

Behavioural inference from signal processing using animal-borne multi-sensor loggers: a novel solution to extend the knowledge of sea turtle ecology

Lorène Jeantet, Víctor Planas-Bielsa, Simon Benhamou, Sebastien Geiger, Jordan Martin, Flora Siegwalt, Pierre Lelong, Julie Gresser, Denis Etienne, Gaëlle Hiélard, et al.

▶ To cite this version:

Lorène Jeantet, Víctor Planas-Bielsa, Simon Benhamou, Sebastien Geiger, Jordan Martin, et al.. Behavioural inference from signal processing using animal-borne multi-sensor loggers: a novel solution to extend the knowledge of sea turtle ecology. Royal Society Open Science, 2020, 7 (5), pp.200139. 10.1098/rsos.200139. hal-02613340

HAL Id: hal-02613340

https://hal.science/hal-02613340

Submitted on 9 Sep 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Behavioural inference from signal processing using animal-borne multi-sensor loggers: a novel solution to extend the knowledge of sea turtle ecology

- 1 Lorène Jeantet¹, Víctor Planas-Bielsa², Simon Benhamou³, Sebastien Geiger¹, Jordan
- 2 Martin¹, Flora Siegwalt¹, Pierre Lelong¹, Julie Gresser⁴, Denis Etienne⁴, Gaëlle Hiélard⁵,
- 3 Alexandre Arque⁵, Sidney Regis¹, Nicolas Lecerf¹, Cédric Frouin¹, Abdelwahab
- 4 Benhalilou⁶, Céline Murgale⁶, Thomas Maillet⁶, Lucas Andreani⁶, Guilhem
- 5 Campistron⁶, Hélène Delvaux⁷, Christelle Guyon⁷, Sandrine Richard⁸, Fabien Lefebvre¹,
- 6 Nathalie Aubert¹, Caroline Habold¹, Yvon le Maho^{1,2}, Damien Chevallier¹.

7 8

- 9 ¹Institut Pluridisciplinaire Hubert Curien, CNRS–Unistra, 67087 Strasbourg, France.
- ²Centre Scientifique de Monaco, Département de Biologie Polaire, 8 quai Antoine Ier, MC
- 11 98000 Monaco.
- 12 ³Centre d'Écologie Fonctionnelle et Évolutive, CNRS, Montpellier, France & Cogitamus Lab.
- ⁴DEAL Martinique, Pointe de Jaham, BP 7212, 97274 Schoelcher Cedex, France.
- ⁵Office de l'Eau Martinique, 7 Avenue Condorcet, BP 32, 97201 Fort-de-France, Martinique,
- France.
- ⁶Association POEMM, 73 lot papayers, Anse a l'âne, 97229 Les Trois Ilets, Martinique.
- ⁷ DEAL Guyane, Rue Carlos Finley, CS 76003, 97306 Cayenne Cedex, France.
- ⁸Centre National d'Etudes Spatiales, Centre Spatial Guyanais, BP 726, 97387 Kourou Cedex,
- 19 Guyane.

2021

22 *Corresponding author: Lorène Jeantet lorene.jeantet@iphc.cnrs.fr

23

- 24 Keywords: supervised learning algorithms, accelerometer, sea turtle, animal-borne
- 25 camera, behavioural classification, marine ecology

Abstract

The identification of sea turtle behaviours is a prerequisite to predicting the activities and time-budget of these animals in their natural habitat over long term. However, this is hampered by a lack of reliable methods that enable the detection and monitoring of certain key behaviours such as feeding. This study proposes a combined approach that automatically identifies the different behaviours of free-ranging sea turtles through the use of animal-borne multi-sensor recorders (accelerometer, gyroscope and time-depth recorder), validated by animal-borne video-recorder data. We show here that the combination of supervised learning algorithms and multi-signal analysis tools can provide accurate inferences of the behaviours expressed, including behaviours that are of crucial ecological interest for sea turtles, such as feeding and scratching. Our procedure uses multi-sensor miniaturized loggers that can be deployed on free-ranging animals with minimal disturbance. It provides an easily adaptable and replicable approach for the long-term automatic identification of the different activities and determination of time-budgets in sea turtles. This approach should also be applicable to a broad range of other species and could significantly contribute to the conservation of endangered species by providing detailed knowledge of key animal activities such as feeding, travelling and resting.

1 Introduction

It is essential to assess the feeding behaviours of free-ranging animals in order to estimate their time budgets, and thus understand how these animals maximise their fitness [1,2]. However, investigating the foraging behaviour of sea turtles in their natural environment remains a significant challenge as it is impossible to obtain long-term behavioural data through visual observations alone. Although some studies have provided relevant information on sea turtle diet through post-mortem stomach content analysis or the deployment of animal-borne video-recorders [3–5], the proportion of time that sea turtles allocate to feeding activities on the long term remains unknown. Time-depth recorders (TDR) have been used to record the dive profiles and durations of free-ranging sea turtles and have provided insights into their underwater activities [6–8]. However, a number of authors have underlined the limits of focussing on dive profile, as foraging activity cannot be distinguished from transit or resting phases [9,10]. The joint use of TDR and video recorders revealed that the typical dive types described in [11,12] could not be associated with specific activities such as travelling, resting or foraging [13,14].

Devices combining miniaturised tri-axial accelerometers and TDR were described as a powerful tool to improve the identification of fine-scale behaviours in animals that cannot be easily monitored by visual observation [15–17]. Such devices have been deployed to study the behaviour and dive patterns of loggerheads (*Caretta caretta*, 18), green turtles (*Chelonia mydas*,19) and leatherbacks (*Dermochelys coriacea*, 20) during the inter-nesting period. However, the interpretation of the acceleration signals used in these studies to identify sea turtle behaviours in water was not validated by simultaneous visual observation, possibly resulting in misidentification and significant biases in the interpretation of the data.

A new approach was therefore necessary to reliably identify the underwater behaviours of free-ranging sea turtles without using direct visual observation (which is usually impossible) or video recordings, which are limited to short-term studies (a few hours) because of their high power consumption. Accelerometers permit the identification of feeding activity and time budget in marine animals such as seals and penguins by recording head movements that are likely to correspond to prey captures [21–23]. For the same purpose, accelerometers have been placed on the beak [24–27] or the top of the head [28] of sea turtles to record beak-openings and capture attempts. However, the position of the device was a significant disturbance for the individuals and could not be considered for long-term use (up to several

weeks). It was therefore crucial to develop a protocol for the long-term recording and identification of sea turtle feeding activities that minimizes disturbance to the animals whilst making optimal use of the subtle variations in data acquired by loggers that are mounted on the carapace rather than the head.

Further work is needed to validate the identification of sea turtle underwater behaviours by data acquired by animal-borne sensors. In particular, before attempting to provide new insights about the at-sea behaviours of sea turtles in natural conditions, one needs is to automatically and correctly identify these behaviours, including those that are hard to detect but play a key role such as feeding, from data acquired in a way that minimizes the disturbance of equipped animal. The aim of our study is therefore to develop a new approach fulfilling this need. In this framework, we will use the results we obtained about turtles' behaviours only to illustrate the output of our approach without giving them any biological significance. Although sea turtle behaviours have mainly been inferred from combined acceleration and depth data, the additional use of a gyroscope (which records angular velocity) can provide further relevant information in remote behavioural identification [29– 31]. Thus, we deployed loggers combining an accelerometer, a gyroscope and a TDR on the carapace of free-ranging immature green turtles. This equipment was linked to a videorecorder that was mounted in the logger device to provide visual evidence that could validate logger interpretations of behaviours, given that our approach ultimately aims to infer behaviours solely through logger use. Surface behaviours were identified separately from depth data. The study tested a set of methods to infer diving behaviours from the signals provided by the accelerometer, gyroscope and TDR, including automatic segmentation and supervised learning algorithms. The validity of our approach was tested through the use of confusion matrices and by comparing the inferred activity budgets with those obtained from video recordings.

