The accurate measurement of SL requires the determination of source range. Short  
83 ranges can be easily observed *in aquaria*; however, internal tank reflections and  
84 reverberation often contribute significant complications towards SL calculation. *In*  
85 *situ* SLs of fish calls are rarely reported (Cato, 1998, McCauley, 2001) and *in situ*  
86 visual confirmation of vocalization at a known reference distance is even less  
87 frequent. One example of such audio and visual recording was of an individual  
88 sciaenid, a silver perch (*Bairdiella chrysoura*), fortuitously caught on camera at the  
89 time of vocalization (Sprague and Luczkovich, 2004). For Sciaenidae, this elusive  
90 behavior is in part due to their propensity to vocalize in turbid waters after sunset,  
91 thereby inhibiting discrimination of source range by visual methods. Thus the  
92 identification of source position or range remains the restricting factor in the accurate  
93 determination of *in situ* fish call SL.

94

95 Fish calls offer significant information about the caller to the intended recipient and,  
96 inadvertently, the observer. The size of the individual can be related to both the call  
97 SL and spectral peak frequency (Connaughton *et al.*, 2000). However, calls emitted  
98 by the same fish are not all necessarily of the same SL. For example, Lagadere and  
99 Mariani (2006) proposed that the short grunts of French meagre (*A. regius*) are  
100 weaker than the long grunts. Similar results from *in situ* *A. japonicus* calls have been

101 observed, but it was not confirmed whether pressure wave amplitude differences were  
102 due to range, weaker swim bladder twitches or multiple ray-path interference (Parsons  
103 *et al.*, 2006, 2009). Therefore measurement of many SLs may be required to  
104 categorize a fish call. It is the aim of this study to determine SL ranges for each  
105 category of call exhibited by *A. japonicus* in an aggregation in Mosman Bay, Swan  
106 River.

107

## 108 **II METHODS**

109

110 A passive acoustic hydrophone array was deployed in Mosman Bay (Figure 1) to  
111 localize individual fish within a spawning aggregation of *A. japonicus*. In Mosman  
112 Bay the river banks descend rapidly to a 21 m deep channel comprising a sand/silt  
113 substrate, a few artificial reefs and several depressions, some of which reach 22 m  
114 deep at high tide. At four locations calibrated, omni-directional, HTI 90-U and 96-  
115 min hydrophones (Hi-Tech Inc., MS, USA) were attached either to an autonomous  
116 sea-noise logger (developed at Curtin University of Technology, Western Australia  
117 and the Defence Science and Technology Organisation (DSTO)), or to Sony Digital  
118 Audio Tape (DAT) recorders. The deployment duration encompassed an *A. japonicus*  
119 evening spawning period in approximately 21 m of flat water above a relatively  
120 uniform silt substrate riverbed. Recordings were sampled at 10,416 Hz and were time  
121 synchronized on a half hourly basis by the implosion of a light bulb at a known  
122 location and depth (Parsons *et al.*, 2009). To limit sound energy from sources other  
123 than the *A. japonicus* calls (Parsons, 2010), hi- and low-pass filters of 50 and 1000 Hz  
124 were applied to the data.

125

126 Calls were localized by the arrival-time difference technique (Cato, 1998) and  
127 location error ellipses provided for each call. The localization of calls and associated  
128 error ellipsoids from the hydrophone array allowed the determination of source  
129 ranges, the details of which were outlined by Parsons (2010) and Parsons *et al.*  
130 (2009). To minimize noise error, calls analyzed for SL in this study were recorded at  
131 a time of relatively little background noise, an example of which is shown in Figure 2.

132

133 *A. japonicus* have been shown to emit three categories of swim bladder driven calls  
134 (Parsons, 2010, Parsons *et al.*, 2006, 2009, 2010; Figure 3). Each of these call  
135 categories varies in duration and has been suggested to have different associated  
136 behaviors. Categories were therefore analyzed separately to observe variations in SL.  
137 Category 1 calls are short, comprising 2-5 swim bladder pulses. Category 2 calls  
138 comprise 11-32 pulses while Category 3 calls are a series of 1-4 pulse calls in quick  
139 succession.

140

141 Sound pressure level (SPL) or sound level  $L_p$  is a logarithmic measure of the effective  
142 sound pressure of a sound relative to a reference value. It is measured in decibels (dB)  
143 above a standard reference level:

144 
$$L_p = 10 \log_{10} \left( \frac{p_{rms}^2}{p_{ref}^2} \right) = 20 \log_{10} \left( \frac{p_{rms}}{p_{ref}} \right) \text{ dB} \quad (1)$$

145 where  $p_{rms}$  is the root-mean-square pressure and  $p_{ref}$  is the reference pressure.

146

147 To accurately determine fish call SL it is necessary to first remove the background  
148 noise. For this purpose *A. japonicus* calls and background noise were considered as

149 incoherent signals. By Parseval's Theorem, the time-averaged squared total pressure  
 150 recorded by the logger was equal to the sum of the time-averaged squared partial  
 151 pressure of each constituent signal (Sprague and Luczkovich, 2004). The level of fish  
 152 call signal ( $L_s$ ), once background noise is removed, is given as:

$$153 \quad L_s = 10 \log_{10} \left( 10^{\frac{L_{s+n}}{10}} - 10^{\frac{L_n}{10}} \right) \quad (2)$$

154 where  $L_{s+n}$  was the level in dB re 1  $\mu$ Pa of the overall signal and  $L_n$  is the background  
 155 noise level (McCauley, 2001).

156

157 The method of call energy level analysis in this study is based on theory protocols  
 158 from McCauley (2001). For a plane wave, which is taken to be a good approximation  
 159 for recordings, the energy per unit area (or "energy flux"), over the duration  $T$  of a  
 160 signal where  $T_0$  and  $T_e$  denote the signal start and end, is given by:

$$161 \quad \frac{E}{A} = \int_{T_0}^{T_e} I dt = \int_{T_0}^{T_e} \frac{p_{s+n}^2(t)}{\rho c} dt \quad (3)$$

162 where  $E$  is the signal energy;  $A$  the unit area; and  $I$  the intensity, defined by the  
 163 combined signal and noise pressure  $p_{s+n}$ , and the characteristic acoustic impedance  $\rho c$ .

164 The mean-squared pressure values ( $\overline{p_{s+n}^2}$ ) of the total received signal can be derived  
 165 as:

$$166 \quad \overline{p^2} = \frac{1}{T} \int_{T_0}^{T_e} p^2(t) dt \quad (4)$$

167 Thus, from Equation 4 the 'equivalent signal energy',  $E_s(t)$  is given by:

168 
$$E_s(t) = \int_0^T p_{s+n}^2(t) dt - \int_{T_n}^{T_n+T} p_n^2(t) dt \quad (5)$$

169 and the peak-to-peak pressures ( $p_{pp}$ ) are given as:

170 
$$p_{pp} = \max[p(t)] + |\min[p(t)]| \quad (6)$$

171 where  $\max[p(t)]$  and  $\min[p(t)]$  are the respective maximum positive and minimum  
 172 negative values from the pressure wave form (McCauley, 2001, Southall *et al.*, 2007).  
 173 Figure 4 displays three of the steps involved in analyzing the acoustic pressure  
 174 attributable to a call and determining the frequency band over which the majority of  
 175 call energy occurs. A digitized segment of the recording, including the call and  
 176 encompassing a minimum of 500 sample points either side of the call, was converted  
 177 to pressure wave form (Figure 4A). An estimation of the 'sound exposure level' (SEL)  
 178 was made by integrating the pressure squared over the duration of the call in the form  
 179 of:

180 
$$E_s(t) = T_s \sum_{i=1}^{i=k} (p_{(s+n)_i}^2 - \overline{p_n^2}) \quad (7)$$

181 where  $T_s$  is the signal length,  $p_{(s+n)_i}$  is the  $i^{\text{th}}$  element of the pressure wave form  
 182 (containing signal and noise),  $k$  is the last sample point and  $\overline{p_n^2}$  is the mean-squared  
 183 noise pressure from noise samples, obtained from the level of  $p^2$  immediately before  
 184 or after the call (McCauley, 2001). Pressure levels within the 5% and 95% region of  
 185 the total cumulative energy curve (Figure 4B) were calculated (Malme *et al.*, 1986)  
 186 thus standardizing the averaging time to that at which 90% of the energy from the  
 187 entire signal (less noise) had passed. The call length was taken as the time for 90% of  
 188 the signal energy to pass. A power spectral density of each call was produced to



189 observe spectral peak frequencies compared with each calculated energy level (Figure  
190 4C) and aid confirmation of repetitive calling by individual fish (Parsons *et al.*, 2009).

