# A peer-reviewed version of this preprint was published in PeerJ on 19 September 2018.

<u>View the peer-reviewed version</u> (peerj.com/articles/5459), which is the preferred citable publication unless you specifically need to cite this preprint.

Mattern T, McPherson MD, Ellenberg U, van Heezik Y, Seddon PJ. 2018. High definition video loggers provide new insights into behaviour, physiology, and the oceanic habitat of a marine predator, the yelloweyed penguin. PeerJ 6:e5459 <a href="https://doi.org/10.7717/peerj.5459">https://doi.org/10.7717/peerj.5459</a>



1	High definition video loggers provide new insights into behaviour,
2	physiology, and the oceanic habitat of marine top predators
3	Thomas Mattern <sup>1,2,*</sup> , Mike McPherson <sup>3</sup> , Ursula Ellenberg <sup>2</sup> , Yolanda van Heezik <sup>1</sup> , Philip J. Seddon <sup>1</sup>
4	<sup>1</sup> Department of Zoology, PO Box 56, University of Otago, Dunedin, NZ
5	<sup>2</sup> Department of Ecology, Environment and Evolution, La Trobe University, Melbourne, Australia
6	<sup>3</sup> 7 Adams Bottom, Leighton Buzzard, Bedfordshire, England
7	* Author for correspondence (t.mattern@eudyptes.net)
8	KEYWORDS
9	animal-borne video loggers, marine top-predators, diving behaviour, benthic habitat, predator-prey
10	interactions, penguins
11	Summary Statement
12	The use of full HD animal-borne video loggers can provide information about a wide range of
13	behavioural, physiological and environmental aspects of marine top-predators' biology.
14	
15	
16	
17	



## 18 Abstract

- Camera loggers are increasingly used to examine behavioural aspects of free-ranging
  animals. However, often video loggers are deployed with a focus on specific behavioural
  traits utilizing small cameras with a limited field of view, poor light performance and video
  quality. Yet rapid developments in consumer electronics provide new devices with much
  improved visual data allowing a wider scope for studies employing this novel methodology.
- 2. We developed a camera logger that records full HD video through a wide-angle lens, providing high resolution footage with a greater field of view than other camera loggers. Main goal was the analysis of foraging behaviour of a marine top-predator, the Yellow-eyed penguin in New Zealand, in the context habitat characteristics. Frame-by-frame analysis allowed accurate timing of prey pursuits and time spent over certain seafloor types. Similarly, it was possible to time breathing intervals between dives and quantify exhalation events during prey events, a previously undescribed behaviour. Screen overlays facilitate analysis of flipper angles and beat frequencies throughout various stages of the dive cycle.
- 3. The recorded video footage showed that prey species were associated with certain seafloor types, revealed different predator evasion strategies by benthic fishes, and highlighted varying energetic consequences for penguins pursuing certain types of prey. Flipper movement analysis confirmed decreasing effort during descent phases as the bird gained depth, and that ascent was principally passive. Breathing episodes between dives were short (<1 s) while the majority of the time was devoted to subsurface scanning with a submerged head.</p>
- 4. Video data recorded on free-ranging animals not only provides a wealth of information recorded from a single deployment but also necessitates new approaches with regards to analysis of visual data. Here, we demonstrate the diversity of information that can be gleaned from video logger data, if devices with high video resolution and wide field of view are utilized.



## 44 Introduction

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

61

62

63

64

65

66

67

68

69

Examining the at-sea behaviour of marine animals has long been a challenging endeavour. Direct visual observations of behaviour are almost impossible, especially when most of it happens under the ocean's surface. In recent decades, advances in telemetry technologies and the emergence of bio-logging hardware have provided the means to track marine animals and reveal their foraging behaviour in great detail. Starting in the 1970s with rather crude location estimates and limited data quality recorded by unwieldy devices that could only be used on large animals, advancements in micro-electronics have resulted in ever smaller and more accurate loggers to pinpoint an animal's position to within a few metres and record their diving depths with oceanography-grade precision (Wilmers et al. 2015). New technologies such as accelerometers and gyroscopes further refined methods to study marine habitat use (e.g. Noda et al. 2014). Yet, placing dive metrics into a complex behavioural and environmental context can be difficult; ideally a reference framework based on direct observations is used to match up dive metrics and actual behaviours (e.g. Moreau et al. 2009; Volpov et al. 2016). So the original dilemma of having to make direct observations of marine animal behaviours still persists. Animal-borne video recorders offer the means to overcome this problem. In recent years animal-borne camera systems have made it possible to log in situ observations of behaviour from the animal's point of view (Moll et al. 2007). For example, deployment of lightweight video cameras on flying birds provided new perspectives on prey pursuit in falcons (Kane & Zamani 2014) and revealed how albatrosses use the presence of killer whales to locate prey (Sakamoto et al. 2009). No other animal group has been more subject to deployment of video recording devices in recent years than marine animals. By overcoming the observational barrier at sea, video loggers are providing copious amounts of novel data that range from identification of feeding strategies (Takahashi et al. 2008) and previously unknown food sources (Thiebot et al., in review), to social interactions such as group foraging (Sutton et al. 2015) or kleptoparasitism (Handley & Pistorius 2015). Video data also offers the means to calibrate other bio-logging data (Watanabe & Takahashi 2013; Gómez-Laich et al. 2015).



