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Abstract: Telemetry investigations to gather essential information about fish migrations are strongly reliant on the behaviour, energetics, condition and survival of the animals being unaltered by the tagging procedure. Twaite shad (*Alosa fallax* Lacépède; 'shad') is a threatened clupeid fish for which there is a considerable knowledge gap on their anadromous movements. They are also reported to be sensitive to handling and anaesthesia, resulting in practical difficulties in tag implantation; previous investigations externally attached tags without sedation. The aim of this study was to incrementally refine the acoustic-tagging protocol for shad via application of a previously un-tried anaesthetic (i.e. tricaine methanesulphonate (MS-222)) and by surgical implantation of the tag in the peritoneal cavity. All captured shad ($n = 25$) survived handling, anaesthesia and tagging, and were detected moving upstream after release. Surgically implantation ($n = 5$) was significantly faster than externally mounting the tag ($n = 20$) and time to recover was similar. Total upstream movement, total movement, residence time in receiver array and speed of upstream movement were statistically similar for externally and internally tagged fish. Post-spawning, a large proportion (68 %) of tagged fish returned to the estuary, downstream of the receiver array. Internal tagging under anaesthesia is recommended for studying anadromous movements of shad, given welfare benefits during surgery and once at liberty, thus increasing the likelihood of tagged fish performing natural behaviours. Further, implantation of tags programmed to last many years enables multiple spawning migrations by the same individuals to be studied, which would lead to substantial advances in ecological knowledge and potentially reduce the total number of tagged fish.

1 **Refinement of acoustic-tagging protocol for twaite shad *Alosa fallax* (Lacépède), a**
2 **species sensitive to handling and sedation**

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9 **ABSTRACT**

10 Telemetry investigations to gather essential information about fish migrations are reliant on
11 the behaviour, condition and survival of the animals being unaltered by the tagging
12 procedure. Twaite shad (*Alosa fallax* Lacépède; 'shad') is a threatened clupeid fish for which
13 there is a considerable knowledge gap on their anadromous movements. They are also
14 reported to be sensitive to handling and anaesthesia, resulting in practical difficulties in tag
15 implantation; previous investigations externally attached tags without sedation. The aim of
16 this study was to incrementally refine the acoustic-tagging protocol for shad *via* application
17 of a previously un-tried anaesthetic (i.e. tricaine methanesulphonate (MS-222)) and by
18 surgical implantation of the tag in the peritoneal cavity. All captured shad ($n = 25$) survived
19 handling, anaesthesia and tagging, and were detected moving upstream after release.
20 Surgically implantation ($n = 5$) was significantly faster than externally mounting the tag ($n =$
21 20) and time to recover was similar. Total upstream movement, total movement, residence
22 time in receiver array and speed of upstream movement were statistically similar for
23 externally and internally tagged fish. Post-spawning, a large proportion (68 %) of tagged fish
24 returned to the estuary, downstream of the receiver array. Internal tagging under
25 anaesthesia is recommended for studying anadromous movements of shad, given welfare
26 benefits during surgery and once at liberty, thus increasing the likelihood of tagged fish
27 performing natural behaviours. Further, implantation of tags programmed to last many years
28 enables multiple spawning migrations by the same individuals to be studied, which would

29 lead to substantial advances in ecological knowledge and potentially reduce the number of
30 fish tagged.

31 *Keywords*

32 Anadromous; Animal welfare; Iteroparous; Regulated procedure; Surgical implantation;
33 Telemetry

34 **1. Introduction**

35 Fish telemetry investigations are routinely performed to gather essential information about
36 migrations, habitat use, predator–prey interactions and responses to anthropogenic impacts,
37 to help protect species and the environments they inhabit (Hussey et al., 2015). Such
38 studies are reliant on the behaviour, condition and survival of the animals being unaltered by
39 the tagging procedure (Cooke et al., 2013). This has resulted in a considerable amount of
40 work to identify maximum tag burden, optimal tag implantation location and most appropriate
41 methods of anaesthesia (Broadhurst et al., 2009; Ross & Ross, 2009). There have been
42 considerable refinements in internal tagging procedures, with tags often retained for the
43 lifetime of the fish with minimal long-term impact (Jepsen et al., 2002; Bridger and Booth,
44 2003; Cooke et al., 2011). External tag attachment remains important in some studies and
45 species, including those considered to be sensitive to handling (Jepsen et al., 2015;
46 Johnson et al., 2015). However, tags can become fouled, increase drag during swimming,
47 cause irritation and harm as the fish grow, potentially impairing individual behaviour and
48 increasing mortality risk (Mulcahy, 2003; Cooke et al., 2013; Jepsen et al., 2015).