191 The received level (RL) can be related to the SL and simple transmission loss (TL) in  
192 the form of:

$$193 \quad RL = A \log_{10} r + SL \quad (8)$$

194 where  $A$  is the transmission loss constant,  $r$  is the slant range (the shortest range  
195 between two points of differing altitude; Urick, 1983). RL is thus assumed a linear  
196 function of  $\log_{10}r$ . Linear regression of RL against  $\log_{10}r$  (Walpole and Myers, 1985)  
197 was carried out to provide estimates of the TL constant ( $A$ ) and the SL. Correlation  
198 values ( $R^2$ ) were calculated to determine the variance in RLs explained by the  
199 regression models. The 95% confidence intervals for the estimates of  $A$  and  $B$  were  
200 calculated using the methods described by Walpole and Myers (1985), as were 95%  
201 confidence intervals for predictions of received levels from sources at any given range  
202 along the regression curve. These are shown as the 95% confidence interval  
203 boundaries on the transmission loss plots in this report.

204

205 The estimate of losses due to geometrical spreading provides a minimum loss on  
206 which to base initial calculations (Sprague and Luczkovich, 2004). Although surface  
207 reflections were observed in the reported data, in the context of SL calculation  
208 spherical spreading was considered as a minimum estimate for transmission losses for  
209 calls in 20 m depth water, at ranges of less than 100 m (Cato, 1980). Losses due to  
210 absorption and water movement were considered of negligible impact. The estimated  
211 transmission loss trends were compared to spherical ( $20\log_{10}(r)$ ) and cylindrical  
212 ( $10\log_{10}(r)$ ) spreading losses, where  $r$  is measured in meters (Urick, 1983), to help  
213 validate the regression models.

214

215 Signal SL is often presented in a variety of formats and to help comparison of the  
216 results reported here with other data the regression models were applied to estimate  
217 SLs not only in SPL (dB re 1 $\mu$ Pa at 1 m), but also sound exposure level (SEL) and  
218 peak-to-peak pressure for each call category. Sound exposure is the integral of the  
219 pressure squared over the duration of the call and SEL is given in dB re 1 $\mu$ Pa<sup>2</sup>.s at 1  
220 m. Additionally, as TL is not always calculated during estimates of signal SL a final  
221 method of estimating SL was to backstep from a RL at known range to a range of 1 m  
222 using spherical spreading  $20\log_{10}r$  as the TL. All SLs were estimated to the reference  
223 pressure at a range of 1 m from the source.

224

### 225 **III RESULTS**

226

227 Parsons *et al.* (2009) localized 213 calls (65 and 148 Category 1 and 2 calls,  
228 respectively) using the array. Several calls of each category included overlap with  
229 adjoining calls, background vessel noise, or insufficient number of sample points for  
230 noise removal, and were discarded. The remaining calls, which contained sufficient  
231 noise sample points, were analyzed to determine received SPLs. Of the total localized  
232 calls, 53 Category 1 and 112 Category 2 calls at ranges of approximately 20 to 100 m  
233 offered signals of sufficient clarity to analyze SPLs. One fish, in particular, was  
234 tracked (Parsons *et al.*, 2009), which produced 65 Category 2 calls over a 4-minute  
235 period for SPL analysis in this study. Source levels and regression-determined  
236 transmission losses of call categories are shown in Table I and are discussed with the  
237 results of each call category.

238

239 Figure 5 illustrates the effects of transmission loss with range for localized Category 1  
240 calls. Category 1 call wave forms originating from different fish were often disparate  
241 in structure (Parsons, 2010). These calls varied, not only in the number of pulses, but  
242 in the pressure amplitude of those pulses. In 62% of the Category 1 calls the  
243 maximum amplitude of the first pulse was less than 80% of that of the second pulse  
244 (Parsons, 2010). The linear regression of SPLs from all the 53 Category 1 calls  
245 produced a SL of  $163 \pm 16$  dB re  $1\mu\text{Pa}$  at 1 m and estimated spreading losses of  $25.4$   
246  $\log_{10}(r)$  (Figure 5 and Table I). The transmission losses evident in the data of the  
247 Category 1 calls more closely resembled that of spherical spreading than cylindrical  
248 spreading (Figure 5, compare the slope of the regression determined continuous line  
249 with the dashed and dot-dash lines of spherical and cylindrical spreading,  
250 respectively). Correlation of call SPLs with the transmission loss curve was lower  
251 than any other call category (Table I,  $R^2 = 0.41$ ).

252

253 Category 2 calls exhibited little variation of regression-calculated SL (Figure 6,  $R^2 =$   
254  $0.82$ ). Similar to Category 1, the Category 2 call transmission losses more closely  
255 resembled spherical than cylindrical spreading losses (Figure 6, continuous, dash and  
256 dot-dashed lines, respectively). The Category 2 call SL was determined as  $172 \pm 3.6$   
257 dB re  $1\mu\text{Pa}$  at 1 m (Table I). Similar to the tracked individual reported by Parsons *et*  
258 *al.* (2009), SPLs from all Category 2 calls exhibited ray multi-path interference  
259 features, evident as the received signal level oscillating with range. This interference  
260 appeared to be more pronounced in the SPLs of the single caller at ranges between 50  
261 and 100 m (Figure 6, circles).

262

263 During localization recordings an individual fish emitting Category 3 calls  
264 approached the bottomed hydrophone. The range was determined by geometry from  
265 the arrival-time difference between the initial wave form peak associated with the call  
266 and the first surface reflection of this peak received by the hydrophone (Parsons *et al.*,  
267 2010). One of the calls was emitted from within 1.6 meters of the hydrophone and  
268 therefore the fish must have been swimming close to the riverbed. For range  
269 determination purposes it was assumed that the fish continued at the same depth.  
270 Localization of Category 2 calls has shown that they were generally emitted from  
271 positions on, or near, the riverbed (Parsons *et al.*, 2009). The individual fish provided  
272 both single (n = 11) and double (n = 17) pulse calls for SL analysis, at a variety of  
273 ranges up to 16 m. Ranging of Category 3 calls of greater than 2 pulses was not  
274 conducted.

275

276 Based on the assumption that the individual remained in the same orientation  
277 throughout its calls, the recordings as the fish swam past the hydrophone provided a  
278 comparison of SL between orientations of head towards and away. It was assumed  
279 that the fish was swimming forwards and not drifting backwards with the current at an  
280 alternate orientation. The consistency of call time with range suggests that a direct  
281 route at approximately  $0.5 \text{ ms}^{-1}$  was the case. Figure 7 displays the received SPL with  
282 range for the fish swimming towards, and past, the hydrophone. The 'o' and 'x'  
283 markers indicate 1 and 2 pulse calls respectively, emitted as the fish approaches  
284 (dashed line) and departs (dot-dashed line). Once the fish was approximately 16  
285 meters past the hydrophone the calls were emitted in such quick succession, with very  
286 little range variation, that neither toward or away orientation was assumed. The least

287 squares regression curve for all Category 3 calls, together with 95% confidence limits  
288 are shown in Figure 7 by the continuous and dotted lines respectively.

289

290 The regression model for all Category 3 calls estimated a SL of  $157 \pm 5.2$  dB re  $1\mu\text{Pa}$   
291 at 1 m, lower than both Category 1 and 2 calls. Calculated SLs of calls employing  
292 two swim bladder pulses in both orientations were greater than those of the single  
293 pulse calls (6.2 and 2.4 dB re  $1\mu\text{Pa}$  at 1 m greater towards and away from the  
294 hydrophone respectively). In both types of call the SLs were greater with fish facing  
295 towards the hydrophone (greater by 4.7 and 8.5 dB re  $1\mu\text{Pa}$  at 1 m for single and  
296 double pulse calls respectively; Table I, Figure 7). The sample size was small for the  
297 Category 3 calls, and even smaller when grouped into the 1 and 2 pulse calls and the  
298 orientation. Considering the sample size and the large overlapping confidence  
299 intervals (partially as a result of the sample size; Table I) these observations are not  
300 statistically significant. However, it was only the fortuitous passing of the calling fish  
301 at close range that allowed the inference of orientation and therefore the grouping of  
302 these Category 3 calls. This is an unlikely event and difficult to reproduce, especially  
303 without inducing behavioral bias, thus the differences in SL between the four types of  
304 Category 3 calls are reported as a point of note only.