118119

120

121

94

95

96

97

98

99

100

101

102

103

104

105

106

107

108

109

110

111112

113

114

115

116

117

2 Material and Methods

2.1 Ethics statements

This study meets the legal requirements of the countries in which the work was carried out and follows all institutional guidelines. The protocol was approved by the "Conseil National de la Protection de la Nature" (http://www.conservation-nature.fr/acteurs2.php?id= 11), and the French Ministry for Ecology, Sustainable Development and Energy (permit number: 2013154-0037), which acts as an ethics committee in Martinique. The fieldwork was carried out in strict accordance with the recommendations of the Prefecture of Martinique in order to minimize the disturbance of animals (Authorisation n°201710-0005).

2.2 Data collection from free-ranging green turtles

The field work was carried out from February 2018 to May 2019 in Grande Anse d'Arlet (14°50'N, 61°09'W), Martinique, France. We deployed CATS (Customized Animal Tracking Solutions, Germany) devices for periods ranging from several hours to several days on free-ranging immature green turtles. A CATS device is comprised of a video-recorder (1920 x 1080 pixels at 30 FPS) combined with a tri-axial accelerometer, a tri-axial gyroscope and a TDR (Supplementary Material, Fig. 1). The maximum battery capacity was considered to provide a recording capacity of 18 h of video footage and 48 h for other data. These devices were programmed to record acceleration and angular velocity (gyroscope) at a frequency of 20 or 50 Hz according to the recording capacity of the logger (the 50 Hz data were subsampled at 20 Hz using a linear interpolation to homogenise the sample). Depth was recorded at 1 Hz using a pressure sensor with a range from 0 to 2000 m and 0.2m accuracy.

The relatively shallow depths of the area allowed free divers to capture the turtles manually, as described in Nivière et al. (2018) [32]. Once an individual had been caught, it was placed on a boat and identified by scanning its PIT (Passive Integrated Transponder) or tagged with a new PIT if it was unknown. It was then weighed and its carapace length was measured (Supplementary Material, Table S1). The device was attached to the carapace using four suction cups. Air was manually expelled from the cups, which were held in place by the use of a galvanic timed release system. The eventual dissolving of these releases by sea water and the positive buoyancy of the device (23.3 x 13.5 x 4 cm for 0.785 kg) led to the remote release of the device several hours later. Devices were recovered by geolocation of their Argos SPOT-363A tag (MK10, Wildlife Computers Redmond, WA, USA) with a goniometer (RXG-134, CLS, France). Instruments were deployed on 37 individuals, but complete datasets including video, acceleration, gyroscope and depth values were only recovered for 13 individuals (Supplementary Material, Table S1).

2.3 Processing of video data and behavioural labelling

The video footage was watched to identify the various behaviours and determine their starting and ending times to the closest 0.01 s. Acceleration, angular velocity and depth data corresponding to each behavioural phase were visualized using R software (version 3.5.3) and the package *rblt* (Fig 1 & 2; 33). The 46 resulting behaviours were clustered into categories according their similarities (the definition of the various behaviours is available in Supplementary Material, Table S2). We retained seven main expressed categories for the multi-sensor signals, namely "Breathing", "Feeding", "Gliding", "Swimming", "Resting", "Scratching" and "Staying at the surface". All other behaviours were very infrequent and were grouped in an eighth category labelled "Other".

2.4 Analysis of the angular velocity and acceleration data

The device was installed on the carapace in a tilted position along a longitudinal axis to obtain video images of the head. This results in biased values of accelerations and angular speeds for the surge (i.e. back-to-front) and heave (bottom-to-top) body axes, which therefore had to be corrected (see R-script in Supplementary Material). The static acceleration vector (i.e. the component due to gravity) $\overline{\bf a}=(\overline{a}_x,\overline{a}_y,\overline{a}_z)$ was obtained by separately averaging the acceleration values $(a_x, a_y \text{ and } a_z)$ on the surge, sway (right-to-left) and heave axes, respectively, over a centred running temporal window set to $\Delta t=2$ s. which was the smallest window resulting in a norm, $\|\overline{\bf a}\|$ that remains close to 1 g (9.98 m/s²) for almost all measures. The Dynamic Body Acceleration was then computed as DBA = $\sqrt{\bf d}^2$, where $\bf d=a-\overline{a}$ is the dynamic acceleration vector [34]. Similarly the Rotational Activity was computed as RA = $\sqrt{\bf g}^2$, where $\bf g=(g_x,g_y,g_z)$ is the angular velocity $(g_x,g_y,and g_z)$ correspond respectively to the values of roll, pitch and yaw per unit time provided by the gyroscope).

2.5 Segmentation of the multi-sensor dataset

The automatic identification of the labelled behaviours from the multi-sensor signals required the segmentation of the dataset into homogenous behavioural bouts with respect to a given variable. We started by relying on the depth data to distinguish the dives, defined as depths exceeding 0.3 m for at least 5 s, from the surface periods. We attributed the surface periods to either "Breathing" or "Staying at the surface", according to whether the turtle remained at the surface for less or more than 6 s, respectively. We then distinguished between the various

possible diving behaviours by using a changepoint algorithm, the Pruned Exact Linear Time (PELT) algorithm (R package changepoint; 35), in which the "pen.value" parameter, which corresponds to the additional penalty in the cost function for each additional partition of the data, can be manually adjusted. We tested different values and retained those which resulted in the best balance between obtaining homogenous behavioural bouts and limiting oversegmentation. We first detected depth changes over 3 s of each dive (function cpt.mean, penalty = "Manual", pen.value = 5) to obtain segments which were labelled as "ascending", "descending" or "flat" depending on whether the vertical speed was > 0.1 m.s⁻¹, <-0.1 m.s⁻¹ or between these two values, respectively. These ascending and descending segments were further segmented based on the DBA mean and variance (function cpt.meanvar, penalty = "Manual", pen.value = 50) in order to distinguish between the swimming and gliding phases of these segments. The green turtle is a grazing herbivore which mainly feed on seagrass and algae [36]. The head movements occurring during feeding activities are easily detected by gyroscopes and/or accelerometers set directly on the head, but are rarely detected when these sensors are placed on the carapace. We did however note that the carapace tended to display pitch oscillations when the turtle pulled on the seagrass, an activity that we refer to hereafter as "Grabbing" (Fig 2). Accordingly, we further segmented the "flat" segments based on the variance of g_y (angular speed in the animal's sagittal plane; function *cpt.var*, penalty = "Manual", pen.value = 20) to pinpoint this behaviour. Each segment was then labelled as either the behavioural category that was expressed for at least 3/5 of its duration, or as "Transition" if several behaviours were involved with none of them occurring for 3/5 of the behavioural bout. Thus, the overall procedure classified multi-sensor signals into nine categories comprised of surface behaviours ("Breathing" and "Staying at the surface") which were identified using depth data alone, diving behaviours ("Feeding, "Gliding", "Resting", "Scratching" and "Swimming") and also "Other" and "Transition", for which supervised learning algorithms were required.

214

215

188

189

190

191

192

193

194

195

196

197

198

199

200

201

202

203

204

205

206

207

208

209

210

211

212

213

2.6 Identification of the diving behaviours by supervised learning algorithms

We trained five supervised machine learning algorithms – (1) Classification And Regression Trees (CART), (2) Random Forest (RF), (3) Extreme Gradient Boosting (EGB), (4) Support Vector Machine (SVM), and (5) Linear Discriminant Analysis (LDA) – to associate the eight diving behaviour categories with the corresponding patterns of different input variables. They are the most commonly used classifiers in behaviour recognition and are considered to be relevant in ecology studies [17,37,38]. These algorithms were applied to our data using the R packages *rpart* [39] for CART, *randomForest* [40] for RF (*n*=300, mtry=14), *xgboost* [41] for EGB (num class=8, eta=0.3, max depth=3), *e1071* [42] for SVM and *MASS* [43] for LDA.