49 Twaité shad *Alosa fallax* (Lacépède) ('shad' hereafter) is an anadromous clupeid fish
50 species that was once abundant and widespread across Europe (Arahamian et al., 2003).
51 Their populations have, however, declined considerably in the last century. Causal factors
52 relate primarily to anthropogenic disturbances, especially the construction of weirs in the
53 lower reaches of rivers that reduce access to spawning areas (Jolly et al., 2012). The
54 species is listed on Appendix III of the Bern Convention and Annexes II and V of the EU
55 Habitats Directive. Despite their conservation importance, their anadromous spawning

56 migration remains under-studied primarily due to difficulties tagging shad, a species reported
57 to adversely react to handling and sedation (with 2-phenoxyethanol) that results in high
58 mortality rates (Rooney and King, 2014; Breine et al., 2017). To overcome these challenges,
59 recent investigations have externally mounted acoustic tags without sedation because it is
60 less invasive and thought to be quicker than surgical implantation (Rooney and King, 2014;
61 Breine et al., 2017). Although these studies were successful, Breine et al. (2017)
62 recommended further research on the effects of anaesthesia, handling and tagging on shad.

63 The aim of this study was to refine the acoustic-tagging protocol for shad, giving due
64 consideration to their sensitivity to handling and sedation, to provide short-term welfare
65 benefits during surgery and long-term welfare benefits while at liberty, thus enabling
66 expression of natural behaviours. Objectives were to: (1) refine the external tag attachment
67 protocol of Breine et al. (2017) *via* application of previously un-tried anaesthetic (i.e. tricaine
68 methanesulphonate (MS-222)); (2) further refine the procedure by surgically implanting the
69 tag within the peritoneal cavity; and (3) quantify the impacts of the tagging methods through
70 comparison of shad movement. As shad are iteroparous and, potentially, philopatric (King
71 and Roche, 2008), implantation of tags programmed to last many years enables multiple
72 spawning migrations by the same individuals to be studied, which would lead to substantial
73 advances in ecological knowledge.

74 **2. Methods**

75 *2.1. Fish capture and iterative tagging process*

76 The refinement of the shad tagging protocol was completed during the 2017 shad spawning
77 migration in the River Severn, Western England (Fig. 1). Twenty-five shad were captured
78 from two locations, downstream of Maisemore ($n = 8$) and Upper Lode weirs ($n = 17$), with
79 23 captured by angling (small lure with single barbless hook) and two with a seine net (30-m
80 long, 2-m deep and 10-mm mesh) (Table 1). Tagging was an iterative process involving
81 small batches of fish to minimise the number of fish with compromised welfare if tagging was
82 unsuccessful and to enable refinements between batches. Thus, the initial 3 captured fish
83 were externally tagged under general anaesthesia (batch 1), with tagging only

84 recommencing once a receiver 14.8-km upstream of the release location revealed the fish
 85 had recovered sufficiently to continue their upstream movement. The decision to commence
 86 surgically implanting tags in the body cavity (batch 4) was only taken after a further 11 shad
 87 had been successfully tagged externally (batch 2 and 3). The final six fish (batch 5) were
 88 tagged externally because there was no opportunity to establish if the internally tagged fish
 89 (batch 4) had been detected on the receiver upstream of the release location.