305

306 Figure 8 shows the distribution of SLs for each category based on the determined  
307 transmission losses to the receiver for each category, which were back calculated from  
308 the recorded SPLs and range. It is thought that the lack of variance in Category 2 call  
309 SL distribution was due to the fact that more than half the calls were emitted by one  
310 fish. By comparison, the Category 1 call SLs were derived from a number of fish. In  
311 addition, the variability observed in short calls (between 1 and 4 pulses) has a greater

312 effect than that found in Category 2 long calls (11-30 pulses). The SL distribution of  
313 all Category 3 calls was affected by the differing orientation of the fish and pulse  
314 number.

315

316 The SL regression models were repeated to determine the SEL and peak-to-peak  
317 pressure SLs for each category (Table II). In each category the difference between  
318 SPL and SEL is determined by the call lengths. This is because the root-mean-square  
319 SPL (dB re 1  $\mu$ Pa) is equal to the SEL (dB re 1  $\mu$ Pa<sup>2</sup>.s) minus  $10\log_{(10)}(\text{call length})$ .

320

321 Mean background noise levels during the course of these recordings were observed at  
322 108 dB re 1 $\mu$ Pa (over the same 50-1000 Hz bandwidth as the call frequency analysis).  
323 However, during the course of Mosman Bay recordings in this study noise levels  
324 which include vessel traffic and/or calling *A. japonicus* have been known to reach 148  
325 dB re 1 $\mu$ Pa. The background noise has significant impact on the detection range of a  
326 call to both intended recipient and observer (Urlick, 1983, Sprague and Luczkovich,  
327 2004). For simple comparative purposes a maximum detection range of the  
328 regression determined SL was calculated for each call category (Table III). This is the  
329 range at which the signal would have attenuated to background noise level using  
330 signal processing based on the estimated transmission losses and also using spherical  
331 spreading as the transmission loss. It does not use statistical analysis of the  
332 probability of signal detection or account for fish hearing critical ratios and so could  
333 not be used as an estimation of the ranges over which fish may detect calls. The ratio  
334 of detection ranges of a call with and without the noise of a passing vessel, however,  
335 is comparable and shows that, even under near ideal detection conditions,

336 anthropogenic activity can reduce detection of such calls from hundreds of meters, to  
337 less than 10 m.

338

#### 339 **IV DISCUSSION**

340

341 A simple method to determine fish call SL would be concurrent calibrated audio and  
342 visual recording, at a known distance from the hydrophone, similar to that reported by  
343 Sprague and Luczkovich (2004). However, visual confirmation of behaviorally  
344 unbiased, *in situ*, fish calls, in dark or turbid waters, is improbable at such short  
345 ranges. In typical SPL calculations the origin is considered as a point source. Due to  
346 visibility, a caller in dark or turbid waters would be required to be at such close range  
347 that it may no longer be considered as a point source. Furthermore, the observed  
348 disparities in SL, due to different fish and interference suggests that a range of  
349 observed call SPLs is required to model the species SLs. Therefore precisely locating  
350 individuals from call arrival-time differences to determine accurate source ranges of  
351 species-specific calls is the most appropriate method for recording SLs.

352

353 Estimated SLs of 163, 172 and 157 dB re 1 $\mu$ Pa at 1 m with associated confidence  
354 limits for *A. japonicus* call Categories 1, 2, and 3 have been established.  
355 Transmission losses observed in the linear regression models for Categories 1 and 2  
356 were within acceptable range of spherical spreading to ‘practical’ spreading loss  
357 (Coates, 1990) and significantly greater than that of cylindrical spreading. Spherical  
358 spreading has been considered satisfactory to ranges of around 50 m in water depths  
359 of approximately 20 m (Cato, 1998).

360

361 Due to the variability of the call structure Category 1 calls displayed comparatively  
362 low correlation with the least squares regression transmission loss model, compared to  
363 that of the other call categories. The difference in amplitude between the first two  
364 pulses of a call varied significantly throughout all three call categories. This variation  
365 had greater impact on SLs of short calls comprising fewer pulses. Therefore  
366 determined confidence limits of 148 and 179 dB re 1 $\mu$ Pa at 1 m with a correlation of  
367  $R^2 = 0.42$  for the best fit ( $25.39\log_{10}(r)$ ) transmission losses were deemed a  
368 reasonable estimate of SL range for Category 1 calls.

369

370 SLs increased through Categories 3, 1 and 2 as more pulses were included in calls.  
371 This was due, in part, to an increase in amplitude over the first three pulses of several  
372 calls. Parsons (2010) highlighted that in many long Category 2 calls the first one, two  
373 or three pulses were of lower amplitude than subsequent pulses. To corroborate this,  
374 double pulse Category 3 calls when the fish was orientated towards the hydrophone  
375 produced SLs comparable with those of the Category 1 calls (Table I). Lagadere and  
376 Mariani (2006) observed similar traits in the short calls of *Argyrosomus regius* and  
377 suggested these were less intense than long grunts, a hypothesis in agreement with  
378 findings of this study. After the initial pulses the detected maximum amplitude often  
379 reached a plateau, and as a result, Category 2 call SLs were more consistent than those  
380 of Category 1. It is suggested that the high correlation of Category 3 call SL within  
381 the single and double pulse trends were due to the single source origin. Whether this  
382 trait is characteristic of an individual is unknown.

383

384 Source levels of each *A. japonicus* category of call were significantly greater than the  
385 reported SLs of calls from other species. *In situ*, silver perch (*Bairdiella chrysoura*)



386 and oyster toadfish (*Opsanus tau*) reportedly have maximum SLs of 135 and 132 dB  
387 re 1 $\mu$ Pa at 1 m, respectively (Sprague and Luczkovich, 2004; Barimo and Fine, 1998),  
388 and typically grow to total lengths of 30 and 43.2 cm, respectively (Robins and Ray,  
389 1986). As call SLs are related to fish size (Connaughton *et al.*, 2000) and these  
390 species are considerably smaller than *A. japonicus* captured in the Swan River (greater  
391 than 1 m; Farmer, 2008), the comparatively lower SLs are to be expected. Cato  
392 (1980) and McCauley (2001) reported SLs of 148 ( $\pm$  4) and 149 ( $\pm$  2) dB re 1 $\mu$ Pa at 1  
393 m, respectively, from sources speculated to be *Protonibea diacanthus*, a Sciaenidae of  
394 similar sound production mechanism to *A. japonicus*. Both of these reports comprised  
395 calls of similar spectral peak frequencies (250 - 400 Hz) to those of *A. japonicus*  
396 (Parsons, 2010, Parsons *et al.*, 2006). *P. diacanthus* caught in a recent study around  
397 the Northern Territory were of comparable length to the Swan River *A. japonicus*  
398 (Phelan, 2008). However, sampling conducted in Cape York Peninsula waters in  
399 1999 and 2000 yielded predominantly 750-799 and 600-649 mm specimens,  
400 respectively (Phelan *et al.*, 2008). It is not known whether the sounds reported by  
401 Cato (1980) and McCauley (2001) originated from similar sized fish.