72

73

74

75

76

77

78

79

80

81

82

83

84

85

86

87

88

89

90

91

92

93

94

95

What most of these studies have in common is their focus on specific behavioural traits while providing limited information about the environment the behaviours occurred in. This is principally due to limitations of the video hardware used, which has to be small and light-weight so as to not overly impede the study animal's movement capabilities (Ludynia et al. 2012) and hence behaviour. As a result, video quality (i.e. image resolution, field of view/FOV) is sacrificed in favour of smaller cameras. However, with the rise in popularity of action cams on the consumer market, new video devices have recently become available with high definition video capabilities and wide-angle optics, suitable for deployment even on smaller marine animals such as penguins. This leap in quality has significant implications for the study of marine animals as it not only allows monitoring of wideranging aspects of behaviour, but also provides new opportunities for the visual analysis of the environment the animals use. This is particularly relevant in species that forage at the seafloor where video data can provide extensive information about the benthic habitat (Watanuki et al. 2008). The yellow-eyed penguin (Megadyptes antipodes) in New Zealand is known to be a benthic forager (Mattern et al. 2007) that feeds primarily on demersal fish species (van Heezik & Davis 1990; Moore et al. 1995). It has been suggested that this strategy might come at the expense of reduced behavioural flexibility, with subsequent vulnerability to changes in the marine environment (Mattern et al. 2007). In particular, degradation of seafloor ecosystems in the wake of commercial bottom fisheries are suspected to influence yellow-eyed penguin foraging success and population developments (Browne et al. 2011; Mattern et al. 2013). While the species' at-sea movement and diving behaviour has been subject to a number of studies in the past decades (Moore et al. 1995; Mattern et al. 2007, 2013), information about their benthic habitat is scarce. To be able to assess the extent to which penguin behaviour and foraging success correlates with the composition of the benthic habitat, we developed a camera logger that records full HD videos through wide-angle lenses. The main focus of our study was to assess the suitability of the device for the visual analysis of penguin prey pursuit behaviour and characteristics of the benthic ecosystem. However, the deployment revealed far more information than was anticipated. The video data PeerJ Preprints | https://doi.org/10.7287/peerj.preprints.2765v2 | CC BY 4.0 Open Access | rec: 6 Mar 2017, publ: 6 Mar 2017 Peer Preprints

100

101

103

104

105

106

107

108

109

110

111

113

114

115

116

117

118

119

120

- provided novel insights into physiological aspects of the penguin's diving activities and allowed us to
- 97 draw conclusions about prey capture techniques. In this paper, we summarise our findings,
- demonstrate analytical approaches to evaluate animal-borne video data, and highlight the multi-
- 99 disciplinary potential of full HD video loggers.

# Materials and methods

### Study site and species

The Yellow-eyed penguin, classified as "Endangered" by the IUCN Redlist (BirdLife International

2012), is one of five penguin species endemic to the New Zealand region and occurs on the sub-

Antarctic Auckland and Campbell Islands as well as the south-eastern coastlines of New Zealand's

South Island and Stewart Island (Seddon et al. 2013). This study was carried out at the Boulder Beach

complex, Otago Peninsula, South Island, New Zealand (45.90°S, 170.56°E). Penguins from this site

have been subject to foraging studies in the past decade that have suggested substantial impact of

bottom trawling activities on the yellow-eyed penguins' at-sea movements (Mattern et al. 2013).

The research was approved by the University of Otago's Animal Ethics Committee (#11/14) and was

permitted by the NZ Department of Conservation (45799-FAU).

#### Video logger & deployment

112 We developed a high-definition video logger (dimensions LxWxH, 80x40x23mm; weight: 48g) which

is combined with a time-depth recorder (TDR, 12x31x11mm, 6.5g; AXY-depth, Technosmart Ltd. Italy)

and a GPS logger (modified i-gotU, GT-120, Mobile Action Technology Inc., Taiwan, 22x31x11mm,

12g). The latter two devices were combined into a single unit by gluing the AXY-depth to the longer

side of the GPS device. Camera and logger combination were then attached individually in line to the

lower back of the penguin using adhesive tape (Wilson et al. 1997). Additional drag of the devices

was principally limited to the cameras frontal area (Bannasch et al. 1994). The camera logger consists

of a modified Mobius action-cam with a 130° wide-angle lens (www.mobius-actioncam.com). In

order to achieve the smallest and lightest device possible, the camera electronics, video sensor and

PeerJ Preprints | https://doi.org/10.7287/peerj.preprints.2765\( \mathbb{Z} \) CC BY 4.0 Open Access | rec: 6 Mar 2017, publ: 6 Mar 2017

122

123

124

125

126

127

128

129

130

131

132

133

134

135

136

137

138

139

140

141

142

143

144

145

lens were removed from the casing and the battery replaced with a 1200 mAh Lithium Polymer battery to extend recording time. A small bespoke timer board was developed to allow the camera to be fired at a pre-determined time. The recording then ran until the battery fell below the minimum operating voltage of the camera (ca. 2-4 hours). Connections were provided to allow programming the alarm and also to access the camera's USB port for managing camera setting, extracting the video data and recharging the battery. The board was isolated electrically to prevent the contacts from shorting as sea-water is conductive. Activation of the interface was achieved using a Hall-effect device. An Arduino-based interface was developed to allow the current date/time and alarm time to be set. The camera was programmed to record video data at a resolution of 1920x1080 pixels (1080p) at a frame rate of 30 frames per second. Video data was recorded in H.264 MPEG4 format and stored on a 32GB MicroSD card. The camera was encased in epoxy resin to prevent water damage and provide the device with a hydrodynamic shape to reduce additional drag (Culik et al. 1994). After device recovery, data was downloaded through the camera's USB interface. Since the logger stores video data as a series of full frame images ('progressive scan'), it was possible to conduct a frame-by-frame analysis to accurately time components of the bird's behaviour – i.e. breathing intervals, flipper beat frequencies and amplitudes – as well as time spent over certain benthic habitats. Video analysis was conducted in professional editing software (Adobe Premiere Pro CS 6, Adobe Systems Inc., San Jose, CA, USA) which allows video scrubbing and provides the option to display frame number in the preview timer. The video logger was deployed on a breeding male Yellow-eyed penguin tending two chicks on 17 December 2015. The bird left on a single foraging trip on 18 December before the device was recovered on 19 December.

### Analysis of behaviours

**Prey pursuits.** We defined the beginning of a prey pursuit as the moment when the penguin markedly accelerated while swimming along the seafloor; the end was reached when the penguin



decelerated again to its previous cruise speed (if no prey was caught), or when the prey item was 146 147 swallowed completely. Acceleration and deceleration were associated with temporary blurring of the 148 video footage due to irregular body movement, allowing for exact timing of prey pursuits. Where 149 possible, prey species were identified from frames providing a clear view of the prey item. 150 Beyond prey interactions, the video data offered the opportunities to analyse physiological aspects 151 of the penguin's behaviour. 152 **Surface breathing.** We timed breathing events when the penguin was at the surface following a dive. 153 Noting frame numbers when the bird raised its head out of the water before lowering below the 154 surface again made it possible to determine time the penguin was able to respire (https://vimeo.com/179414575#t=145). 155 156 Flipper movements. During dives, flipper beat frequencies (beats per minute, BPM) were determined 157 by counting the number of frames required to complete one flipper beat cycle, beginning the count when the flipper angle reached its maximum upward inclination and ending with the frame prior to 158 159 the subsequent maximum upward inclination. In the video editing software we overlaid a template 160 indicating 10, 30, 50, 70 and 90 degree angles radiating from the base of the flippers on the video 161 data (https://vimeo.com/179414575). This allowed us to visually determine maximum amplitude of 162 each flipper beat to the nearest 5°. Analysis of benthic habitat 163 164 For all dives, the benthic habitat was classified according to sediment type (fine sand, coarse sand 165 with shell fish fragments, gravel), sediment structure (flat, sediment ripples) and composition of the 166 epibenthic communities. For the latter, we used a presence/absence approach in which the 167 occurrence of brittlestars (Ophiuroidea), anthozoans (anemones and soft corals), and horse mussels 168 (Atrina zelandica) within a 30-frames time window. Future deployments with a functional GPS logger 169 can be used for more elaborate analysis of the benthic habitat, e.g. the creation of biodiversity 170 indices.