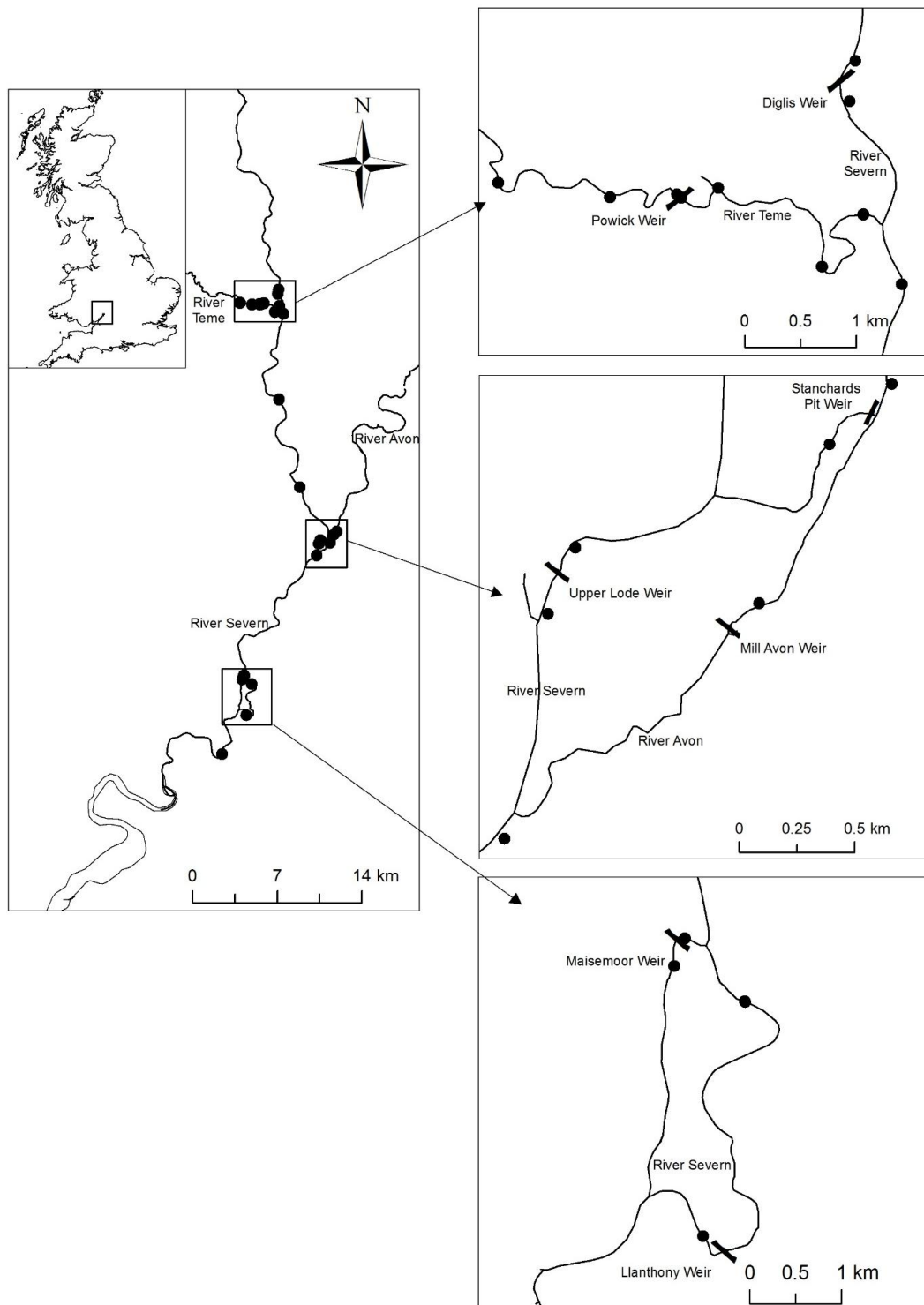
90 Table 1. Capture date, sample size and capture, release (DS = downstream, US = upstream)
 91 and tag locations of twaite shad tagged in five batches on the River Severn.

Batch	Date	<i>n</i>	Capture location	Release location	Tag location
1	11/5/17	3	DS Maisemore Weir	US Maisemore Weir	External
2	17/5/17	5	DS Upper Lode Weir	US Upper Lode Weir	External
3	17/5/17	6	DS Upper Lode Weir	DS Upper Lode Weir	External
4	22/5/17	5	DS Maisemore Weir	US Maisemore Weir	Internal
5	31/5/17	6	DS Upper Lode Weir	DS Upper Lode Weir	External