221

222

223

224

225

226

227

228

229

230

231

232

233

234

235

236

237

238

239

240

241

242

243

244

245

246

247

248

249

250

251

252

253

For each segment, the algorithms were fed with four descriptive statistics (mean, minimum, maximum, and variance) computed for the three linear acceleration values (a_x , a_y and a_z), for the three angular speeds values (g_x, g_y) and g_z), and for DBA and RA. We also included the difference between the last and first depth values, and the duration of each segment. The fact that "Feeding" was characterised by high-frequency oscillations, in particular in terms of pitch speed (Fig 2), but also (although less obviously) in terms of roll speed and surge/sway accelerations, enabled us to distinguish this behaviour from the others. To do so, we filtered the raw values of g_x , g_y , a_x and a_y through a running window of 1 s and subtracted the obtained smoothed values from the respective raw values, then calculated the local mean of the obtained high frequency signals. We then computed the squared differences between the high frequency values and their respective local means in order to characterize these oscillations. The mean and the maximum value of these squared differences for each segment were then added to the list of variables used to feed the algorithms, i.e. 42 variables for each segment. Such a number of variables may be characterised numerous correlations. However, machine learning algorithms are less sensitive than classical regression methods to correlation in the explanatory variables. Nevertheless, for a simpler interpretation purpose, we looked for some possible reduced set of variables that may reach the same accuracy as the full data set, but we did not find any convincing one that had a close performance to the full data set. As the focus was more on predictability than interpretability (as is usual the case in machine learning), we kept all the 42 variables.

2.7 Validation of the automatic behavioural inferences

To estimate the ability of our procedure to correctly infer the behaviours of sea turtles based on acceleration, angular velocity and depth data, we repeatedly performed 2/3:1/3 splits of the sample of 13 individuals, with nine individuals retained for the learning phase and the remaining four individuals used to validate the outcome. From the 715 possible combinations, we retained the 358 combinations in which "Feeding" and "Scratching" were not underrepresented in the training dataset (i.e. when more than 60% of total feeding and scratching segments were present, i.e. 1145 and 868, respectively). Nevertheless, the number of "Feeding" and "Scratching" segments was much lower than those attributed to "Resting" and

"Swimming" (17325 and 9795 segments, respectively). As an unbalanced training dataset can hinder the performance of supervised learning algorithms [44], we set an upper limit at 1000 segments per behaviour for the training dataset. These segments were randomly selected for the over-expressed categories at each training trial.

For each trial, we evaluated the efficiency of the different methods by computing the number of well-identified behaviours (true positive, TP, and true negative, TN) and of behaviours considered to be misclassified (false negative, FN, and false positive, FP) into a confusion matrix. We calculated three indicators for each behaviour: (1) "Sensitivity" = TP/(TP+FN), also called *true positive rate*, *hit rate* or *recall*, measures the ability of a method to detect the target behaviour among other behaviours; (2) "Precision" = TP/(TP+FP), also called *positive predictive value*, measures the ability of a method to correctly identify the target behaviour; and (3) "Specificity" = TN/(TN+FP), also called *selectivity* or *true negative rate*, measures the ability of a method to avoid wrongly considering other behaviours as the target behaviour. We also computed "Accuracy" = (TP+TN)/(TP+TN+FP+FN), which measures the ability of a method to correctly identify all behaviours as a whole.

Furthermore, to possibly improve the performance and/or minimise the variance of behavioural inferences, we also relied on the "Ensemble Methods" [45,46], which consisted of combining the results obtained with the five supervised machine learning algorithms. We tested two such methods. The first was the "Voting Ensemble" (VE), which retained the most frequently predicted behaviour. The second involved a "Weighted Sum" (WS), where weights were given to the different predicted behaviours, based on "Precision" (weighting based on Sensitivity and Specificity was also tested but gave poor results). In order to highlight the best method to automatically identify the diving behaviours and particularly the feeding behaviours, we used Anova to compare the mean global accuracy obtained for the 375 combinations of the seven classifiers (CART, SVL, LDA, RF, EGB, VE and WE). As the result of the Anova showed significant effects, we ran pairwise comparisons of mean performance using the Tukey HSD test.

Finally, the individual activity budgets were inferred by computing the proportion of time involved in the various surface behaviours ("Breathing" and "Staying at the surface") inferred from depth data, and the proportion of time dedicated to diving behaviours ("Feeding", "Gliding", "Other", "Resting", "Scratching", and "Swimming"), inferred using the best classifier (Fig. 3). The inferred activity budgets were compared to those observed in video recordings.

3 Results

A total of 66.2 hours of video were recorded, with a maximum of 14.6 hours for one individual (Table 1). The seven specific behavioural categories retained for the analysis ("Breathing", "Feeding", "Gliding", "Resting", "Scratching", "Staying at the surface" and "Swimming") represented 99% of the total duration. Only the two shortest deployments were not associated with a feeding event while the maximum duration of feeding represented only 8% of the recording time of the individual. The catching of jellyfish was observed only occasionally in three individuals. This behaviour represented only 0.1% of the total feeding duration of the thirteen individuals, whilst the rest of the feeding time for those individuals was used for grazing on seagrass. For the others, feeding consisted only of grazing on seagrass. "Scratching" was particularly expressed by one turtle, and represented 13% of its observed time.

The seven classifiers identified the five specific behavioural categories on which we focused ("Feeding", "Gliding", "Resting", "Scratching" and "Swimming") and two additional categories, "Transition" and "Other", with an accuracy ranging from 0.91 to 0.95. The highest score was obtained with WS and the lowest one with SVM. The Tuckey HSD test indicated that the RF, VE and EGB outputs were not significantly different (0.935, 0.932 and 0.932, respectively). All classifiers identified the behavioural category with a low false positive rate (< 0.1 for the best classifiers; Fig.4). Few segments were wrongly identified as "Feeding" with the WS method, which thus obtained the lowest false positive rate (with respect to other classifiers) for this behaviour. The best true positive rates, for the seven classifiers, were obtained in the "Scratching" category despite its low occurrence in the dataset, meaning that this behaviour was relatively well identified when it occurred.

The activity budget, representing the percentage of the total mean time allocated to each behavioural category, showed similar proportions between the predictions and the observations (Fig. 5 & 6). This result highlights the ability of our method and the WS model to predict the behaviours of immature green turtle in natural conditions. The main differences between the observed and predicted activity budgets were seen in the "Resting" and "Swimming" behaviours (Fig. 5 & 6). These differences were small and represented less than 3% of the total observed time (Table 2). "Feeding" and "Scratching" were under-represented in our models and consequently their difference between predicted—actual time represent roughly 1% of the total observed time. Their low expression for some individuals led to an important percentage difference with respect to the observed time of the behaviour even if they were predicted in small proportion. The results obtained for each individual are available

in supplementary material (Supplementary Material, Table S3). With a very low true positive rate, the predicted time of "Transition" represented on average 0.2% of the total observation time. Thus the overall procedure was able to reliably infer the seven mainly expressed behaviours of the immature green turtles.

326

327

328

329

330

331

332

333

334

335

336

337

338

339

340

341

342

343

344

345

346

347

348

349

350

351

352

353

354

322

323

324

325

4 Discussion

This is the first study to validate the use of acceleration, gyroscope and TDR signals for inferring free-ranging green turtle behaviours. In previous studies, carapace-mounted accelerometers were used to describe swimming behaviours and buoyancy regulation in sea turtles [19,20,47,48] in specific contexts where signals associated to "Swimming" and "Gliding" could be visually identified, or were used to estimate sea turtle activity levels in terms of DBA [18,49]. The possibility to rely on accelerometers and other carapace-mounted sensors such as TDRs and gyroscopes to infer behaviours of free-ranging sea turtles had not been explored in detail until now due to the lack of a validation process, which is critically important for this kind of approach [50]. The validation process described in the present study has enabled us to provide an overall procedure permitting the reliable inference of the seven most commonly expressed behaviours of the free-ranging green turtle (namely "Breathing", "Feeding", "Gliding", "Resting", "Scratching", "Staying at the Surface" and "Swimming"), and thus inferring the fine-scale activity budgets of animals whose populations are currently under anthropogenic pressures which jeopardize their future [51,52]. This inference is essential if we wish to compare how these animals allocate their time between different activities according to natural and anthropogenic pressures such as available resources, environmental changes or tourism. When combined with GPS data, this protocol could identify the areas where sea turtles concentrate their activities and thus help to delineate protected areas in order to limit human disturbances.