Peer Preprints

173

174

175

176

177

178

179

180

181

182

183

184

185

186

187

188

189

190

191

192

193

194

A selection of edited video clips demonstrating the various behaviours and habitat types descried 172

above can be accessed via https://vimeo.com/album/4103142.

## Dive data analysis

Dive data recorded by the TDR was analysed following methods described in detail in Mattern et al. (2007). Key dive parameters determined were maximum depth reached, duration of the dive event and its three main phases (i.e. descent, bottom phase, ascent) as well as vertical velocities during descent and ascent. Dives were classified as pelagic or benthic dives using dive profile characteristics, where near horizontal bottom phases with little vertical variance as well as consistent maximum dive depths on consecutive dives were used as cues for diving along the seafloor. This approach was validated by recorded video data.

Statistical analysis was carried out in R (R Development Core Team 2008).

# Results

Foraging trip length, diving events and video coverage.

The day following camera deployment, the penguin performed a 10.7 hour-long foraging trip. The first dive event was recorded at 5:30 hrs and the last event concluded at 16:10 hrs. The bird performed 286 dives of which 159 dive profiles matched the criteria for benthic dives (Figure 1). Median dive depth reached during benthic dives was 54.4m (range: 4.8-62.1 m, n=159) whereas the majority of pelagic dives occurred in the upper 10m of the water column (median: 7.8m, range: 0.5-31.7 m, n=127): camera footage confirmed these to be principally travelling behaviour (https://vimeo.com/179414642). For the first 3½ hours of the foraging trip (05:30-09:00 hrs) the bird performed mainly pelagic dives, indicating primarily travelling behaviour towards its main foraging grounds; the remaining hours (09:00-16:10 hrs) the bird principally devoted its time to benthic diving (Figure 1). The camera operated continuously from 11:00:22 hrs to 13:01:43 hrs. Due to occasional frame loss when data were written to memory, total length of the recorded footage amounted to 2

196

197

198

199

200

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

217

218

219

220

hours 8 seconds). 46 complete dives were video recorded which corresponds to 16% of all dive events; of these 32 dives were benthic dives. However, dives were longer during the middle of the day so that camera footage covered 25% of the trip's cumulative dive time. Prey pursuits & capture A total of 20 prey pursuits was recorded at the seafloor; 14 of these resulted in successful capture of either opalfish (Hemerocoetes monopterygius, 10 specimens) or blue cod (Parapercis colias, 2 specimens); prey species could not be identified during two captures, but the penguin's searching behaviour and ease of ingestion suggested these were opalfish (Figure 2). All of these prey pursuits occurred at the sea floor with the penguin swimming very close to the bottom (https://vimeo.com/179414724). During the camera operation time, the penguins spent 5.7 minutes on prey pursuit, which corresponds to 19% of the total time the bird foraged along the seafloor (29.9 minutes) and 6% of its total dive time (89.9 minutes); 3.8 minutes were devoted to pursuing and capturing opalfish; 46 seconds were used for the two blue cod captures, and the remaining 1.2 minutes were unsuccessful prey pursuits (Figure 2). Two main prey pursuit strategies became apparent that were associated with prey species. When catching opalfish, the penguin would glide closely above the seafloor, sometimes briefly accelerating before starting to hover over a certain spot while repeatedly pecking at the substrate until the prey item was captured (https://vimeo.com/179414724). During encounters with blue cod prolonged pursuits ensued during which fish zigzagged at a fast pace along the seafloor (https://vimeo.com/179414724#t=2m46s). In one instance the fish was caught as it appeared to seek shelter at the base of a horse mussel protruding from the substrate (https://vimeo.com/179414724#t=2m55s). An unsuccessful prey pursuit of blue cod ended with the fish escaping under what appeared to be a half-buried back plate of a dishwasher (https://vimeo.com/179414777). A third blue cod encounter occurred just seconds after a successful capture of an opalfish; it seems likely that the resulting prolonged bottom time and oxygendemanding prey pursuits drove the penguin to carry the fish to the surface at an almost vertical angle

PeerJ Preprints | https://doi.org/10.7287/peerj.preprints.2765\( \mathbb{Q} \) | CC BY 4.0 Open Access | rec: 6 Mar 2017, publ: 6 Mar 2017

Peer Preprints

245

as indicated sun disc's central position in the frame; the fish was ultimately dropped at the surface 221 222 (https://vimeo.com/179414724#t=3m07s). Benthic habitat 223 224 During the video logger's operating time, the penguin spent 29.9 minutes foraging along the seafloor. 225 The majority of the penguin's bottom time (90%) was spent over coarse sand, whereas time spent 226 over fine sand (7%) and gravel (0.9%) was negligible (Figure 2). Two thirds of the bottom time (65.9%) 227 was spent over sand ripples, the remaining time (34.1%) the bird foraged over flat ground. Brittle 228 stars and anthozoans were present in most areas visited by the penguin with the former being 229 present in 22.5 mins (75%) of the benthic video footage while the latter occur for a total of 17.9 mins 230 (60%). Horse mussels were present for a total of 9.3 minutes (31%) of the bottom time. 231 Prey encounters were associated with certain benthic habitat types. All prey encounters occurred 232 over coarse sand although the sediment structure differed depending on prey species. Opalfish were 233 principally encountered on sediment ripples (93.6% of the total prey pursuit time, 234 https://vimeo.com/179414724), while flat bottom habitat played a more important role during blue 235 cod pursuits (52.8% of pursuit time, <a href="https://vimeo.com/179414724#t=2m32s">https://vimeo.com/179414724#t=2m32s</a>). With regards to 236 epibenthic characteristics, brittle stars and anemones were present during the majority of the prey 237 pursuit times for both fish species (Figure 3). However, horse mussels were present only during blue 238 cod pursuits (81.4% of pursuit time). 239 Flipper movement 240 When descending to the sea floor the penguin propelled itself with fast, strong flipper strokes that 241 got progressively slower and less pronounced with time and, thus, increasing depth (flipper 242 amplitude:  $\rho$ =-0.83,  $F_{1.363}$ =791.8, p<0.001, BPM:  $\rho$ =-0.36,  $F_{1.363}$ =55.2, p<0.001, Figure 4a&b; https://vimeo.com/179414575). In contrast, ascending was principally passive with the penguin using 243 244 its natural buoyancy to return to the surface, occasionally aided by a few strokes in the early stages