92

93 *2.2. External and internal tagging procedures*

94 Prior to tagging, acoustic tags (20-mm long x 7-mm diameter (V7), 1.6-g weight in air and
 95 29-mm long x 9-mm diameter (V9), 4.7-g weight in air; www.vemco.com) were activated and
 96 tested with a hand-held detector to verify they were transmitting; weight in air did not exceed
 97 2% of fish mass. Following capture, fish were briefly held in water filled containers (100 L)
 98 prior to their general anaesthesia (MS-222; 0.4-g per 10-L of water). All fish were inspected
 99 for signs of pre-existing injury and disease; no captured fish were excluded from tagging.
 100 Whilst being sedated, the fish were measured (fork length, nearest mm; mean \pm S.D.: 354 \pm
 101 37 mm, range = 302–420 mm), and scale sample and a fin biopsy taken (for use in
 102 complementary studies). The influence of the anaesthetic was visually assessed using body,
 103 opercula and eye movements, with fish only removed following their lack of a response to
 104 touch, loss of ability to balance and the cessation of pectoral fin and eye movements.



105
 106 Figure 1. A map of acoustic receiver locations (black dots) in the River Severn catchment,
 107 including impediments to fish migration (black lines). Maisemore and Llanthony weirs
 108 represent the tidal limit, and Maisemore and Upper Lode weirs were capture locations.

109 Externally mounted tags were attached using surgical thread (Ethilon) passed
 110 through the dorsal musculature using hollow needles and held in place using a rubber plate

111 and aluminium sleeves (as per Breine et al., 2017). Surgically implanted tags were
112 disinfected with providone-iodine and rinsed with saline solution before being implanted into
113 the body cavity through a ventro-lateral incision made with a scalpel, anterior to the muscle
114 bed of the pelvic fins. The incision was closed with an absorbable monofilament suture. Fish
115 were held in a clean V-shaped foam support and their eyes were covered with a damp cloth
116 during surgery. All fish were treated in compliance with the UK ASPA (1986) Home Office
117 licence number PPL 60/4400.

118 After surgery, fish were transferred to a damp sling for weighing (to 25 g; mean \pm
119 S.D. = 547 ± 173 g, range = 300–850 g) and then returned to the river, being held whilst they
120 orientated towards the flow and were only released when they had regained balance, body
121 reflexes and swimming ability. This was considered preferable to holding fish in tanks with
122 water circulation and aeration, as shad have been recorded to die during transportation and
123 at fish farms (Clough et al., 2004). Fish were released upstream of Maisemore Weir ($n = 8$),
124 downstream of Upper Lode Weir ($n = 12$) and upstream of Upper Lode Weir ($n = 5$) as part
125 of the wider investigation (Table 1). Catchment-wide migration was examined using 23
126 strategically located acoustic receivers (Vemco; www.vemco.com) (Fig. 1); no fish were
127 detected on the most upstream receivers.

128 2.3. *Data analysis*

129 Time taken for anaesthesia, surgery and recovery when externally and internally tagging
130 shad was compared using t-tests (non-normal data (Shapiro test) were log-transformed). It
131 was not possible to recapture tagged shad to assess general health and condition, external
132 tag fouling or healing of incisions for internally implanted tags. Instead, movements of fish in
133 the river were used as evidence that the fish had recovered from handling, anaesthesia and
134 surgery. Specifically, the amount of upstream movement (i.e. sum of all upstream
135 movements), total movement (i.e. sum of all up and downstream movements), and
136 residence time in the receiver array (i.e. number of days from release to first detection on
137 last receiver) were calculated for each fish. In addition, the speed of upstream movement
138 between receivers was calculated (distance between receivers / last detection on upstream

139 receiver – first detection on downstream receiver). The movements of fish in batches 1 and
 140 4, captured and released at the same location but with external and internal tag attachment,
 141 were compared using t-tests (non-normal data (Shapiro test) were log-transformed) to
 142 quantify impacts of the tagging methodology. Both movement and speed metrics represent
 143 minimum estimates, as they are measured at the resolution of receiver separation, thus back
 144 and forth movements between receiver detection area are undetected. The fates of
 145 individual fish were broadly separated into those that returned to the estuary and those that
 146 were assumed to have died in the river, though the latter could not be separated from tag
 147 failure or loss, and the potential cause of death could not be determined (e.g. tagging
 148 induced, natural predation event, tagging-induced predation event or natural mortality after
 149 spawning). Data analysis was performed primarily in Microsoft Excel and statistical
 150 comparisons performed using R statistical software (version 3.4.3, R Core Team 2017), with
 151 movement speed analysis in the V-Track package (Campbell et al., 2012).