402

403 Each study has shown significant variability in the estimated SLs. Regardless, the *A.*  
404 *japonicus* SLs are greater than those of previously reported species, as are the SLs of  
405 other similar sized Sciaenidae, where the call structure is similar (Cato, 1980;  
406 McCauley, 2001). While the two previous reports did not test as many samples as this  
407 study, the difference in SL is large and the authors suggest several possible reasons  
408 for this:

409

410 (1) The sizes of *P. diacanthus* callers were unknown and if of considerably smaller  
411 size may contribute to the intensity difference;

412

413 (2) Previous reports of fish SLs have estimated transmission losses at  $20\log_{10}(r)$ , in  
414 contrast with this study where transmission losses were calculated through best fit  
415 regression and found to be greater than spherical spreading (Table III). The  
416 substitution of  $20\log_{10}(r)$  for transmission loss reduced the *A. japonicus* SL for  
417 categories 1, 2 and 3 to 153, 165 and 156 dB re  $1\mu\text{Pa}$  at 1 m, respectively (Table II).  
418 If the SEL is calculated using a  $20\log_{10}(r)$  transmission loss the determined SL  
419 becomes  $162 \pm 2.7$  dB re  $1\mu\text{Pa}^2\cdot\text{s}$  at 1 m, more comparable with the 150 dB re  $1\mu\text{Pa}^2\cdot\text{s}$   
420 at 1 m reported by McCauley (2001) for the speculated *P. diacanthus*;

421

422 (3) The most comparable sound between those of *A. japonicus* above and those  
423 reported for *P. diacanthus* differ in length and structure. McCauley (2001) noted that  
424 the *P. diacanthus* calls persisted for longer than a second, much longer than the 0.35-s  
425 *A. japonicus* calls. The amount of energy required to produce calls increases with  
426 duration for the same SPL (Mitchell *et al.*, 2008). Hence for the same amount of  
427 energy use, shorter sounds would have higher SLs. The *P. diacanthus* calls reduced  
428 in amplitude in the latter half of the call (McCauley, 2001), similar to those of the  
429 Atlantic cod (*Gadhus morhua*). Nilsson (2004) attributed this reduction to buildup of  
430 lactic acid in the sonic muscle. So the greater call length and possible lactic acid  
431 buildup are suggested to account for the lower SL.

432

433 This study has provided some observations of *in situ* fish call directivity. The  
434 estimated SL of Category 3 calls recorded as the fish passed the hydrophone displayed

435 a 5 to 8.5 dB difference between orientations. If the assumption of forward motion is  
436 correct, and the individual was not swimming against a current drifting backwards  
437 past the hydrophone, calls were greater in front of the caller than behind. This finding  
438 is in contrast with Barimo and Fine (1998), who hypothesized that directivity in sound  
439 production of *O. tau* would be reduced in the direction of the fish ears and observed a  
440 3 to 5 dB re 1 $\mu$ Pa decline in SL moving from the rear to the front of the fish. The  
441 reason for this difference is unknown, though if the *A. japonicus* Category 3 caller  
442 was swimming into, and drifting backwards with the current the directivity would  
443 result in a reduced SPL in front of the fish. Although the sample size of different  
444 Category 3 calls is small and the variance large (in part due to the small sample size),  
445 the differences in SL between orientation is notable. This would not be expected  
446 given that the wavelength of each call is large compared with the range to the  
447 receiver, but is a phenomenon similar to that reported by Barimo and Fine (1998).

448

449 Although *A. japonicus* calls contain energy across the frequency bandwidth from 50  
450 to 1000 Hz, the tone burst dominates the call such that often one or more of the  
451 spectral peak frequencies are 10 dB re 1 $\mu$ Pa greater than other amplitude modulated  
452 frequencies. As a result ray multi-path interference was observed in the SLs (Figure  
453 6). This was observed to be more prominent at the farther regions of the test ranges;  
454 however, this may be due to the cluster of calls emitted by one fish rather than close  
455 range measurements where the effect was masked by the variability of different  
456 callers.

457

458 Source level relative to background noise is a significant factor in determining the  
459 maximum range at which a call can be detected by its intended recipient. Table III

460 has shown some coarse estimates of maximum range at which a call would have  
461 attenuated to background noise level based purely on a) the transmission losses  
462 estimated in this study and b) transmission losses estimated using spherical spreading.  
463 It does not use statistical analysis of the probability of signal detection, frequency  
464 bandwidths used by fish when detecting sounds or account for fish hearing critical  
465 thresholds and so could not be used as an estimation of the ranges over which fish  
466 may detect calls. To assess such communication ranges requires considerable effort  
467 and has been deemed future work. However, from a management perspective such  
468 detection ranges may have specific impacts on the spawning success rates of an  
469 aggregation. Wind driven waves can also produce background noise at a level similar  
470 to that of passing vessels and it is possible that the fish have adapted to continue their  
471 behavior despite increased noise levels. However, waves which would produce noise  
472 of that level are rare compared with the vessel presence in Mosman Bay.

473

474 *A. japonicus* in Mosman Bay form aggregations of low density calling by comparison  
475 with other reports (Sprague and Luczkovich, 2004). Thus individuals can often be  
476 detected at comparatively long ranges (Parsons *et al.*, 2009). During periods of low  
477 background noise, such as those reported here, regression models predict a signal  
478 processing detection range of approximately 120, 400 and 110 m for calls of  
479 Categories 1, 2 and 3 respectively. However, periods of high background vessel noise  
480 were frequent at the study site. Vessel noise of water skiers passing directly above  
481 calling fish has reportedly masked calls at distances of less than the water depth  
482 (Parsons *et al.*, 2009) and received vessel noise greater than 150 dB re 1 $\mu$ Pa would  
483 mask all calls at distances of near 1 m (assuming spherical spreading). The ratio of  
484 detection ranges between those identified with and without the noise of a passing

485 vessel (Table III) are comparable and show that, even under near ideal detection  
486 conditions, anthropogenic activity can significantly reduce detection ranges of such  
487 calls. Anthropogenic noise therefore poses a significant impact which needs assessing  
488 to determine what, if any, effect it has on spawning success.

489

## 490 **V CONCLUSIONS AND RECOMMENDATIONS**

491

492 To utilize the SL of *A. japonicus* calls is to begin the process of absolute biomass  
493 estimates. Hydrophone array studies have provided an insight into separation  
494 distances and distributions of calling *A. japonicus* in Mosman Bay. Proposed future  
495 work involves the combination of the determined SL ranges and distributions with  
496 call counting techniques to estimate the number of calling *A. japonicus* within the  
497 hydrophone detection range. The issue of determining the ratio of calling/non-calling  
498 fish remains to be resolved. In an ideal world this would involve an isolated  
499 aggregation of randomly distributed sized males and females of known number. Thus  
500 the number of calls could be related to the overall population. In the absence of this  
501 possibility, ratios will have to be repeatedly obtained from *aquaria*.

502

503 The accurate determination of ranges at which fish can detect calls is an important  
504 step towards assessing the impact of anthropogenic noise on communication. This  
505 requires statistical analysis of the probability of signal detection and an evaluation of  
506 the critical hearing ratio of the fish involved. Ranges given in this paper are only an  
507 identification that, at the SLs of *A. japonicus* calls in Mosman Bay, passing vessels  
508 can reduce detection, even using advanced signal processing, from hundreds of meters  
509 to less than 10 m.

510

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512

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521

522 **VI REFERENCES**

523

524 Barimo, J. F. and Fine, M. L. (1998). "Relationship of swim-bladder shape to the  
525 directionality pattern of underwater sound in the oyster toadfish," *Can. J. Zool.* **76**,  
526 134-43.

527

528 Cato, D. H. (1980). "Some unusual sounds of apparent biological origin responsible  
529 for sustained background noise in the Timor Sea," *J. Acoust. Soc. Am.* **68**, 1056-60.

530

531 Cato, D. H. (1998). "Simple methods of estimating source levels and locations of  
532 marine animal sounds," *J. Acoust. Soc. Am.* **104**, 1667-78.