of the ascent (flipper amplitude:  $\rho$ =-0.08,  $F_{1,74}$ =0.5, p=0.488; BPM:  $\rho$ =-0.52,  $F_{1,74}$ =0.5, p<0.001, Figure



4c&d) and no observable flipper movements towards the end of the dive 246 247 (https://vimeo.com/179414575#t=1m49s). Despite differences in flipper movement between the two transit phases of a dive, the vertical velocities recorded by the TDR did not differ significantly 248 249 (mean descent velocity: 1.45±0.28 m/s, mean ascent velocity: 1.36±0.57 m/s, n=159 dives, Welch's t-250 test:  $t_{232}$ =1.73, p=0.09). 251 During the bottom phase flipper amplitudes and beat frequencies showed no correlation with 252 relative bottom time (flipper amplitude:  $\rho$ =-0.08,  $F_{1,74}$ =0.5, p=0.488; flipper BPM:  $\rho$ =-0.52,  $F_{1,74}$ =0.5, 253 p<0.001, Figure 4c&d). This is owing to the fact that bottom phases consisted of a mix of searching 254 behaviour and high speed prey pursuit (https://vimeo.com/179414575#t=0m33s). While searching 255 the penguin showed lower flipper beat frequencies (133±48 BPM, n=809) paired with greater flipper 256 amplitudes (53°±14°) when compared to prey pursuit (BPM: 162±44, n=113,  $t_{232}$ =-13.37, p<0.001; 257 amplitude: 45°±7°, *t*<sub>152</sub>=6.39, p<0.001). Surface breathing & underwater exhalation 258 Frame counts of the video footage during 10 random selected surface periods between dives showed 259 260 that the penguin lifted its head out of the water to breathe for only brief moments (average 261 duration: 0.77±0.22 s, n=193); for the majority of the time at the surface the bird kept its head under 262 water (1.53±1.19 s, n=182) (https://vimeo.com/179414575#t=2m25s). Duration of breathing 263 intervals increased with ongoing duration of the surface period (Pearson correlation:  $\rho$ =0.45, 264  $F_{1,191}$ =47.4, p=<0.001, Fig 3) indicating increased respiration activity in preparation for the next dive (Figure 3). 265 During the dive, exhalation regularly occurred at the onset of phases with increased acceleration (i.e. 266 267 prey pursuit). Such exhalations were brief but performed with substantial force; air was jetted from 268 the nostrils as a fine gas spurt (https://vimeo.com/179418254). During the passive phase of the 269 ascent, the penguin frequently exhaled as indicated be a stream of large bubbles released from the 270 nostrils. The bird released substantial amounts of air on the last few meters immediately prior to

Peer Preprints

reaching the surface (<a href="https://vimeo.com/179414575#t=2m18s">https://vimeo.com/179414575#t=2m18s</a>). While some of this air may have
been released from the plumage (c.f. Davenport *et al.* 2011) bubbles seem principally to originate
from the frontal head region; there was no visible major gas release from the penguin's back region.

## Discussion

The high-quality video footage provided a substantial amount of new insights into the foraging behaviour of Yellow-eyed penguins and their benthic habitat, while the device did not appear to substantially affect the penguin's underwater mobility.

#### Device effects

Attaching external recording devices to diving animals always comes at the cost of compromising their streamlined body shape (e.g. Ludynia et al., 2012), a problem that can be mitigated via device shape, size and attachment position (Bannasch *et al.* 1994). At the surface there were no indications that the penguin was negatively affected by the device; the bird did not exhibit balancing problems which externally attached devices can cause in smaller species (Chiaradia *et al.* 2005), nor did it peck at the device frequently which suggests aberrant behaviour (Wilson & Wilson 1989). Moreover, the number of successful prey captures further suggests that the bird's foraging capabilities were not drastically affected by the video logger. The bird was one of the few breeders that raised two chicks to fledging in an otherwise poor breeding season.

## Predator-prey interactions & prey species importance

In line with previous descriptions of yellow-eyed penguins as primarily benthic foragers (Mattern *et al.* 2007), the penguin's prey pursuit and captures recorded during the camera operation indeed all occurred at the sea floor. Swimming very close to the seafloor could serve several purposes. It could be a strategy to flush out benthic prey that blends in with the substrate, but it could also mean the penguin has a greater chance to see its prey from the side, and thus reduce the effect of prey camouflage. Opalfish, for example, are very well camouflaged and very difficult to make out from above (Roberts *et al.* 2015). This species seems to principally rely on its camouflage as means of PeerJ Preprints | https://doi.org/10.7287/peerj.preprints.2765**12** | CC BY 4.0 Open Access | rec: 6 Mar 2017, publ: 6 Mar 2017

297

298

299

300

301

302

303

304

305

306

307

308

309

310

311

312

313

314

315

316

317

318

319

320

predator avoidance since none of the opalfish captures involved a chase. In contrast, during both successful blue cod encounters extended high-speed chases ensued before the fish was ultimately captured. Blue cod and opalfish differ significantly in their anatomy with the small, slender opalfish presumably lacking the physical prowess for prolonged swimming when compared to muscular blue cod (Roberts et al. 2015). When facing an air breathing predator, the latter strategy is likely advantageous as the predator's increased energy requirements for pursuit make escape a more likely outcome for the prey. The penguin's hasty ascent and subsequent failure to consume a blue cod it captured after a 22-second-long chase demonstrates the efficacy of this evasion strategy. Both opalfish and blue cod have previously been found to be among the most important prey items in the Yellow-eyed penguin's diet (van Heezik 1990b; Moore & Wakelin 1997). While both fish species have comparable energetic values (~20 kJ g<sup>-1</sup>, Browne et al., 2011), the body mass of opalfish is considerably lower when compared to blue cod (van Heezik 1990a,b). So it is possible that the energy gain from catching blue cod justifies the expenditure to catch it, while the easier-to-catch opalfish might need to be caught in larger quantities. However, recent studies suggest that blue cod might be suboptimal prey for chick-rearing yellow-eyed penguins due to their size (Browne et al. 2011; Mattern et al. 2013) so that the penguins ability to locate prey such as opalfish might be a decisive factor with regards to reproductive success.