152 3. Results

153 All 25 fish caught during the investigation survived capture, handling, sedation and tagging,
 154 and were assessed as being in satisfactory condition prior to be returned to the river. The
 155 time taken for anaesthesia was similar ($t = -0.054171$, d.f. = 5.5144, $P = 0.959$) whereas
 156 internal implantation was significantly faster than external attachment (t -test on logged data;
 157 $t = -88.36$, d.f. = 32.372, $P < 0.001$), both usually within four minutes (Table 2). The mean
 158 time to recover was also similar (t -test on logged data; $t = -1.9709$, d.f. = 7.8191, $P = 0.085$),
 159 and the longest recovery did not exceed six minutes for either treatment group (Table 2).

160 Table 2. Time (seconds; mean \pm 95% C.I. (min.–max.)) taken for anaesthesia, surgery and
 161 recovery when externally and internally tagging shad with acoustic tags.

Procedure stage	External ($n = 20$)	Internal ($n = 5$)
Anaesthesia	112 \pm 12 (60–182)	113 \pm 28 (70–150)
Surgery	113 \pm 10 (83–179)	117 \pm 12 (104–136)
Recovery	149 \pm 28 (85–356)	196 \pm 54 (140–301)

162

163 All shad were detected moving upstream in fresh water, i.e. against the flow. Of all
 164 the batches, the first batch of fish (external tag) had the greatest mean upstream movement
 165 (61.1 ± 51.7 km) and mean total movement (122.9 ± 95.2 km), whereas the fourth batch
 166 (internal tag) spent the longest mean time in the river (21.4 ± 8.8 days) and fastest mean
 167 speed of upstream movement (1.10 ± 0.32 m/s) (Table 3). Fish in batches 1 and 4 were
 168 captured and released at the same location with external and internal tags, respectively, and
 169 had similar upstream movements (*t*-test on logged data; $t = 0.095988$, d.f. = 3.7202, $P =$
 170 0.926), total movements (*t*-test on logged data; $t = 0.31356$, d.f. = 4.3419, $P = 0.768$), times
 171 in the river (*t*-test; $t = -0.61932$, d.f. = 5.5427, $P = 0.560$) and speed of upstream movements
 172 (*t*-test; $t = 2.1894$, d.f. = 6, $P = 0.0711$) (Table 3). The individual fish with the greatest
 173 upstream (138.0 km) and total movements (281.4 km), and longest time in the river (29.8
 174 days) had an internal tag, whereas the fastest upstream movements (1.79 m/s) was by a fish
 175 that had an external tag.

176 Table 3. Mean \pm 95% C.I. (min.–max.) upstream movement (km), total movement (km),
 177 residence time in the receiver array (days) and speed of upstream movement (m/s) for shad
 178 tagged in five batches on the River Severn.

Batch	Upstream movement (km)	Total movement (km)	Time in river (days)	Speed of upstream movements (m/s)
1	61.1 ± 51.7 (27.7–113.1)	122.9 ± 95.2 (60.4–218.5)	18.3 ± 4.4 (13.9–21.2)	0.60 ± 0.19 (0.50–0.80)
2	16.4 ± 11.4 (4.0–37.7)	50.8 ± 26.5 (19.0–96.5)	12.8 ± 5.0 (6.6–23.3)	0.54 ± 0.14 (0.30–0.73)
3	14.4 ± 11.7 (1.0–33.9)	46.2 ± 28.7 (5.7–91.4)	8.4 ± 4.5 (0.2–16.2)	0.51 ± 0.17 (0.31–0.77)
4	58.0 ± 39.6 (30.7–138.0)	112.1 ± 83.6 (51.0–281.4)	21.4 ± 8.8 (9.3–29.8)	1.10 ± 0.32 (0.72–1.52)
5	15.5 ± 11.6	49.0 ± 16.4	8.9 ± 5.7	1.09 ± 0.38

(2.0–38.6)

(28.3–73.7)

(1.5–19.1)

(0.55–1.79)

179

180 Seventeen (68%) of the tagged shad performed a downstream migration to the
181 estuary between 25 May and 21 June 2017, 14.7 ± 3.9 days after tagging. Eight fish were
182 assumed to have died in the river (though tag failure or loss could not be ruled out) but were
183 tracked for a similar amount of time, i.e. 10.6 ± 8.2 days. The one exception (external tag)
184 was last detected 5 h after release, 5.7 km upstream of its release location. Four fish
185 (external = 2 and internal = 2) were last detected in the vicinity of a suspected spawning
186 location 9–27 days after release, three of which moved downstream after release and
187 subsequently returned to fresh water. Three fish (external = 2 and internal = 1) were last
188 detected moving downstream 5, 7 and 12 days after release, each having moved a minimum
189 of 18.7, 4.0 and 36.3 km, respectively, in an upstream direction while in fresh water.