533

534 Coates, R. F. W. (1990). Underwater Acoustic Systems, John Wiley & Sons Inc.,  
535 New York, 188 pp.  
536

537 Connaughton, M. A., Taylor, M. H. and Fine, M. L. (2000). "Effects of fish size and  
538 temperature on weakfish disturbance calls: Implications for the mechanism of sound  
539 generation," J. Exp. Mar. Biol. Ecol. **203**, 1503-12.  
540

541 Farmer, B. M. (2008). Comparisons of the biological and genetic characteristics of the  
542 Mulloway *Argyrosomus japonicus* (Sciaenidae) in different regions of Western  
543 Australia, PhD Thesis, Murdoch University. 216 pp.  
544

545 Fine, M. L., Winn, H. E. and Olla, B. L. (1977). '*Communication in fishes*', in TA  
546 Sebeok (ed.), How animals communicate, Indiana University Press, Bloomington. pp.  
547 472-518.  
548

549 Lagardere, J. P. and Mariani, A. (2006). "Spawning sounds in the meagre  
550 *Argyrosomus regius* recorded in the Gironde Estuary, France," J. Fish Biol. **69**, 1697-  
551 702.  
552

553 Luczkovich, J. J., Sprague, M. W., Johnson, S. E. and Pullinger, C. (1999).  
554 "Delimiting spawning areas of weakfish *Cynoscion regalis* (family sciaenidae) in  
555 Pamlico Sound, North Carolina using passive hydroacoustic surveys," Bioacoustics,  
556 **10**, 143-60.  
557

558 Malme, C. I., Smith, P. W. J. and Miles, P. R. (1986). Characterisation of geophysical  
559 acoustic survey sounds' OCS Study MMS-86-0032 Prepared by BBN Laboratories  
560 Inc., Cambridge, for Battelle Memorial Institute to the Department of the Interior,  
561 Mineral Management Service, Pacific Outer Continental Shelf Region, Los Angeles.  
562

563 McCauley, R. D. (2001). Biological sea noise in northern Australia: Patterns of fish  
564 calling, PhD Thesis, James Cook University of North Queensland. 290 pp.  
565

566 Mitchell, S., Poland, J. & Fine, M.L. (2008). "Does muscle fatigue limit  
567 advertisement calling in the oyster toadfish *Opsanus tau*?" Ani. Behav. 76, 3, 1011-  
568 1016.  
569

570 Myrberg, A. A. J. & Spires, J. Y. (1972). "Sound discrimination by the bicolor  
571 damselfish (*Eupomacentrus Partitus*)," J. Exp. Mar. Biol. Ecol. 57, 727-735.  
572

573 Nilsson, J. (2004). "Acoustic behaviour of spawning cod (*Gadus morhua*),"  
574 Candidatus scientiarum Thesis, University of Bergen, Bergen, Norway. 135 pp.  
575

576 Parsons, M. J. G. (2010). Passive acoustic techniques for monitoring fish  
577 aggregations, In "An investigation into active and passive acoustic techniques to study  
578 aggregating fish species," PhD Thesis, Curtin University, pp. 131-297.  
579

580 Parsons, M. J. G., McCauley, R. D. and Mackie, M. C. (2006). "Spawning sounds of  
581 the mulloway (*Argyrosomus japonicus*)," Noise in Progress, First Australasian



582 Acoustical Societies Conference, Christchurch, New Zealand. November 26<sup>th</sup> -29<sup>th</sup>.  
583 pp. 401-6.

584

585 Parsons, M. J. G., McCauley, R. D., Mackie, M. C., Siwabessy, P. J. and Duncan, A.  
586 J. (2009). "Localisation of an individual mullock (*Argyrosomus japonicus*) within a  
587 spawning aggregation and their behavioural characteristics throughout a diel  
588 spawning period," ICES J. Mar. Sci. **66**, 1007-14.

589

590 Parsons, M. J. G., McCauley, R. D., Mackie, M. C., and Duncan, A. J. (2010).  
591 "Ranging close proximity mullock (*Argyrosomus japonicus*) calls with a single  
592 hydrophone," Acoust. Australia, **39**, 95-101

593

594 Phelan, M. (2008). Assessment of the implications of target fishing on black jewfish  
595 (*Protonibea diacanthus*) aggregations in the Northern Territory, Fisheries Research  
596 and Development Corporation, FRDC Report No. 2004/004. No. 91. 115 pp.

597

598 Phelan, M., Gribble, N. A. and Garrett, R. N. (2008). "Fishery biology and  
599 management of *Protonibea diacanthus* (Sciaenidae) aggregations in far Northern  
600 Cape York Peninsula waters," Cont. Shelf Res. **28**, 2143-51.

601

602 Robins, C. R. and Ray, G. C., (1986). A field guide to Atlantic coast fishes of North  
603 America. Houghton Mifflin Company, Boston, U.S.A. 354 p.

604

605 Southall, B. L., Bowles, A. E., Ellison, W. T., Finneran, J. J., Gentry, R. L., Greene  
606 Jr., C. R., Kastak, D., Ketten, D. R., Miller, J. H., Nachtigall, P. E., Richardson, W. J.,

607 Thomas, J. A. and Tyack., P. L. (2007). "Marine mammal noise exposure criteria:  
608 Initial Scientific Recommendations," *Aquat. Mam.*, **33**, 415-77.  
609

610 Sprague, M. W. and Luczkovich, J. J. (2002). "Quantifying Species-Specific  
611 Contributions to the Overall Sound Level," An International Workshop on the  
612 Applications of Passive Acoustics in Fisheries, Massachusetts Institute of  
613 Technology, Cambridge, MA, pp. 147-53.  
614

615 Sprague, M. W. and Luczkovich, J. J. (2004). "Measurement of an individual silver  
616 perch *Bairdiella chrysoura* sound pressure level in a field recording," *J. Acoust. Soc.*  
617 *Am.* **116**, 3186-91.  
618

619 Ueng, J. P., Huang, B. Q. and Mok, H. K. (2007). "Sexual differences in the spawning  
620 sounds of the Japanese Croaker *Argyrosomus japonicus* (Sciaenidae)," *Zool. Stud.*,  
621 **46**, 103-10.  
622

623 Urick, R. J. (1983). *Principles of underwater sound*, 3rd edn, MacGraw-Hill, New  
624 York. 451 pp.  
625

626 Walpole, R. E. and Myers, R. H. (1985). '*Linear regression and correlation*', in  
627 *Probability and Statistics for Engineers and Scientists*, third edition, Macmillan  
628 Publishing Company, New York. pp. 315-36.  
629

630 Winn, H. E. (1964). '*The biological significance of fish sounds*', in WN Tavolga (ed.),  
631 *Marine Bioacoustics*, vol. 2, Pergamon Press, Sydney. pp. 213-31.

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634

635 Table I. Extrapolated call sources levels for each category of *A. japonicus* call  
636 from least squares linear regression. Values display SPL source levels and equivalent  
637 spreading losses together with 95% confidence limits and the curve correlation with  
638 data points.

Call Category	Orientation	Number Calls	Source Level (dB re 1 $\mu$ Pa) ( $\pm$ 95% confidence limits)	Transmission Loss (log(r)) ( $\pm$ 95% confidence limits)	$R^2$	
Category 1	All	N/A	53	163 (148, 179)	-25.39 (-35, -16)	0.42
Category 2	Individual	N/A	65	172 (163, 180)	-23.94 (-30, -17)	0.61
	All	N/A	112	172 (168, 176)	-23.74 (-26, -22)	0.82
	All	N/A	28	157 (152, 162)	-23.04 (-27, -19)	0.88
Category 3	One pulse	Towards	7	156 (151, 162)	-18.67 (-26, -11)	0.89
		Away	4	152 (144, 159)	-19.17 (-27, -11)	0.98
	Two pulse	Towards	3	163 (98, 227)	-27.53 (-102, 47)	0.96
		Away	10	154 (150, 158)	-18.81 (-24, -14)	0.93

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653 Table II. Values of source levels with standard deviation, based on recorded  
 654 values. Source levels (dB re 1 $\mu$ Pa at 1 m) using  $20\log_{(10)}(r)$  losses are shown. Data is  
 655 also for 3 type of source level as they are often reported (SPL, SEL equivalent energy  
 656 and peak-to-peak pressures). For each method and call category the calculated source  
 657 level, transmission loss curve constant and correlation coefficient are shown. Mean  
 658 call lengths for each category are also shown.  
 659