#### Benthic environment

Judging from the total time the bird spent over a benthic environment dominated by coarse sand and sediment ripples (65.9% of total bottom time) as well as almost exclusive encounters of opalfish over such habitat (Figure 2), it can be assumed that the penguin focussed principally on this species. Blue cod encounters were associated with the presence of horse mussels. These large bivalves protrude from the seafloor and provide hard substrate for other epibenthic taxa, thereby increasing local benthic biodiversity (Cummings *et al.* 1998). Benthic habitat with increased benthic biodiversity is generally more attractive to a variety of benthic fish species, most likely due to enhanced feeding



322 shelter to avoid capture (https://vimeo.com/179414777). 323 The majority of prey pursuits occurred in areas that featured anthozoans, principally sea anemones 324 (Figure 2). Anemones are known to play an important role as refugia and feeding habitats for small 325 fish (Elliott 1992) and could therefore be another indicator for locally increased biodiversity. Brittle 326 stars on the other hand, although equally abundant, seemed to be of lesser relevance with regards to 327 prey encounters. So it appears that examining the composition of the benthic habitat alone might 328 enable assessment of which prey types penguins are foraging for, though more data is required 329 before conclusions can be drawn. However, this already hints at the potential for wide-ranging 330 habitat analysis of at-sea movements in benthic top predators, provided that spatial distribution of 331 the different benthic habitats can be obtained. While in our specific case, no such habitat maps exist, 332 planned further deployments of video loggers are expected to provide the necessary environmental 333 information. 334 Deploying video loggers on penguins could enable detailed mapping of the benthic habitat within the 335 species home ranges. Yellow-eyed penguins are known to have preferred individual foraging areas 336 often with little overlap between birds (Moore 1999). Moreover, the birds tend to often dive along 337 the seafloor when swimming towards their foraging grounds (Mattern et al. 2007) so that camera 338 logger data in combination with GPS information can be used to establish spatial biodiversity indices 339 and benthic habitat maps. 340 The outer ranges of the marine habitat of Yellow-eyed penguins from the Otago Peninsula is subject 341 to bottom fisheries which have a profound effect on benthic ecosystems (e.g. Hinz et al., 2009; 342 Queirós et al., 2006; Schratzberger and Jennings, 2002). Yellow-eyed penguins have been found to 343 forage in the wake of trawl fisheries, potentially to the detriment of their reproductive success 344 (Mattern et al. 2013). Changes in sediment structure and epibenthic biodiversity as a result of 345 bottom trawl disturbance likely negatively affect the penguins' foraging success (Browne et al. 2011).

conditions (Cranfield et al. 2001). Our video data also suggests that the fish use the bivalves as

347

348

349

350

351

352

353

354

355

356

357

358

359

360

361

362

363

364

365

366

367

368

369

370

Camera loggers can help to determine how much of the penguins' foraging habitat has been compromised by fishing activities and what the consequences are for this species' foraging behaviour and success.

Beyond investigations of behaviour in a wider environmental context, our study also shows the potential application of camera loggers for the investigation of physiological aspects of marine animals.

#### Flipper movements

Our observations of flipper movements, i.e. strong flipper movements at the beginning of a dive that decrease with depth, and cessation of flipper movements during ascent, align with findings reported in other penguins. Using accelerometers, Sato et al. (2002) found that King penguins showed vigorous flipper beating at the beginning of a dive to counter positive buoyancy. With increasing depth, air volume in the penguin's body becomes compressed, reducing its buoyancy so that fewer flipper beats are required. That this also applies to flipper amplitude (Fig 4) was not detectable by using body acceleration as the only measure. A more elaborate system of sensors and magnets attached to flippers was used on Magellanic penguins which allowed the recording of both flipper amplitudes and beat frequencies (Wilson & Liebsch 2003). However, the system proved to be prone to failure, rendering the use of back-mounted wide-angle cameras a much more reliable alternative. Flipper beat frequencies and amplitudes are directly related to energy expenditure (Kooyman & Ponganis 1998; Sato et al. 2011). They provide the means for the quantification of energy budgets (Wilson & Liebsch 2003) and subsequently can be used to assess individual fitness in relation to foraging success and subsequent reproductive performance (Kooyman & Ponganis 1998). We provide proof that the ascent phase in penguins is largely passive, as has been suggested using both accelerometers and magnets (Sato et al. 2002; Wilson & Liebsch 2003). Sato et al. (2002) concluded that during ascent penguins benefit from expanding air volume in their body which increases their buoyancy as they get closer to the surface. Penguins also actively slow down their

Peer Preprints

371

372

373

374

375

376

377

378

379

380

381

382

383

384

385

386

387

388

389

390

391

392

393

394

395

ascent and it was argued that this could be achieved by increasing the attack angles of their flippers to increase drag (Sato *et al.* 2002). Judging from body movements apparent in the video data during the ascent phases we suggest that the yellow-eyed penguin indeed adjusted flipper attack angles while ascending, although this seems to be more for steering. Based on the video footage it appears that the birds might have used controlled exhalation towards the end of the ascent to control speed (<a href="https://vimeo.com/179414575#t=2m18s">https://vimeo.com/179414575#t=2m18s</a>).