We tested seven classifiers (LDA, SVM, CART, RF, EGB, VE and WS) to compare their strengths and weaknesses in automatic behavioural identification based on TDR, acceleration and gyroscopic data. The classifiers identified the seven behavioural classes with a global accuracy ranging from 0.91 to 0.95, which is comparable to the accuracy reached in other similar studies [17,53,54]. The WS classifier performed better than the base and VE classifiers: clearly, assigning precision-based weights to the base classifier' predictions improved the behavioural classification. The decrease we observed in the false positive rate for rare behaviours through the use an ensemble method for "Feeding" in this study has also

been highlighted by Brewster et al. (2018) [37]. Ensemble methods are mainly used because they reduce the variance of behaviour classification [53,55] and thus increase the global accuracy. However, they involve a higher computational cost and require a reliable setting up of base learners.

355

356

357

358

359

360

361

362

363

364

365

366

367

368

369

370

371

372

373

374

375

376

377

378

379

380

381

382

383

384

385

386

387

388

The use of supervised machine-learning has become common to automatically identify behaviours from data provided by animal-borne loggers [17,50,56]. Indeed, the development of fast personal computers and of free user-friendly computing libraries made it possible to easily apply these 'black box' algorithms to huge amounts of data. The machine-learning approach has thus turned out to be a very powerful tool for identifying well-characterized behaviours (in terms of signal) such as locomotion [56–58] and resting [59–61]. However, it appears to be rather inefficient when seeking to identify behaviours with confusing signal characteristics. Examples include feeding and grooming in pumas [62], pecking in plovers [63] or foraging in fur seals [64]. Although one could expect that feeding machine-learning algorithms with big data should provide the most accurate predictive rules [16,65,66], Wilson et al. (2018) [67] showed that a classification method based on a good understanding and careful examination of the acceleration signal actually gives better results in terms of computational time and of accuracy than non-optimized machine learning. Accordingly, the mixed approach developed in this study fed machine-learning algorithms with a number of derived signals which were specifically elaborated to pinpoint specific hard-to-detect behaviours when alternative simpler means based on a single or a few parameters appeared to be effective. This method allowed us to identify key behaviours such as feeding and scratching, which had previously been either misidentified or not identified at all due to the lack of discriminative signals in the raw data obtained from raw acceleration and/or gyroscopic data obtained with loggers fixed to the carapace of the turtle. Although our choice of derived signals makes our approach specific to sea turtles, this principle can be applied to numerous species if the different signals are considered with care before the study.

When carrying out automatic behavioural identification from multi-sensor data using supervised learning algorithms, one of the main difficulties is the segmentation of the multi-sensor data to obtain homogeneous segments that are representative of the various behavioural categories. To date, most studies divided the multi-sensor data into segments using fixed-time segments [68–70] or a sliding sample window with a fixed length [38,71]. However, several studies testing the size of the window showed that it influences classification accuracy and the identification of short behaviours [53,72–74]. Indeed, an individual can express both short and long behaviours, such as burst swimming in lemon

sharks or a prey capture in Adélie penguins compared to normal swimming behaviour [37,75]. Whilst the use of long fixed segments dramatically increases the proportion of inhomogeneous segments, using short segments may prevent the detection of certain key signals such as low-frequency oscillations. A hierarchical, adapted segmentation procedure therefore seems to be a more judicious choice. This consists of splitting behaviours into groups based on signals that are easily interpretable in a dichotomic way (variables such as depth were used to attain this in our study). A change-point algorithm can be used to achieve a more specific segmentation based on other signals, with a possible ad hoc adjustment of the contrast is required to evaluate whether two successive values do or do not belong to the same segment (such as the manual penalty of the PELT algorithm). In this paper, we demonstrate this approach for the green turtle (Fig. 3, R-script in Supplementary Material), but there is no reason it could not be easily adapted for other species. This will certainly necessitate the identification of the optimal hyper-parameters as well as the informative signals for the segmentation according to the species, but the approach of combining automated segmentation and machine learning methods with well thought out descriptive variables should apply as well.

The approach we proposed thus offers promising perspectives for inferring behaviours of animals that cannot be easily observed in the wild though the automatic analysis of large amounts of raw data acquired over long periods by miniaturised (low disturbance) loggers such as high-frequency tri-axial accelerometers and gyroscopes. It provides a number of adaptable principles that enable the efficient use of machine learning algorithms to automatically identify fine-scale behaviours in sea turtles, and may be used for a wide range of species. The automated and reliable identification of the various behaviours permits a rapid inference of the time budget of the animals under study. Identifying how much time the studied animals dedicate to activities such as feeding, travelling and resting can be of relevance when seeking to understand how individuals attempt to maximise their fitness in a given environment. This approach could therefore be a key tool in understanding the ecology of endangered species and make a significant contribution to their conservation.

5 Data accessibility statement

The R-script to visualize the raw acceleration, gyroscope and depth profile associated with the observed behaviours of the immature green turtles have been uploaded as part of the supplementary material. The same is true for the R-script to automatically identify sea turtle

- behaviour from the labelled data. The datasets containing the acceleration, gyroscope and
- depth recordings of the 13 immature green turtles as well as their observed behaviours are
- available within the Dryad Digital Repository: https://doi.org/10.5061/dryad.hhmgqnkd9.
- 425 URL for reviewer:
- 426 https://datadryad.org/stash/share/j3x2UusNTI1OWqLsJ1mUOq7wMm3t3PAGMzwAMuYM
- 427 FO0

429 6 Author contributions

- DC, HD and SR contributed conception and design of the study. LJ, DC, JM, FS, JG, DE,
- 431 GH, AA, YLM, PL, SR, FL, NL, NA, CF, FV, ES, MB, FF, AB, CM, TM, MD, CG, SB and
- AB contributed to data acquisition. LJ, SG and SB performed the data acceleration analysis
- and LJ and VPB the statistical analysis. LJ wrote the first draft of the manuscript and SB,
- VPB, FS and DC contributed critically to subsequent versions.

435

436 7 Funding

- 437 This study was co-financed by the FEDER Martinique (European Union, Conventions
- 438 2012/DEAL/0010/4-4/31882 & 2014/DEAL/0008/4-4/ 32947), DEAL Martinique
- 439 (Conventions 2012/DEAL/0010/4-4/31882 & 2014/DEAL/0008/4-4/32947), the ODE
- 440 Martinique (Convention 014-03-2015), the CNRS (Subvention Mission pour
- 1'Interdisciplinarité), the ERDF fund (Convention CNRS-EDF- juillet2013) and the Fondation
- de France (Subvention Fondation Ars Cuttoli Paul Appell). Lorene Jeantet's Ph.D scholarship
- was supported by DEAL Guyane and CNES Guyane.

444

445

8 Acknowledgments

- 446 This study was carried out within the framework of the Plan National d'Action Tortues
- Marines de Martinique et Guyane Française. The authors also appreciate the support of the
- 448 ANTIDOT project (Pépinière Interdisciplinaire Guyane, Mission pour l'Interdisciplinarité,
- CNRS). The authors would like to thank the DEAL Martinique and Guyane, the CNES, the
- ODE Martinique, the ONCFS Martinique and Guyane, the ONEMA Martinique and Guyane,
- 451 the SMPE Martinique and Guyane, the ONF Martinique, the PNR Martinique, the Surfrider
- 452 Foundation, Carbet des Sciences, Plongée-Passion, the Collège Cassien Sainte-Claire and the
- 453 Collège Petit Manoir for their technical support and field assistance. We are also grateful to

- 454 the numerous volunteers and free divers for their participation in the field operations. Results
- obtained in this paper were computed on the vo.grand-est.fr virtual organization of the EGI
- 456 Infrastructure through IPHC resources. We thank EGI, France Grilles and the IPHC
- Computing team for providing the technical support, computing and storage facilities. We are
- also grateful to the three anonymous reviewers for their helpful corrections and comments.