		Call Category		
		1	2	3
Source level (dB re 1 $\mu$ Pa)		153 (6)	165 (2)	156 (4)
20log(r) transmission loss (s.d.)				
<b>Sound</b>	Source level (95% c.l.)	163 (148, 179)	172 (168, 176)	157 (152, 162)
<b>pressure level</b>	Transmission loss			
<b>(dB re 1<math>\mu</math>Pa at</b>	(log <sub>(10)</sub> (r)) (95% c.l.)	-25.4 (-34.6, -16.2)	-23.7 (-25.9, -21.6)	-23.0 (-26.6, -19.5)
<b>1 m)</b>	$R^2$	(0.42)	(0.82)	(0.88)
<b>SEL</b>	Source level (95% c.l.)	152 (138, 166)	165 (156, 173)	136 (132, 139)
<b>(dB re 1<math>\mu</math>Pa<sup>2</sup>s</b>	Transmission loss			
<b>at 1 m)</b>	(log <sub>(10)</sub> (r)) (95% c.l.)	-22.9 (-31.1, -14.6)	-21.8 (-27.2, -16.5)	-17.4 (-21.2, -13.5)
	$R^2$	(0.64)	(0.64)	(0.74)
<b>Peak-peak</b>	Source level (95% c.l.)	183 (173, 195)	194 (189, 201)	167 (165, 170)
<b>pressure</b>	Transmission loss			
<b>(dB re 1<math>\mu</math>Pa at</b>	(log <sub>(10)</sub> (r)) (95% c.l.)	-25.2 (-31.7, -18.6)	-27.2 (-30.8, -23.6)	-16.1 (-18.8, -13.5)
<b>1 m)</b>	$R^2$	(0.77)	(0.86)	(0.83)
Mean call length (s) (s.d.)		0.054 (0.021)	0.346 (0.063)	0.018 (0.015)

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665 Table III. Maximum detection ranges (r) for all call categories (using basic signal  
666 processing) when emitted during two levels of background noise (110 and 150 dB re  
667 1 $\mu$ Pa), calculated by assuming the regression calculated TL determined from the data  
668 and losses due only to spherical spreading. Call/vessel noise energy was computed  
669 over broadband spectra. As this calculation does not account for critical hearing  
670 ratios, frequency bandwidths used by *A. japonicus* to detect calls and no probability of  
671 signal detection has been applied, this is a simple calculation of the range at which the  
672 signal attenuates to background noise levels.

Detection Range (m)

Call Category	Source level (dB re 1 $\mu$ Pa at 1 m)	Regression calculated		Spherical transmission	
		transmission loss and noise		loss and noise levels	
		levels (dB re 1 $\mu$ Pa)		(dB re 1 $\mu$ Pa)	
		110	150	110	150
<b>1</b>	<b>163</b>	<b>123</b>	<b>3</b>	<b>231</b>	<b>2</b>
<b>2</b>	<b>172</b>	<b>396</b>	<b>8</b>	<b>660</b>	<b>7</b>
<b>3</b>	<b>157</b>	<b>112</b>	<b>2</b>	<b>117</b>	<b>1</b>

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681 Figure 1. Map of Western Australia with a magnification of the Swan River and  
682 Mosman Bay, highlighting the location of the hydrophone array.

683

684 Figure 2. Spectrogram of 20 seconds of *A. japonicus* call period from a  
685 hydrophone positioned on the riverbed in 18 m of flat water. Individual *A. japonicus*  
686 calls (grey-black areas) are clearly visible above minimum background noise (white  
687 areas). Spectrum produced using a 1024 point Hanning window with 0.5 overlap and  
688 frequency resolution of 5.09 Hz.

689

690 Figure 3. Example wave forms of Category 1 short calls (A), Category 2 long  
691 calls (B) and single pulse Category 3 calls (C) as recorded by a 'bottomed'  
692 hydrophone.

693

694 Figure 4. Example sound pressure level (dB re 1 $\mu$ Pa) calculation of an *A.*  
695 *japonicus* Category 2 long call. A) Wave form of example call with noise removal  
696 zone 500 samples before (circles) and after (squares) shown. Crosshairs mark the  
697 points at which 5 and 95% of the total energy occurs within the analyzed region. B)  
698 Cumulative energy of the call showing pressure squared per second with 5 and 95%  
699 region markers shown. C) Power spectral density of call.

700

701 Figure 5. Detected sound pressure levels (dB re 1 $\mu$ Pa) with range (log scale) for  
702 53 Category 1 calls. Continuous line illustrates linear regression model of  
703 transmission losses with 95% confidence limits of source level shown (dotted lines).

704

705 Figure 6. Detected sound pressure levels (dB re 1 $\mu$ Pa) with range (log scale) for  
706 Category 2 calls. Calls of a tracked individual fish ( $\circ$ ) and those of all remaining fish  
707 ( $\times$ ) are shown. Continuous line marks the linear regression determined transmission  
708 losses with 95% confidence limits (dotted lines).

709

710 Figure 7. Time of fish calls with  $\log_{10}(r)$  (A) highlighting the order of 1 pulse ( $\circ$   
711 and  $\square$ ) and 2 pulse ( $\times$  and  $+$ ) Category 3 calls. Sound pressure levels (dB re 1 $\mu$ Pa)  
712 against range (log scale) as detected by the bottomed hydrophone (B) as the fish  
713 approached ( $\square$  and  $+$ , dashed line) and then passed ( $\circ$  and  $\times$ , dot-dashed line) the  
714 hydrophone. The positions of the  $\circ$ ,  $\square$ ,  $\times$  and  $+$  illustrate whether the fish was  
715 orientated towards or away from the hydrophone and whether the call was a single or  
716 double pulse call. Least squares regression curve, together with 95% confidence limits  
717 are shown by the continuous and dotted lines respectively. The order of calls is  
718 indicated by arrows. Calls not suitable for range analysis have been omitted.

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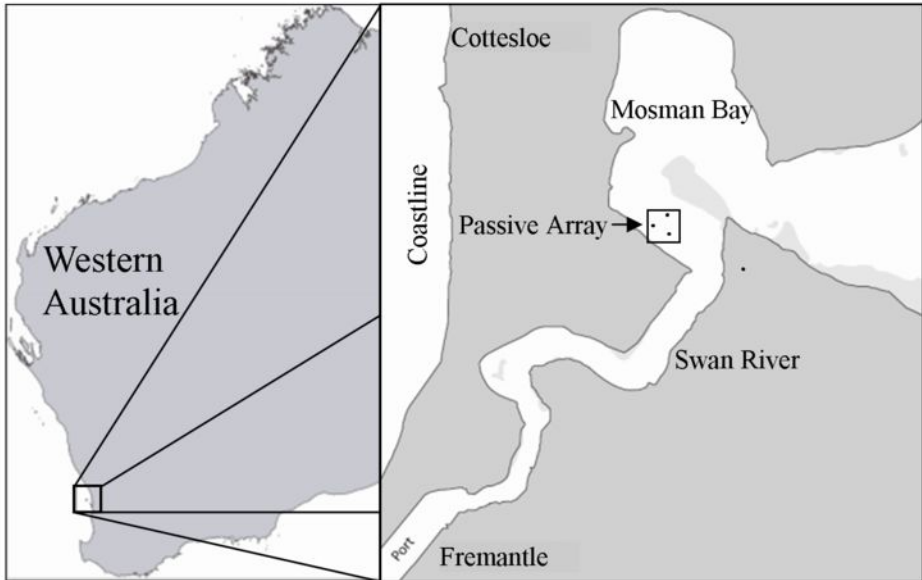
722 Figure 8. Distribution of source levels (dB re 1 $\mu$ Pa at 1 m) from recorded sound  
723 pressure levels for each *A. japonicus* call category based on the estimated  
724 transmission losses only.