#### Respiration

The video data provides new insights into the respiration of Yellow-eyed penguins. To date it was unclear whether penguins exhale regularly while diving. Various studies estimated diving air volume via a penguin's buoyancy calculated from its ascent speeds at the end of dives (Sato et al. 2002, 2011). However, the accuracy of this approach is compromised if the penguins were to exhale prior to their final ascent (Ponganis et al. 2015). The video data clearly showed that the penguin generally exhaled when accelerating during prey pursuit so that models estimating diving air volume via the proxy buoyancy must take acceleration into account. The fact that the penguin exhaled when accelerating probably serves the purpose of reducing blood CO2 and mobilizing O2 from oxygen stores for prey pursuit. Such pursuits must be costly in terms of oxygen consumption as is evident from the observed consecutive prey encounters during one single dive, which resulted in the penguin letting go of the second fish after a rapid ascent to the surface (https://vimeo.com/179414724#t=3m07s). Unlike seals that have been found to exhale when ascending from deep dives, most likely to reduce drop in blood oxygen (Hooker et al. 2005), the penguin principally exhaled during the second half of the ascent possibly indicating adjustment of buoyancy and ascent speed (but see also Davenport et al. 2011). Reoxygination during the surface period in penguins is highly optimized (Wilson et al. 2003). Inhalation events at the surface are brief so that the bird can frequently lower its head into the water, presumably in an effort to look out for potential predators (e.g. sharks, sea lions; Seddon et al., 2013).



Conclusions

396

397

398

399

400

401

402

403

404

405

406

407

408

409

410

411

412

413

414

415

416

417

418

419

the manuscript.

The deployment of a full HD video logger on a Yellow-eyed penguin resulted in a versatile visual data set that provided a variety of information well beyond what was initially intended. Enhanced video quality allows detailed analysis of the benthic environment as well as prey encounter rates and prey composition. In combination with GPS data, the potential for a comprehensive survey of benthic ecosystems is substantial highlighting the multi-disciplinary potential of such data. A large field of view achieved through wide-angle lenses furthermore allows detailed analysis of flipper movements, which to date could only be achieved through elaborate modelling of accelerometer data (Sato et al. 2002, 2011) or use of complicated magnetic logger setups (Wilson & Liebsch 2003). Neither of these setups provided information about exhalation, which appears to play a much more important role during diving than previously thought. When comparing video data recorded here with videos from previously published studies (e.g. Watanabe and Takahashi, 2013) it becomes clear that greater visual fidelity of full HD cameras comes along with a much wider range of quantifiable data. This creates a new opportunity for a more holistic approach to study the diving behaviour of marine animals that integrates behaviour, physiology and their environment. Acknowledgments We would like to thank Horst Mattern, Melanie Young and Jim Watts for help in the field, and to Leon Berard for first preliminary evaluation of the video data. Competing Interest The authors declare no competing or financial interests. Author contribution T.M. designed the study; M.M. and T.M. developed the camera loggers; T.M., P.J.S. and J.v.H. conducted the field work; T.M. and U.E. analysed the data; T.M., M.M., U.E., Y.v.H. and P.J.S. wrote



420 Funding

421 This work was supported by an Otago University Research Grant [PL 112034.01 R.FZ] issued to P.J.S.



422 6. References

423	Bannasch, R., Wilson, R.P. & Culik, B.M. (1994). Hydrodynamic aspects of design and attachment of a
424	back-mounted device in penguins. <b>194</b> , 83–96.
425	BirdLife International. (2012). Megadyptes antipodes. The IUCN Red List of Threatened Species, p.
426	e.T22697800A40186242.
427	Browne, T., Lalas, C., Mattern, T. & Van Heezik, Y. (2011). Chick starvation in yellow-eyed penguins:
428	Evidence for poor diet quality and selective provisioning of chicks from conventional diet
429	analysis and stable isotopes. Austral Ecology, <b>36</b> , 99–108.
430	Chiaradia, A., Ropert-coudert, Y., Healy, M. & Knott, N. (2005). Finding the balance : the effect of the
431	position of external devices on little penguins. <i>Polar Bioscience</i> , <b>18</b> , 46–53.
432	Cranfield, H.J., Carbines, G., Michael, K.P., Dunn, A., Stotter, D.R. & Smith, D.J. (2001). Promising signs
433	of regeneration of blue cod and oyster habitat changed by dredging in Foveaux Strait, southern
434	New Zealand. New Zealand Journal of Marine and Freshwater Research, 35, 897–908.
435	Culik, B.M., Wilson, R.P. & Bannasch, R. (1994). External devices: how important is shape. 118, 353–
436	357.
437	Cummings, V.J., Thrush, S.F., Hewitt, J.E. & Turner, S.J. (1998). The influence of the pinnid bivalve
438	Atrina zelandica (Gray) on benthic macroinvertebrate communities in soft-sediment habitats.
439	Journal of Experimental Marine Biology and Ecology, 228, 227–240.
440	Davenport, J., Hughes, R.N., Shorten, M. & Larsen, P.S. (2011). Drag reduction by air release
441	promotes fast ascent in jumping emperor penguins-a novel hypothesis. Marine Ecology
442	Progress Series, <b>430</b> , 171–182.
443	Elliott, J. (1992). The role of sea anemones as refuges and feeding habitats for the temperate fish
444	Oxylebius pictus. <i>Environmental Biology of Fishes</i> , <b>35</b> , 381–400.

Gómez-Laich, A., Yoda, K., Zavalaga, C. & Quintana, F. (2015). Selfies of Imperial Cormorants 445 446 (Phalacrocorax atriceps): What Is Happening Underwater? PLOS ONE, 10, 1–18. 447 Handley, J.M. & Pistorius, P. (2015). Kleptoparasitism in foraging gentoo penguins Pygoscelis papua. 448 *Polar Biology*, **39**, 391–395. 449 van Heezik, Y. (1990a). Diets of yellow-eyed, Fiordland crested, and little blue penguins breeding 450 sympatrically on Codfish Island, New Zealand. New Zealand Journal of Zoology, 17, 543-548. 451 van Heezik, Y. (1990b). Seasonal, geographical, and age-related variations in the diet of the Yellow-452 eyed Penguin (Megadyptes antipodes). New Zealand Journal of Zoology, 17, 201–212. 453 van Heezik, Y. & Davis, L.S. (1990). Effects of food variability on growth rates, fledging sizes and 454 reproductive success in the Yellow-eyed Penguin Megadyptes antipodes. Ibis, 132, 354–365. 455 Hinz, H., Prieto, V. & Kaiser, M.J. (2009). Trawl disturbance on benthic communities: Chronic effects 456 and experimental predictions. Ecological Applications, 19, 761–773. 457 Hooker, S.K., Miller, P.J.O., Johnson, M.P., Cox, O.P. & Boyd, I.L. (2005). Ascent exhalations of 458 Antarctic fur seals: a behavioural adaptation for breath-hold diving? Proceedings of the Royal 459 Society of London B: Biological Sciences, **272**, 355–363. 460 Kane, S.A. & Zamani, M. (2014). Falcons pursue prey using visual motion cues: new perspectives from 461 animal-borne cameras. Journal of Experimental Biology, 217, 225–234. 462 Kooyman, G.L. & Ponganis, P.J. (1998). The physiological basis of diving to depth: birds and mammals. 463 Annual review of physiology, **60**, 19–32. Ludynia, K., Dehnhard, N., Poisbleau, M., Demongin, L., Masello, J.F. & Quillfeldt, P. (2012). 464 Evaluating the Impact of Handling and Logger Attachment on Foraging Parameters and 465 466 Physiology in Southern Rockhopper Penguins. *PLoS ONE*, **7**, e50429. 467 Mattern, T., Ellenberg, U., Houston, D.M. & Davis, L.S. (2007). Consistent foraging routes and benthic