- 461 9 References
- 462 1. Stephens DW, Krebs JR. 1987 Foraging Theory. Princeton University Press.
- 463 (doi:10.2307/2260475)
- 464 2. Stephens DW, Brown JS, Ydenberg RC. 2007 Foraging: behavior and ecology.
- 465 University of Chicago Press.
- 466 3. Colman LP, Sampaio CLS, Weber MI, de Castilhos JC. 2014 Diet of Olive Ridley Sea
- Turtles, Lepidochelys olivacea, in the Waters of Sergipe, Brazil. Chelonian Conserv.
- 468 *Biol.* **13**, 266–271. (doi:10.2744/CCB-1061.1)
- 469 4. Arthur K, O'Neil J, Limpus CJ, Abernathy K, Marshall G. 2007 Using animal-borne
- imaging to assess green turtle (Chelonia mydas) foraging ecology in Moreton Bay,
- 471 Australia. *Mar. Technol.* **41**, 9–13.
- 472 5. Wildermann NE, Barrios-Garrido H. 2013 First Report of Callinectes sapidus
- 473 (Decapoda: Portunidae) in the Diet of Lepidochelys olivacea. Chelonian Conserv.
- 474 *Biol.* **11**, 265–268. (doi:10.2744/ccb-0934.1)
- 475 6. Lennox RJ et al. 2017 Envisioning the Future of Aquatic Animal Tracking:
- 476 Technology, Science, and Application. *Bioscience* 67, 884–896.
- 477 (doi:10.1093/biosci/bix098)
- 478 7. Hussey NE et al. 2015 Aquatic animal telemetry: A panoramic window into the
- 479 underwater world. *Science* (80-.). **348**, 1255642. (doi:10.1126/science.1255642)
- 480 8. Houghton JDR, Broderick AC, Godley BJ, Metcalfe JD, Hays GC. 2002 Diving
- behaviour during the internesting interval for loggerhead turtles Caretta caretta nesting
- 482 in Cyprus. *Mar. Ecol. Prog. Ser.* **227**, 63–70. (doi:10.3354/meps227063)
- 483 9. Chambault P, de Thoisy B, Kelle L, Berzins R, Bonola M, Delvaux H, Le Maho Y,
- 484 Chevallier D. 2016 Inter-nesting behavioural adjustments of green turtles to an
- 485 estuarine habitat in French Guiana. Mar. Ecol. Prog. Ser. 555, 235–248.
- 486 (doi:10.3354/meps11813)

- 487 10. Chambault P et al. 2016 The influence of oceanographic features on the foraging
- behavior of the olive ridley sea turtle Lepidochelys olivacea along the Guiana coast.
- 489 *Prog. Oceanogr.* **142**, 58–71. (doi:10.1016/j.pocean.2016.01.006)
- 490 11. Hays GC, Adams CR, Broderick AC, Godley BJ, Lucas DJ, Metcalfe JD, Prior AA.
- 491 1999 The diving behaviour of green turtles at Ascension Island. *Anim. Behav.* **59**, 577–
- 492 586. (doi:10.1006/anbe.1999.1326)
- 493 12. Hochscheid S, Godley BJ, Broderick AC, Wilson RP. 1999 Reptilian diving: Highly
- variable dive patterns in the green turtle Chelonia mydas. Mar. Ecol. Prog. Ser. 185,
- 495 101–112. (doi:10.3354/meps185101)
- 496 13. Seminoff JA, Jones TT, Marshall GJ. 2006 Underwater behaviour of green turtles
- 497 monitored with video-time-depth recorders: What's missing from dive profiles? *Mar*.
- 498 *Ecol. Prog. Ser.* **322**, 269–280. (doi:10.3354/meps322269)
- 499 14. Thomson JA, Heithaus MR, Dill LM. 2011 Informing the interpretation of dive profiles
- using animal-borne video: A marine turtle case study. J. Exp. Mar. Bio. Ecol. 410, 12–
- 501 20. (doi:10.1016/j.jembe.2011.10.002)
- 502 15. Laich AGG, Wilson RP, Quintana F, Shepard ELC. 2010 Identification of imperial
- 503 cormorant Phalacrocorax atriceps behaviour using accelerometers. Endanger. Species
- 504 Res. 10, 29–37. (doi:10.3354/esr00091)
- 505 16. Graf PM, Wilson RP, Qasem L, Hackländer K, Rosell F. 2015 The use of acceleration
- to code for animal behaviours; a case study in free-ranging Eurasian beavers Castor
- 507 fiber. *PLoS One* **10**, 1–18. (doi:10.1371/journal.pone.0136751)
- 508 17. Nathan R, Spiegel O, Fortmann-Roe S, Harel R, Wikelski M, Getz WM. 2012 Using
- tri-axial acceleration data to identify behavioral modes of free-ranging animals: general
- 510 concepts and tools illustrated for griffon vultures. J. Exp. Biol. 215, 986–96.
- 511 (doi:10.1242/jeb.058602)
- 512 18. Fossette S, Schofield G, Lilley MKS, Gleiss AC, Hays GC. 2012 Acceleration data
- reveal the energy management strategy of a marine ectotherm during reproduction.
- 514 Funct. Ecol. **26**, 324–333. (doi:10.1111/j.1365-2435.2011.01960.x)
- 515 19. Yasuda T, Arai N. 2009 Changes in flipper beat frequency, body angle and swimming
- speed of female green turtles Chelonia mydas. Mar. Ecol. Prog. Ser. 386, 275–286.
- 517 (doi:10.3354/meps08084)
- 518 20. Fossette S, Gleiss AC, Myers AE, Garner S, Liebsch N, Whitney NM, Hays GC,
- Wilson RP, Lutcavage ME. 2010 Behaviour and buoyancy regulation in the deepest-
- diving reptile: the leatherback turtle. J. Exp. Biol. 213, 4074–4083.

- 521 (doi:10.1242/jeb.048207)
- 522 21. Viviant M, Trites AW, Rosen DAS, Monestiez P. 2009 Prey capture attempts can be
- detected in Steller sea lions and other marine predators using accelerometers. *Polar*
- 524 *Biol.* **33**, 713–719. (doi:10.1007/s00300-009-0750-y)
- 525 22. Watanabe YY, Ito M, Takahashi A. 2014 Testing optimal foraging theory in a penguin-
- 526 krill system. *Proc. R. Soc. B Biol. Sci.* **281**. (doi:10.1098/rspb.2013.2376)
- 527 23. Gallon S, Bailleul F, Charrassin JB, Guinet C, Bost CA, Handrich Y, Hindell M. 2013
- Identifying foraging events in deep diving southern elephant seals, Mirounga leonina,
- using acceleration data loggers. *Deep. Res. Part II Top. Stud. Oceanogr.* **88–89**, 14–22.
- 530 (doi:10.1016/j.dsr2.2012.09.002)
- 531 24. Fossette S, Gaspar P, Handrich Y, Maho Y Le, Georges JY. 2008 Dive and beak
- movement patterns in leatherback turtles Dermochelys coriacea during internesting
- 533 intervals in French Guiana. J. Anim. Ecol. 77, 236–246. (doi:10.1111/j.1365-
- 534 2656.2007.01344.x)
- 535 25. Hochscheid S, Maffucci F, Bentivegna F, Wilson RP. 2005 Gulps, wheezes, and sniffs:
- How measurement of beak movement in sea turtles can elucidate their behaviour and
- ecology. J. Exp. Mar. Bio. Ecol. **316**, 45–53. (doi:10.1016/j.jembe.2004.10.004)
- 538 26. Myers AE, Hays GC. 2006 Do leatherback turtles Dermochelys coriacea forage during
- the breeding season? A combination of data-logging devices provide new insights.
- 540 *Mar. Ecol. Prog. Ser.* **322**, 259–267. (doi:10.3354/meps322259)
- 541 27. Okuyama J, Kawabata Y, Naito Y, Arai N, Kobayashi M. 2009 Monitoring beak
- movements with an acceleration datalogger: A useful technique for assessing the
- feeding and breathing behaviors of sea turtles. Endanger. Species Res.
- 544 (doi:10.3354/esr00215)
- 545 28. Okuyama J et al. 2013 Ethogram of Immature Green Turtles: Behavioral Strategies for
- Somatic Growth in Large Marine Herbivores. PLoS One 8(6): e65783.
- 547 (doi:10.1371/journal.pone.0065783)
- 548 29. Wilson M, Tucker AD, Beedholm K, Mann DA. 2017 Changes of loggerhead turtle (
- 549 Caretta caretta) dive behavior associated with tropical storm passage during the inter-
- nesting period. *J. Exp. Biol.* **220**, 3432–3441. (doi:10.1242/jeb.162644)
- 551 30. Tyson RB, Piniak WED, Domit C, Mann D, Hall M, Nowacek DP, Fuentes MMPB.
- 552 2017 Novel Bio-Logging Tool for Studying Fine-Scale Behaviors of Marine Turtles in
- Response to Sound. *Front. Mar. Sci.* **4**. (doi:10.3389/fmars.2017.00219)
- 554 31. Noda T, Okuyama J, Koizumi T, Arai N, Kobayashi M. 2012 Monitoring attitude and