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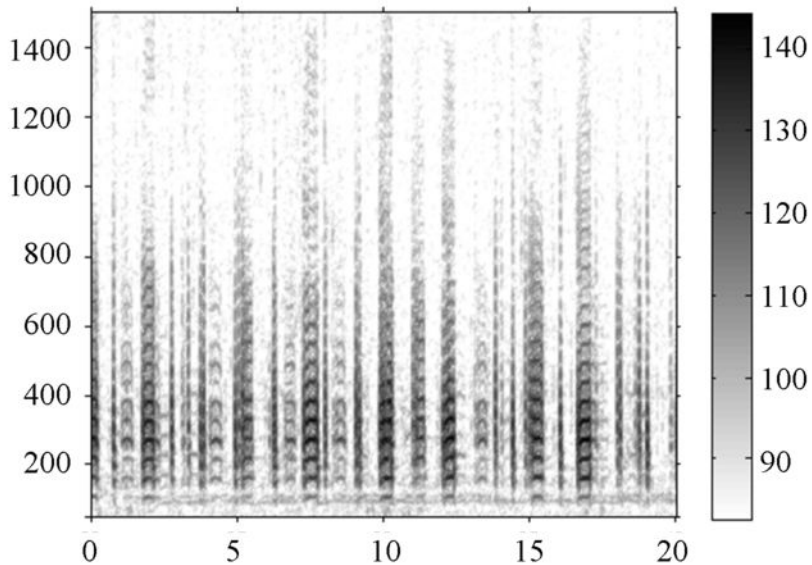
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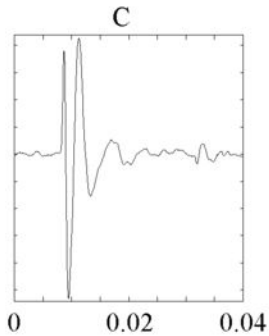
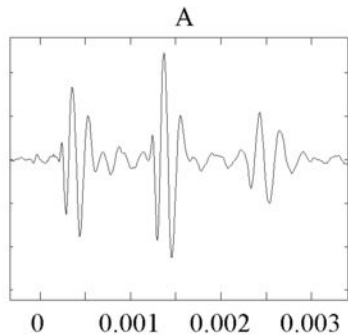
Frequency (Hz)



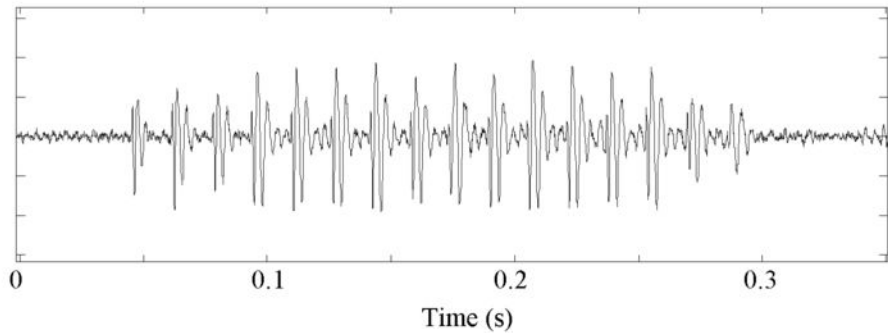
Time (s)

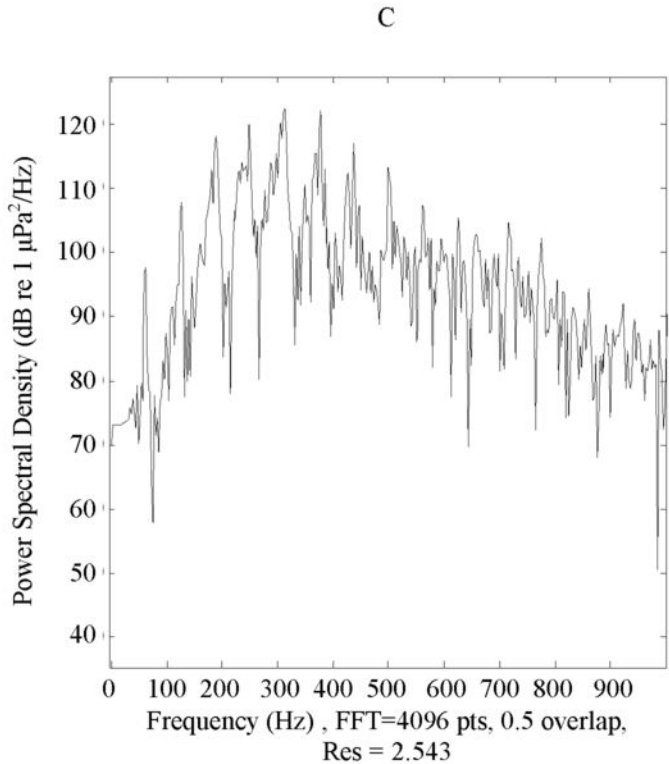
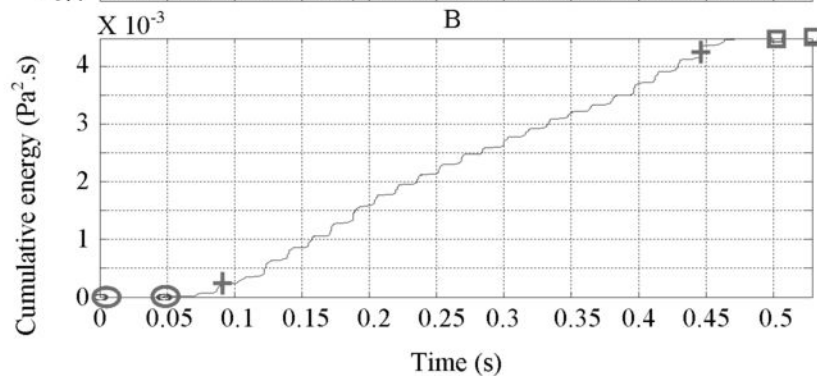
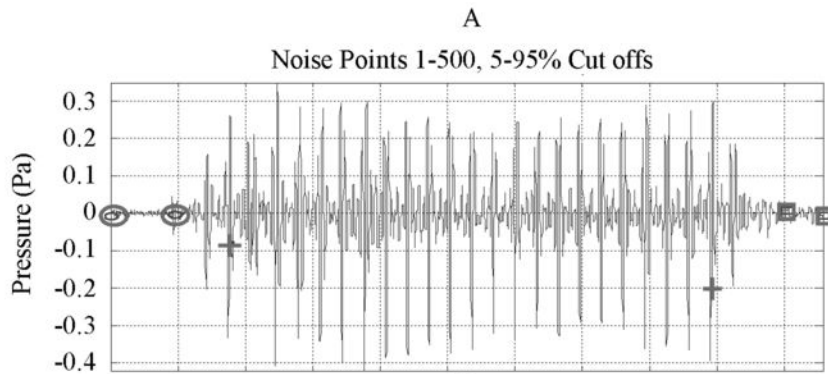
Power spectral density (dB re  $1 \mu\text{Pa}^2/\text{Hz}$ )

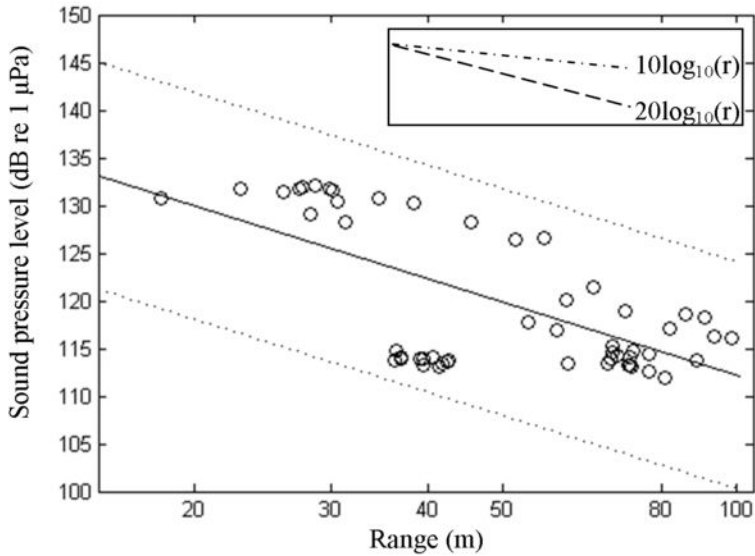
Relative voltage (V)