469 Mattern, T., Ellenberg, U., Houston, D.M., Lamare, M., Davis, L.S., van Heezik, Y. & Seddon, P.J. 470 (2013). Straight line foraging in yellow-eyed penguins: new insights into cascading fisheries 471 effects and orientation capabilities of marine predators. PloS ONE, 8, e84381. 472 Moll, R.J., Millspaugh, J.J., Beringer, J., Sartwell, J. & He, Z. (2007). A new 'view' of ecology and 473 conservation through animal-borne video systems. Trends in Ecology & Evolution, 22, 660-668. 474 Moore, P.J. (1999). Foraging range of the Yellow-eyed penguin Megadyptes antipodes. 27, 49–58. 475 Moore, P.J. & Wakelin, M.D. (1997). Diet of the Yellow-eyed penguin Megadyptes antipodes, South 476 Island, New Zealand, 1991-1993. Marine Ornithology, 25, 17–29. 477 Moore, P.J., Wakelin, M.D., Douglas, M.E., McKinlay, B., Nelson, D. & Murphy, B. (1995). Yellow-eyed 478 penguin foraging study, south-eastern New Zealand, 1991-1993. 41. 479 Moreau, M., Siebert, S., Buerkert, A. & Schlecht, E. (2009). Use of a tri-axial accelerometer for 480 automated recording and classification of goats' grazing behaviour. Applied Animal Behaviour 481 Science, **119**, 158–170. 482 Noda, T., Kawabata, Y., Arai, N., Mitamura, H. & Watanabe, S. (2014). Animal-mounted 483 gyroscope/accelerometer/magnetometer: in situ measurement of the movement performance 484 of fast-start behaviour in fish. J Exp Mar Biol Ecol, 451. 485 Ponganis, P.J., St Leger, J. & Scadeng, M. (2015). Penguin lungs and air sacs: implications for 486 baroprotection, oxygen stores and buoyancy. Journal of Experimental Biology, 218, 720–730. 487 Queirós, A.M., Hiddink, J.G., Kaiser, M.J. & Hinz, H. (2006). Effects of chronic bottom trawling 488 disturbance on benthic biomass, production and size spectra in different habitats. Journal of Experimental Marine Biology and Ecology, **335**, 91–103. 489 490 R Development Core Team. (2008). R: A language and environment for statistical computing.

foraging behaviour in yellow-eyed penguins. Marine Ecology Progress Series, 343, 295-306.

Roberts, C.D., Stewart, A.L. & Struthers, C.D. (2015). The Fishes of New Zealand. Te Papa Press, 491 Wellington, New Zealand. 492 493 Sakamoto, K.Q., Takahashi, A., Iwata, T. & Trathan, P.N. (2009). From the Eye of the Albatrosses: A 494 Bird-Borne Camera Shows an Association between Albatrosses and a Killer Whale in the Southern Ocean. PLoS ONE, 4, e7322. 495 496 Sato, K., Naito, Y., Kato, A., Niizuma, Y., Watanuki, Y., Charrassin, J.B., Bost, C.-A., Handrich, Y. & Le Maho, Y. (2002). Buoyancy and maximal diving depth in penguins. Journal of Experimental 497 Biology, 205, 1189-1197. 498 499 Sato, K., Shiomi, K., Marshall, G., Kooyman, G.L. & Ponganis, P.J. (2011). Stroke rates and diving air 500 volumes of emperor penguins: implications for dive performance. Journal of Experimental 501 Biology, 214, 2854-2863. 502 Schratzberger, M. & Jennings, S. (2002). Impacts of chronic trawling disturbance on meiofaunal 503 communities. Marine Biology, 141, 991-1000. 504 Seddon, P.J., Ellenberg, U. & van Heezik, Y. (2013). Yellow-eyed penguin (Megadyptes antipodes). 505 Penguins: Natural History and Conservation (eds P. Garcia Borboroglu & P.D. Boersma), pp. 91-506 110. University of Washington Press, Seattle & London. 507 Sutton, G.J., Hoskins, A.J. & Arnould, J.P.Y. (2015). Benefits of Group Foraging Depend on Prey Type in 508 a Small Marine Predator, the Little Penguin. PLoS ONE, 10, e0144297. 509 Takahashi, A., Kokubun, N., Mori, Y. & Shin, H.-C. (2008). Krill-feeding behaviour of gentoo penguins 510 as shown by animal-borne camera loggers. Polar Biology, 31, 1291–1294. 511 Thiebot, J.-B., Arnould, J.P.Y., Gómez-Laich, A., Ito, K., Kato, A., Mattern, T., Mitamura, H., Noda, T., 512 Poupart, T., Qunitana, F., Raclot, T., Ropert-Coudert, Y., Sala, J.E., Seddon, P.J., Sutton, G.J., 513 Yoda, K. & Takahashi, A. Predator-borne video loggers emphasize the ecosystem role of jellyfish in across the southern oceans. Current Biology. 514