190 **4. Discussion**

191 During this investigation, twaite shad, a threatened anadromous fish species that is sensitive
192 to handling and sedation, were successfully anaesthetised which enabled tags to be
193 surgically implanted into the peritoneal cavity. These findings are contrary to Rooney and
194 King (2014) who reported mortality of shad anaesthetised with 2-phenoxyethanol and
195 represents a substantial refinement of an accepted tagging protocol (*cf.* Breine et al., 2017).
196 The novel and successful use of MS-222 for shad might be reflective of high variability in
197 species-specific responses to different anaesthetics (e.g. Readman et al., 2017). These
198 refinements have important welfare, ethical and methodological implications for future shad
199 tracking studies.

200 Twaite shad are anadromous and iteroparous. In this study, a large proportion of the
201 tagged fish (68%) migrated downstream to the estuary after undertaking substantial
202 movements upstream and spent an appreciable amount of time in fresh water. This
203 suggested that tagging had little or no impact on their behaviour and that these fish evaded
204 predators (e.g. pike *Esox lucius* L., zander *Sander lucioperca* (L.), otter *Lutra lutra* (L.) and
205 cormorant *Phalacrocorax carbo* L.) and survived spawning. The assumed mortality of

206 individuals that did not return to the estuary (though tag failure or loss could not be ruled out)
207 was considered a result of either natural predation or post-spawning mortality, rather than a
208 direct consequence of being tagged. This is because they performed substantial upstream
209 movements, entered the estuary and returned to fresh water, were last detected at a
210 suspected spawning location and/or residence time was similar to fish that returned to the
211 estuary.

212 A commonly cited advantage of external tagging over surgical implantation is that
213 attachment can be faster (Jepsen et al., 2015; Breine et al., 2017), but internal implantation
214 was significantly faster than external attachment in this investigation. Although there was no
215 evidence of detrimental impacts of externally mounting tags they may have reduced
216 swimming performance through drag or disequilibrium. There are many other long-term
217 benefits of internal implantation to individual fish post-release, including improved tag
218 retention, reduced tissue damage, zero risk of biofouling and zero tag visibility to predators
219 (Cooke et al, 2013; Jepsen et al., 2015). Surgically implanting long-lived tags will also
220 provide substantial advances in ecological knowledge of iteroparous shad by enabling
221 multiple annual spawning migrations of the same individual to be studied. Consequently, the
222 number of fish that need to be tagged could also be reduced, thereby complying with the
223 reduction principle of animal research (Metcalfe and Craig 2011). These refinements should
224 be transferable to other fishes considered sensitive to handling and sedation, and should
225 lead to further refinements in tagging procedures during biotelemetry research.