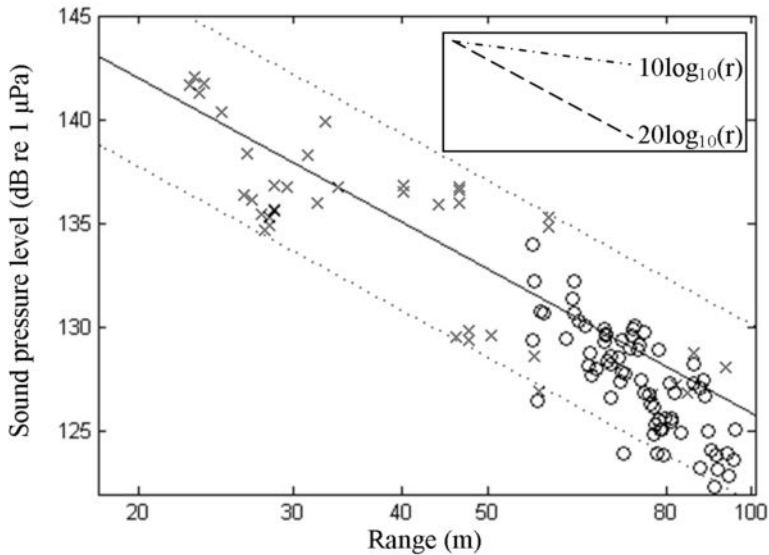


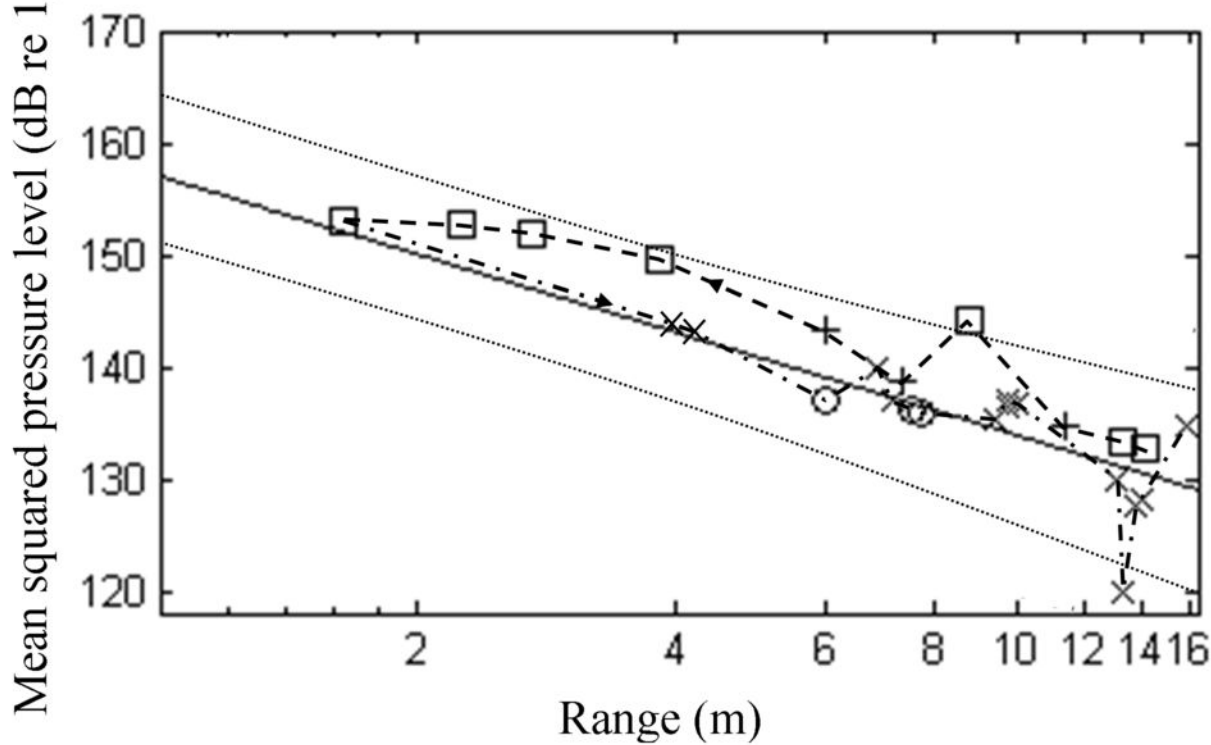
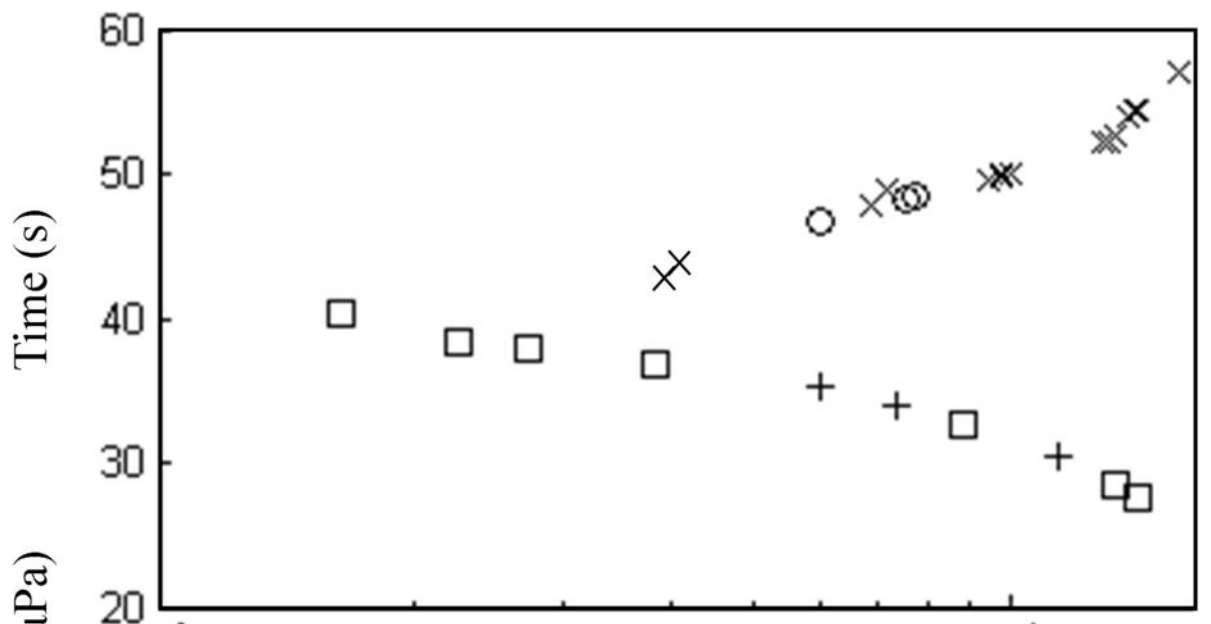
**B**



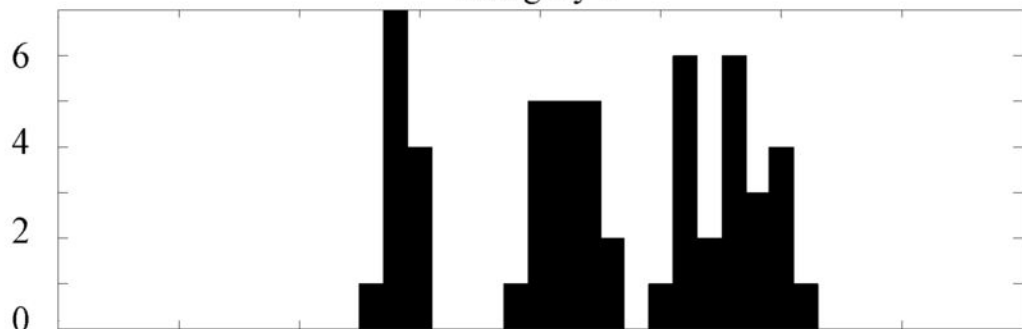




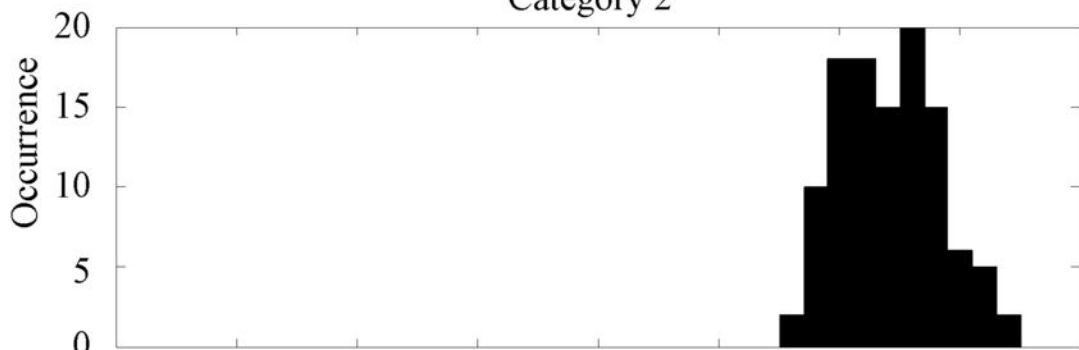




Category 1



Category 2



Category 3