537

516 Marshall, G., Abernathy, K., Semmens, J., Hindell, M.A. & Arnould, J.P.Y. (2016). Dive 517 characteristics can predict foraging success in Australian fur seals (<em&gt;Arctocephalus 518 pusillus doriferus</em&gt;) as validated by animal-borne video. Biology Open, 5, 262 LP-271. 519 Watanabe, Y.Y. & Takahashi, A. (2013). Linking animal-borne video to accelerometers reveals prey capture variability. Proceedings of the National Academy of Sciences, 110, 2199–2204. 520 521 Watanuki, Y., Daunt, F., Takahashi, A., Newell, M., Wanless, S., Sato, K. & Miyazaki, N. (2008). 522 Microhabitat use and prey capture of a bottom-feeding top predator, the European shag, shown by camera loggers. Marine Ecology Progress Series, 356, 283–293. 523 524 Wilmers, C.C., Nickel, B., Bryce, C.M., Smith, J.A., Wheat, R.E. & Yovovich, V. (2015). The golden age 525 of bio-logging: how animal-borne sensors are advancing the frontiers of ecology. Ecology, 96, 1741-1753. 526 527 Wilson, R. & Liebsch, N. (2003). Up-beat motion in swinging limbs: new insights into assessing movement in free-living aquatic vertebrates. Marine Biology, 142, 537–547. 528 529 Wilson, R.P., Pütz, K., Peters, G., Culik, B.M., Scolaro, J.A., Charrassin, J.-B. & Ropert-Coudert, Y. 530 (1997). Long-term attachment of transmitting and recording devices to penguins and other seabirds. 25, 101-106. 531 Wilson, R.P., Simeone, A., Luna-Jorquera, G., Steinfurth, A., Jackson, S. & Fahlman, A. (2003). Patterns 532 533 of respiration in diving penguins: is the last gasp an inspired tactic? The Journal of experimental 534 biology, **206**, 1751–1763. 535 Wilson, R.P. & Wilson, M.T. (1989). A peck activity record for birds fitted with devices. Journal of Field Ornithology, 60, 104-108. 536

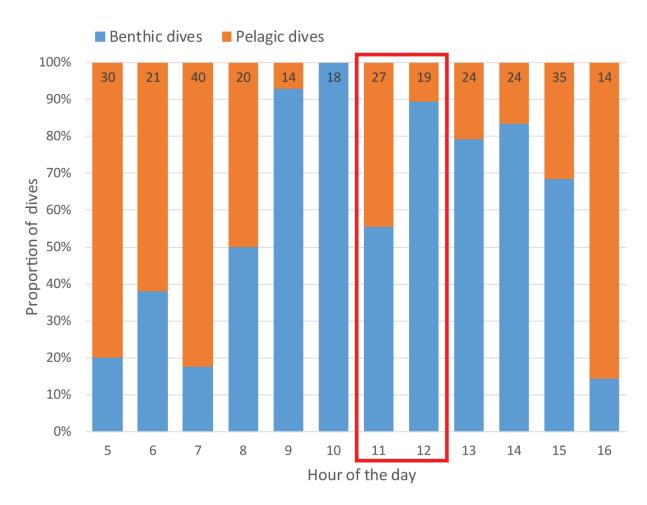
Volpov, B.L., Rosen, D.A.S., Hoskins, A.J., Lourie, H.J., Dorville, N., Baylis, A.M.M., Wheatley, K.E.,

539

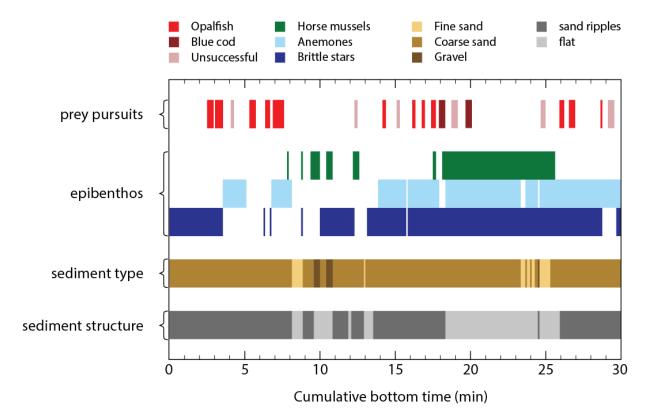
540

541

**Figure 1.** Proportion of benthic and pelagic dives throughout the Yellow-eyed penguin's foraging trip while fitted with a camera logger. Numbers at the top end of bars indicate number of dives performed during the corresponding hour. Red box indicates the hours during which continuous camera footage was recorded.



**Figure 2.** Composition of sediment structure, type and epibenthic communities during the bottom phases of 32 dives performed by a Yellow-eyed penguin fitted with a video camera logger. The x-axis indicates the cumulative time the penguin spent at the seafloor (29.9 minutes) during the 2 hours of camera operation. Coloured horizontal bars indicate duration of periods during which the penguin foraged over certain sediment structures and types, certain epibenthic taxa were present, as well as the length of prey pursuits (including pursuit outcome and prey species).

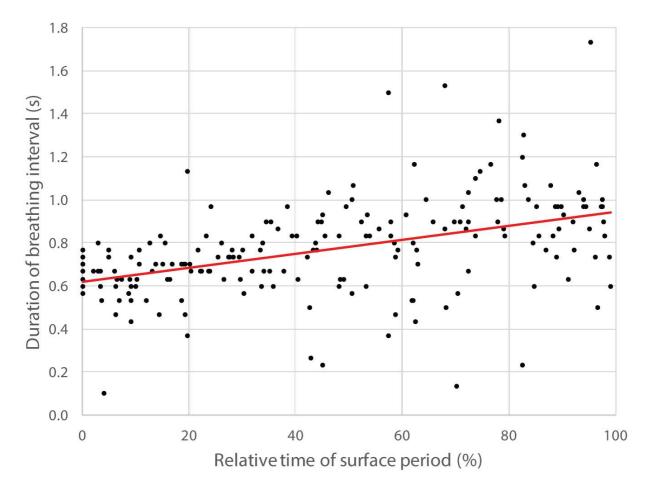


551

552

553

**Figure 3.** Increasing duration of breathing intervals (n=193) during the surface period after 10 randomly selected dives performed by a Yellow-eyed penguin. Note that the x-axis shows relative time to account for varying surface period durations. Red line indicates regression of data (see Results for details).



**Figure 4.** Flipper movements in a Yellow-eyed penguin during the descent, bottom and ascent phases of 10 randomly chosen benthic dives. Graphs in the upper row depict changes in flipper beat frequencies while the lower row consists of graphs showing flipper amplitude (i.e. maximum angle). Red lines indicate regression of the corresponding data (see Results for details). Note that x-axis shows relative durations of the dive phases to account for dive dependent time variations.