- dynamic acceleration of free-moving aquatic animals using a gyroscope. Aquat. Biol.
- **16**, 265–276. (doi:10.3354/ab00448)
- 557 32. Nivière M et al. 2018 Identification of marine key areas across the Caribbean to ensure
- the conservation of the critically endangered hawksbill turtle. *Biol. Conserv.* **223**, 170–
- 559 180. (doi:10.1016/j.biocon.2018.05.002)
- 560 33. Geiger S. 2019 Package 'rblt'., 1–12.
- 34. Wilson RP et al. 2020 Estimates for energy expenditure in free-living animals using
- acceleration proxies: A reappraisal. J. Anim. Ecol. 89, 161-172. (doi:10.1111/1365-
- 563 2656.13040)
- 564 35. Killick R, Fearnhead P, Eckley IA. 2012 Optimal detection of changepoints with a
- linear computational cost. J. Am. Stat. Assoc. 107, 1590–1598.
- 566 (doi:10.1080/01621459.2012.737745)
- 36. Reich KJ, Bjorndal KA, Bolten AB. 2007 The 'lost years' of green turtles: using stable
- isotopes to study cryptic lifestages. *Biol. Lett.* **3**, 712–714.
- 569 37. Brewster LR, Dale JJ, Guttridge TL, Gruber SH, Hansell AC, Elliott M, Cowx IG,
- Whitney NM, Gleiss AC. 2018 Development and application of a machine learning
- algorithm for classification of elasmobranch behaviour from accelerometry data. *Mar.*
- *Biol.* **165**, 62. (doi:10.1007/s00227-018-3318-y)
- 573 38. Ladds MA, Thompson AP, Slip DJ, Hocking DP, Harcourt RG. 2016 Seeing it all:
- 574 Evaluating supervised machine learning methods for the classification of diverse
- otariid behaviours. *PLoS One* **11**, 1–17. (doi:10.1371/journal.pone.0166898)
- 576 39. Therneau T, Atkinson B. 2018 rpart: Recursive Partitioning and Regression Trees. R
- 577 Packag. version 4.1-13., https://CRAN.R-project.org/package=rpart.
- 578 40. Liaw A, Wiener M. 2002 Classification and Regression by randomForest. R news 2,
- 579 18–22. (doi:10.1177/154405910408300516)
- 580 41. Chen T et al. 2018 xgboost: Extreme Gradient Boosting. R Packag. version 0.71.2.,
- 581 https://CRAN.R-project.org/package=xgboost.
- 582 42. Meyer D, Dimitriadou E, Hornik K, Weingnessel A, Leisch F. 2017 e1071: Misc
- Functions of the Department of Statistics, Probability Theory Group (Formerly:
- E1071). R Packag. version 1.6-8, https://CRAN.R-project.org/package=e1071.
- 585 43. Venables W, Ripley B. 2002 Modern Applied Statistics with S. Fourth. New York:
- Springer. See http://www.stats.ox.ac.uk/pub/MASS4.
- 587 44. Japkowicz N. 2000 The Class Imbalance Problem: Significance and Strategies. In
- Proceedings of the 2000 International Conference on Artificial Intelligence (IC-

- 589 AI'2000): Special Track on Inductive Learning, Las Vegas, Nevada.
- 590 (doi:10.1.1.35.1693)
- 591 45. Opitz DW, Maclin R. 1999 Popular Ensemble Methods: An Empirical Study. J. Artif.
- 592 *Intell. Res.* **11**, 169–198. (doi:10.1613/jair.614)
- 593 46. Rokach L. 2010 Ensemble-based classifiers. Artif. Intell. Rev. 33, 1-39.
- 594 (doi:10.1007/s10462-009-9124-7)
- 595 47. Hays GC, Marshall GJ, Seminoff JA. 2007 Flipper beat frequency and amplitude
- 596 changes in diving green turtles, Chelonia mydas. Mar. Biol. 150, 1003–1009.
- 597 (doi:10.1007/s00227-006-0412-3)
- 598 48. Okuyama J, Kataoka K, Kobayashi M, Abe O, Yoseda K, Arai N. 2012 The regularity
- of dive performance in sea turtles: A new perspective from precise activity data. *Anim.*
- 600 Behav. **84**, 349–359. (doi:10.1016/j.anbehav.2012.04.033)
- 601 49. Enstipp MR, Ballorain K, Ciccione S, Narazaki T, Sato K, Georges JY. 2016 Energy
- expenditure of adult green turtles (Chelonia mydas) at their foraging grounds and
- during simulated oceanic migration. Funct. Ecol. 30, 1810–1825. (doi:10.1111/1365-
- 604 2435.12667)
- 605 50. Brown DD, Kays R, Wikelski M, Wilson R, Klimley AP. 2013 Observing the
- unwatchable through acceleration logging of animal behavior. Anim. Biotelemetry 1,
- 607 20. (doi:10.1186/2050-3385-1-20)
- 608 51. Koch V, Nichols WJ, Peckham H, De La Toba V. 2006 Estimates of sea turtle
- 609 mortality from poaching and bycatch in Bahía Magdalena, Baja California Sur,
- Mexico. *Biol. Conserv.* **128**, 327–334. (doi:10.1016/j.biocon.2005.09.038)
- 611 52. Wallace BP, Kot CY, Dimatteo AD, Lee T, Crowder LB, Lewison RL. 2013 Impacts
- of fisheries bycatch on marine turtle populations worldwide: Toward conservation and
- 613 research priorities. *Ecosphere* **4**, 1–49. (doi:10.1890/ES12-00388.1)
- 614 53. Ladds MA, Thompson AP, Kadar JP, Slip D, Hocking D, Harcourt R. 2017 Super
- machine learning: Improving accuracy and reducing variance of behaviour
- classification from accelerometry. Anim. Biotelemetry 5, 1–10. (doi:10.1186/s40317-
- 617 017-0123-1)
- 618 54. Ellis K, Godbole S, Marshall S, Lanckriet G, Staudenmayer J, Kerr J. 2014 Identifying
- Active Travel Behaviors in Challenging Environments Using GPS, Accelerometers,
- and Machine Learning Algorithms. Front. public Heal. 2, 36.
- 621 (doi:10.3389/fpubh.2014.00036)
- 622 55. Ali KM, Pazzani MJ. 1996 Error Reduction through Learning Multiple Descriptions.