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232 **References**

- 233 Aprahamian, M.W., Baglinière, J.L., Sabatié, M.R., Alexandrino, P., Thiel, R., Aprahamian,
234 C.D., 2003. Biology, status, and conservation of the anadromous Atlantic twaite shad
235 *Alosa fallax fallax*. Am. Fish. Soc. Symp. 35, 103–124.
- 236 Breine, J., Pauwels, I.S., Verhelst, P., Vandamme, L., Baeyens, R., Reubens, J. and Coeck,
237 J. 2017. Successful external acoustic tagging of twaite shad *Alosa fallax* (Lacépède
238 1803). Fish. Res. 191, 36–40.
- 239 Bridger, C.J., Booth, R.K., 2003. The effects of biotelemetry transmitter presence and
240 attachment procedures on fish physiology and behavior. Rev. Fish Sci. 11, 13–34.
- 241 Broadhurst, B.T., Ebner, B.C., Clear, R.C., 2009. Radio-tagging flexible-bodied fish:
242 temporary confinement enhances radio-tag retention. Mar. Freshwater Res. 60, 356-
243 360.
- 244 Campbell, H.A., Watts, M.E., Dwyer, R.G., Franklin, C.E. 2012. V-Track: software for
245 analysing and visualising animal movement from acoustic telemetry detections. Mar.
246 Freshwater Res. 63, 815-820.
- 247 Clough, S.C., Lee-Elliott, I.E., Turnpenny, A.W.H., Holden, S.D.J., Hinks, C. 2004. The
248 swimming speeds of twaite shad (*Alosa fallax*). R&D Technical Report W2-049/TR3.
- 249 Cooke, S.J., Midwood, J.D., Thiem, J.D., Klimley, P., Lucas, M.C., Thorstad, E.B., Eiler, J.,
250 Holbrook, C. and Ebner, B.C., 2013. Tracking animals in freshwater with electronic
251 tags: past, present and future. Anim. Biotelem. 1, 5.
- 252 Cooke, S.J., Woodley, C.M., Eppard, M.B., Brown, R.S., Nielsen, J.L., 2011. Advancing the
253 surgical implantation of electronic tags in fish: a gap analysis and research agenda
254 based on a review of trends in intraceolomic tagging effects studies. Rev. Fish. Biol.
255 Fish. 21, 127–51.
- 256 Hussey, N.E., Kessel, S.T., Aarestrup, K., Cooke, S.J., Cowley, P.D., Fisk, A.T., Harcourt,
257 R.G., Holland, K.N., Iverson, S.J., Kocik, J.F., Flemming, J.E.M., Whoriskey, F.G.,
258 2015. Aquatic animal telemetry: a panoramic window into the underwater world.
259 Science 348, 6240.

260 Jepsen, N., Koed, A., Thorstad, E., Baras, E. 2002. Surgical implantation of transmitters in
261 fish: how much have we learnt? *Hydrobiologia* 483, 239–48.

262 Jepsen, N., Thorstad, E.B., Havn, T., Lucas, M.C., 2015. The use of external electronic tags
263 on fish: an evaluation of tag retention and tagging effects. *Anim. Biotel.* 3, 49.

264 Johnson, M.W., Diamond, S.L., Stunz, G.W., 2015. External attachment of acoustic tags to
265 deepwater reef fishes: an alternate approach when internal implantation affects
266 experimental design. *Trans. Am. Fish. Soc.* 144, 851–859.

267 Jolly, M.T., Aprahamian, M.W., Hawkins, S.J., Henderson, P.A., Hillman, R., O'Maoiléidigh,
268 N., Maitland, P.S., Piper, R., Genner, M.J., 2012. Population genetic structure of
269 protected allis shad (*Alosa alosa*) and twaite shad (*Alosa fallax*). *Mar. biol.* 159, 675-
270 687.

271 King, J.J., Roche, W.K. 2008. Aspects of anadromous Allis shad (*Alosa alosa* Linnaeus) and
272 Twaite shad (*Alosa fallax* Lacépède) biology in four Irish Special Areas of
273 Conservation (SACs): status, spawning indications and implications for conservation
274 designation. *Hydrobiologia* 602, 145–154.

275 Metcalfe, J.D. and Craig, J.F., 2011. Ethical justification for the use and treatment of fishes
276 in research: an update. *J. Fish Biol.* 78, 393-394.

277 Mulcahy, D.M., 2003. Surgical implantation of transmitters into fish. *ILAR journal* 44, 295-
278 306.

279 R Core Team (2017). R: A language and environment for statistical computing. R
280 Foundation for Statistical Computing, Vienna, Austria.

281 Readman, G.D., Owen, S.F., Knowles, T.G., Murrell, J.C., 2017, Species specific
282 anaesthetics for fish anaesthesia and euthanasia. *Sci. Rep.-UK* 7, 7102.

283 Rooney, S., King, J., 2014. Presentation: Use of acoustic telemetry to monitor behaviour
284 during the upriver spawning migration of a diadromous fish, the twaite shad (*Alosa*
285 *fallax*). IFM Tagging and Telemetry Workshop Leeds, England, 22–23 July 2014.

286 Ross, L.G., Ross, B., 2009. *Anaesthetic and sedative techniques for aquatic animals*. John
287 Wiley & Sons.