- 623 *Mach. Learn.* **24**, 173–202.
- 624 56. Resheff YS, Rotics S, Harel R, Spiegel O, Nathan R. 2014 AcceleRater: a web
- application for supervised learning of behavioral modes from acceleration
- measurements. *Mov. Ecol.* **2**, 27. (doi:10.1186/s40462-014-0027-0)
- 57. Shepard ELC et al. 2008 Identification of animal movement patterns using tri-axial
- 628 accelerometry. *Endanger. Species Res.* **10**, 47–60. (doi:10.3354/esr00084)
- 629 58. Yoda K, Sato K, Niizuma Y, Kurita M, Naito Y. 1999 Precise monitoring of
- porpoising behaviour of Adélie penguins determined using acceleration data loggers. J.
- 631 Exp. Biol. 202, 3121–3126.
- 632 59. Moreau M, Siebert S, Buerkert A, Schlecht E. 2009 Use of a tri-axial accelerometer for
- automated recording and classification of goats' grazing behaviour. *Appl. Anim. Behav.*
- 634 *Sci.* **119**, 158–170. (doi:10.1016/j.applanim.2009.04.008)
- 635 60. Jeantet L et al. 2018 Combined use of two supervised learning algorithms to model sea
- 636 turtle behaviours from tri-axial acceleration data. J. Exp. Biol. 221.
- 637 (doi:10.1242/jeb.177378)
- 638 61. Shuert CR, Pomeroy PP, Twiss SD. 2018 Assessing the utility and limitations of
- accelerometers and machine learning approaches in classifying behaviour during
- lactation in a phocid seal. *Anim. Biotelemetry* **6**. (doi:10.1186/s40317-018-0158-y)
- 641 62. Wang Y, Nickel B, Rutishauser M, Bryce CM, Williams TM, Elkaim G, Wilmers CC.
- 642 2015 Movement, resting, and attack behaviors of wild pumas are revealed by tri-axial
- 643 accelerometer measurements. *Mov. Ecol.* **3**, 1–12. (doi:10.1186/s40462-015-0030-0)
- 644 63. Bom RA, Bouten W, Piersma T, Oosterbeek K, van Gils JA. 2014 Optimizing
- acceleration-based ethograms: The use of variable-time versus fixed-time
- segmentation. *Mov. Ecol.* **2**, 6. (doi:10.1186/2051-3933-2-6)
- 647 64. Ladds MA, Salton M, Hocking DP, McIntosh RR, Thompson AP, Slip DJ, Harcourt
- RG. 2018 Using accelerometers to develop time-energy budgets of wild fur seals from
- 649 captive surrogates. *PeerJ* **6**, e5814. (doi:10.7717/peerj.5814)
- 650 65. Bidder OR, Campbell HA, Gómez-Laich A, Urgé P, Walker J, Cai Y, Gao L, Quintana
- F, Wilson RP. 2014 Love thy neighbour: Automatic animal behavioural classification
- of acceleration data using the k-nearest neighbour algorithm. PLoS One 9.
- 653 (doi:10.1371/journal.pone.0088609)
- 654 66. Campbell HA, Gao L, Bidder OR, Hunter J, Franklin CE. 2013 Creating a behavioural
- classification module for acceleration data: using a captive surrogate for difficult to
- observe species. J. Exp. Biol. **216**, 4501–6. (doi:10.1242/jeb.089805)

- 657 67. Wilson RP et al. 2018 Give the machine a hand: A Boolean time-based decision-tree
- template for rapidly finding animal behaviours in multi-sensor data. *Methods Ecol.*
- 659 Evol. **2018**, 1–10. (doi:10.1111/2041-210X.13069)
- 660 68. Lagarde F, Guillon M, Dubroca L, Bonnet X, Ben Kaddour K, Slimani T, El mouden
- EH. 2008 Slowness and acceleration: a new method to quantify the activity budget of
- chelonians. *Anim. Behav.* **75**, 319–329. (doi:10.1016/j.anbehav.2007.01.010)
- 663 69. Martiskainen P, Järvinen M, Skön J-P, Tiirikainen J, Kolehmainen M, Mononen J.
- 2009 Cow behaviour pattern recognition using a three-dimensional accelerometer and
- support vector machines. Appl. Anim. Behav. Sci. 119, 32–38.
- 666 (doi:10.1016/j.applanim.2009.03.005)
- 667 70. Shamoun-Baranes J et al. 2012 From sensor data to animal behaviour: An
- oystercatcher example. *PLoS One* 7, e37997. (doi:10.1371/journal.pone.0037997)
- 669 71. McClune DW, Marks NJ, Wilson RP, Houghton JDR, Montgomery IW, McGowan
- NE, Gormley E, Scantlebury M. 2014 Tri-axial accelerometers quantify behaviour in
- the Eurasian badger (Meles meles): Towards an automated interpretation of field data.
- *Anim. Biotelemetry* **2**, 1–6. (doi:10.1186/2050-3385-2-5)
- 673 72. Lush L, Wilson RP, Holton MD, Hopkins P, Marsden KA, Chadwick DR, King AJ.
- 2018 Classification of sheep urination events using accelerometers to aid improved
- measurements of livestock contributions to nitrous oxide emissions. *Comput. Electron.*
- 676 Agric. **150**, 170–177. (doi:10.1016/j.compag.2018.04.018)
- 73. Robert B, White BJ, Renter DG, Larson RL. 2009 Evaluation of three-dimensional
- accelerometers to monitor and classify behavior patterns in cattle. *Comput. Electron.*
- 679 *Agric.* 67, 80–84. (doi:10.1016/j.compag.2009.03.002)
- 680 74. Allik A, Pilt K, Karai D, Fridolin I, Leier M, Jervan G. 2019 Optimization of physical
- activity recognition for real-timewearable systems: Effect ofwindow length, sampling
- frequency and number of features. *Appl. Sci.* **9**. (doi:10.3390/app9224833)
- 683 75. Watanabe YY, Takahashi A. 2013 Linking animal-borne video to accelerometers
- reveals prey capture variability. Proc. Natl. Acad. Sci. U. S. A. 110, 2199–2204.
- 685 (doi:10.1073/pnas.1216244110)
- 686 76. Jeantet L. et al. 2020, Raw acceleration, gyroscope and depth profiles associated with
- the observed behaviours of free-ranging immature green turtles in Martinique, v2,
- Dryad, Dataset, https://doi.org/10.5061/dryad.hhmgqnkd9

690 **10 Tables**

Table 1: Total duration (seconds) of the observed sequences of behavioural categories for the thirteen free-ranging immature green

693 turtles

Behaviour	#1	#2	#3	#4	#5	#6	#7	#8	#9	#10	#11	#12	#13	TOTAL	Relative Importance (%)
Breathing	36	301	37	20	66	87	89	6	57	27	132	75	293	1226	0.51
Feeding	-	1499	162	540	152	1955	70	-	1030	661	6	28	178	6281	2.64
Gliding	-	896	366	524	211	284	1054	102	1257	129	609	372	2271	8075	3.39
Resting	-	10134	7747	4760	5807	11302	19502	711	7190	602	17579	3814	27441	116590	48.95
Scratching	-	512	574	1789	8	903	136	-	64	21	177	94	218	4496	1.89
Staying at the surface	-	898	1396	1546	1394	2541	2955	573	1032	818	1485	582	3246	18465	7.75
Swimming	5279	6522	3801	4082	6421	6005	4800	2026	6354	5493	7739	2760	18895	80178	33.66
Other	-	258	169	283	148	818	136	45	261	209	233	140	188	2887	1.21

Table 2: Average duration of each behaviour shown by the 13 immature green turtles) predicted time vs. observed time. The percentages are expressed with respect to the total individual recorded video duration or to the time the behaviour in question was expressed. The predicted durations of the diving behaviours were obtained using the WS method, and the surfacing behaviours were predicted using depth values.

700	

Behaviour						
Benaviour	predicted (s)	observed (s)	difference (s)	%_total	%_behaviour	
Breathing	99	94	32*	0.2*	46.9	
Feeding	432	497	207	1,1	180.1	
Gliding	936	651	326	1.0	39.1	
Other	92	235	143	0.8	60.2	
Resting	9175	9640	747**	2.7**	6.6	
Scratching	437	354	118	0.7	311.8**	
Staying at the surface	1435	1477	151	0.8	10.8	
Swimming	6206	6256	441	2.4	6.9*	
Transition	48	-	48	0.2*	-	

* The lowest difference obtained among the nine behavioural categories

** The highest difference obtained among the nine behavioural categories

11 Figures

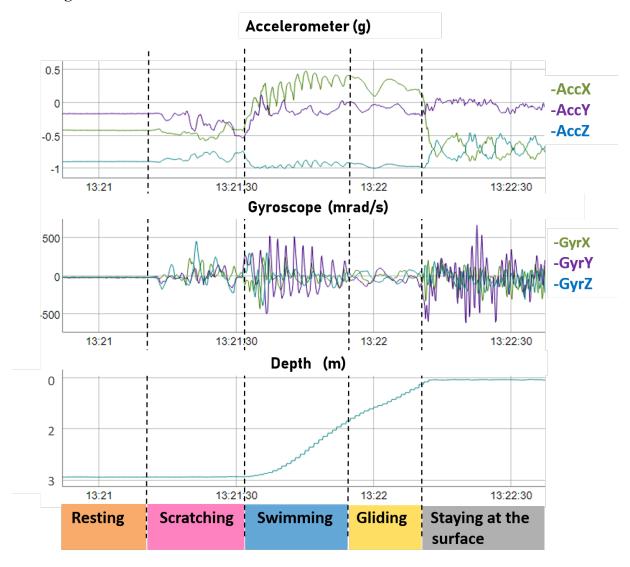


Figure 1. Raw acceleration, gyroscope and depth profiles for several behaviours expressed by turtle #12.

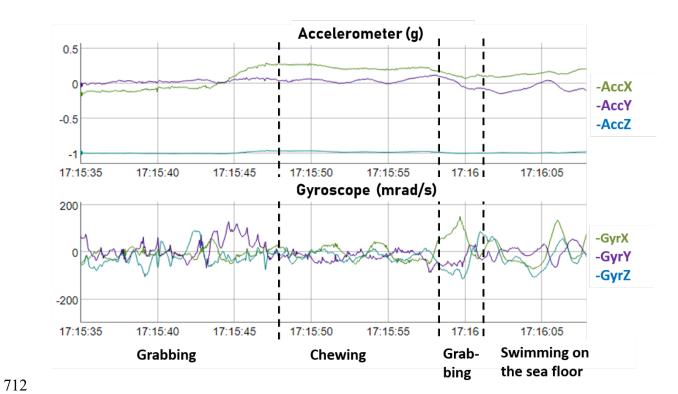


Figure 2: Raw acceleration and gyroscope signals obtained for the feeding behaviours expressed by turtle #6. The definitions of the behaviours are available in Supplementary Material (Table S2).

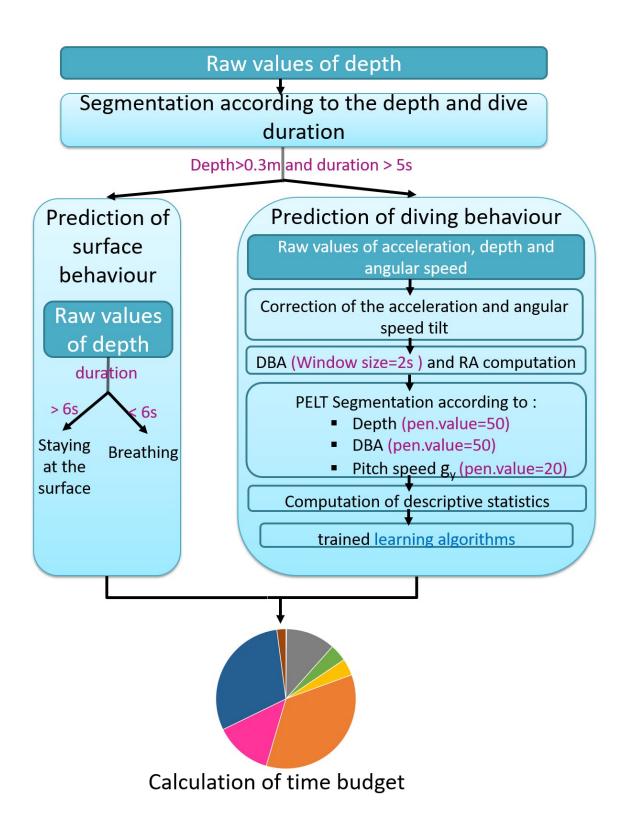


Figure 3. Workflow of automatic behavioural identification using acceleration, angular speed and depth data, as adapted to the green turtle. The hyper-parameters set up specifically for green turtle data are highlighted in purple. The application of this workflow

for other marine species would necessitate the identification of the optimal hyper-parameter values for each species.



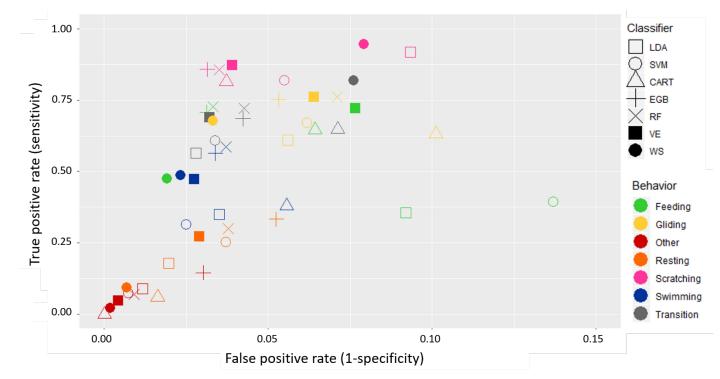


Figure 4. True positive rate vs. the false positive rate obtained with the seven classifiers for the seven diving categories. The symbols show the mean values obtained from 371 combinations of splitting the sample of thirteen individuals into two sub-groups (one of nine individuals for learning and one of four individuals for testing).

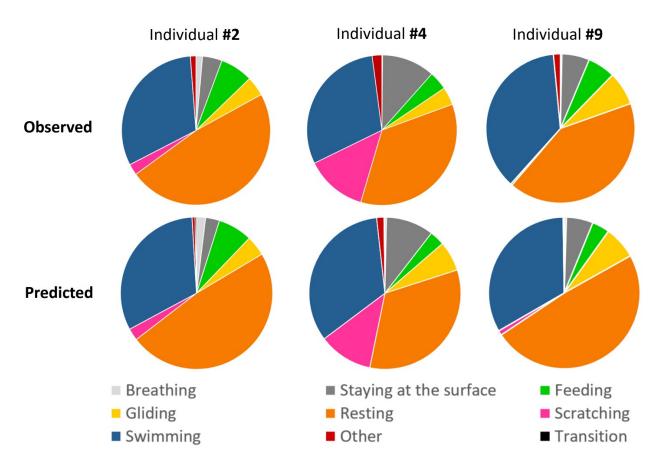


Figure 5. Pie chart of the observed (determined from the video) vs. predicted mean durations of the various behaviours displayed by three free-ranging immature green turtles. The predicted durations of the diving behaviours were obtained using the WS classifier.

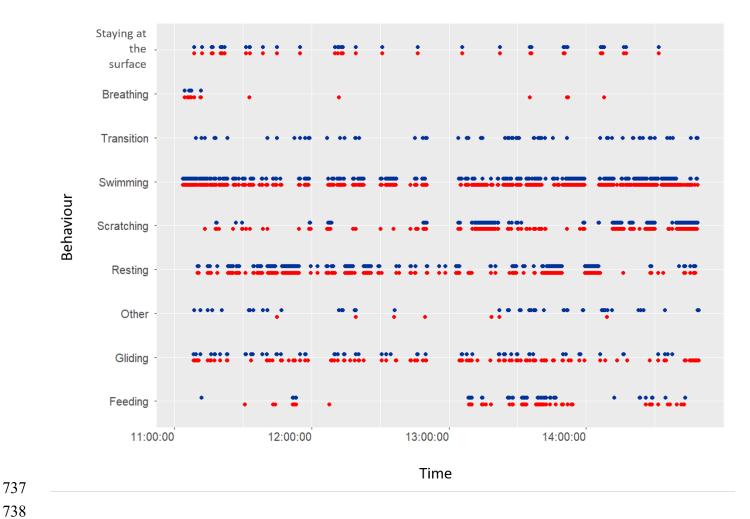


Figure 6. Comparison of the nine main inferred behavioural categories (in red) and of the actually observed ones (in blue) for a few hours for immature green turtle #1. The predicted occurrences of the diving behaviours were obtained using the WS classifier.