



## Canadian Journal of Fisheries and Aquatic Sciences

### Assessing the performance of statistical classifiers to discriminate fish stocks using Fourier analysis of otolith shape

Journal:	<i>Canadian Journal of Fisheries and Aquatic Sciences</i>
Manuscript ID	cjfas-2019-0251.R1
Manuscript Type:	Article
Date Submitted by the Author:	14-Oct-2019
Complete List of Authors:	Smoliński, Szymon; Havforskningsinstituttet; Morski Instytut Rybacki-Panstwowy Instytut Badawczy, Department of Fisheries Resources Schade, Franziska; Johann Heinrich von Thunen-Institut Institut für Ostseefischerei Berg, Florian; Havforskningsinstituttet; University of Bergen, Department of Biological Sciences
Keyword:	Atlantic cod <i>Gadus morhua</i> , Atlantic herring <i>Clupea harengus</i> , fish stock discrimination, machine learning, support vector machines
Is the invited manuscript for consideration in a Special Issue? :	Not applicable (regular submission)

SCHOLARONE™  
Manuscripts

1 **Assessing the performance of statistical classifiers to discriminate fish stocks using**  
2 **Fourier analysis of otolith shape**

3

4 Szymon Smoliński<sup>1,2,\*</sup>, Franziska Maria Schade<sup>3</sup>, Florian Berg<sup>1,4</sup>

5

6 <sup>1</sup> Institute of Marine Research, P.O. Box 1870 Nordnes, 5817 Bergen, Norway

7 <sup>2</sup> Department of Fisheries Resources, National Marine Fisheries Research Institute, Kołłątaja  
8 1, 81-332 Gdynia, Poland

9 <sup>3</sup> Thuenen Institute of Baltic Sea Fisheries, Alter Hafen Süd 2, 18069 Rostock, Germany

10 <sup>4</sup> University of Bergen, Department of Biological Sciences, P.O. Box 7803, 5020  
11 Bergen, Norway

12 \* E-mail: [szymon.smolinski@hi.no](mailto:szymon.smolinski@hi.no)

Draft

**13 Abstract**

14 The assignment of individual fish to its stock of origin is important for reliable stock  
15 assessment and fisheries management. Otolith shape is commonly used as the marker of  
16 distinct stocks in discrimination studies. Our literature review showed that the application and  
17 comparison of alternative statistical classifiers to discriminate fish stocks based on otolith  
18 shape is limited. Therefore, we compared the performance of two traditional and four machine  
19 learning classifiers based on Fourier analysis of otolith shape using selected stocks of Atlantic  
20 cod (*Gadus morhua*) in the southern Baltic and Atlantic herring (*Clupea harengus*) in the  
21 western Norwegian Sea, Skagerrak and the southern Baltic Sea. Our results showed that the  
22 stocks can be successfully discriminated based on their otolith shapes. We observed  
23 significant differences in the accuracy obtained by the tested classifiers. For both species,  
24 support vector machines (SVM) resulted in the highest classification accuracy. These findings  
25 suggest that modern machine learning algorithms, like SVM, can help to improve the  
26 accuracy of fish stock discrimination systems based on the otolith shape.

27

28 **Key words:** Atlantic cod *Gadus morhua*, Atlantic herring *Clupea harengus*, fish stock  
29 discrimination, machine learning, support vector machines

## 30 **1. Introduction**

31 Discrimination of fish stocks is essential for reliable fisheries resource management and is  
32 currently an integral part of modern fish stock assessments (Begg et al. 1999). Many  
33 commercially exploited fish stocks show strong habitat overlaps, resulting in a temporal  
34 mixing. A disregard of stock mixing, particularly when stocks differ in productivity, may lead  
35 to the overexploitation of unique spawning components (Kell et al. 2004; Kerr et al. 2017).  
36 Therefore, individuals from mixed-stock catches need to be assigned to their stock of origin  
37 using reliable stock discrimination methods with high classification accuracy (Cadrin et al.  
38 2014).

39 One widely applied stock discrimination technique involves otoliths; calcium carbonate  
40 structures located in the inner ear of fishes (Campana and Casselman 1993). Otolith shape is  
41 mostly driven by a combination of environmental and genetic factors and contains stock-  
42 specific features, which are usable as a relevant marker of distinct stocks (Vieira et al. 2014;  
43 Berg et al. 2018). In recent years, diverse methods enabling the description of the otolith  
44 shape were developed and tested, such as curvature-based descriptors, wavelets, shape  
45 geodesics or mirroring techniques (Parisi-Baradad et al. 2005; Nasreddine et al. 2009; Harbitz  
46 and Albert 2015). However, otolith outlines are still most frequently investigated with a  
47 mathematical scheme of Fourier decomposition, namely fast Fourier transform or elliptical  
48 Fourier analysis (Stransky 2014). Both fast Fourier transform and elliptical Fourier techniques  
49 decompose shape, which is a polygon of two-dimensional coordinates, into a spectrum of  
50 harmonically related trigonometric curves and calculate coefficients describing each of these  
51 curves (for details see Haines and Crampton 2000; Kuhl and Giardina 1982). Calculated  
52 coefficients may be then used as predictors for the discrimination of fish stocks in  
53 multivariate statistical analysis (Stransky 2014).

54 However, once shape coefficients are extracted, little attention has been paid to apply and  
55 compare performances of alternative statistical systems to assign fish individuals to known  
56 groups (stocks or species) based on their otolith shape. Available classifiers arise from  
57 different fields, like statistics (e.g., linear discriminant analysis), artificial intelligence and  
58 data mining (e.g., decision-trees) or connectionist approaches (e.g., neural networks)  
59 (Fernández-Delgado et al. 2014). Most machine learning (ML) algorithms are not yet part of  
60 the traditional statistical modeling, hence their application in ecology is still scarce (Olden et  
61 al. 2008). However, modern ML algorithms have a high potential to outperform traditional  
62 parametric classifiers in solving real-world classification problems (Fernández-Delgado et al.  
63 2014). They are much more flexible than conventional models and are able to handle the non-  
64 linear relationships and interacting elements that often characterize biological data (Guisan  
65 and Zimmermann 2000). Current computational capabilities and freely available statistical  
66 software allow relatively easy implementation of these modern algorithms and they may be  
67 valuable in the development of fish stock discrimination routines. The advantages of ML  
68 applications have been already considered in other stock discrimination approaches, like in  
69 otolith chemistry (e.g., Mercier et al. 2011) or analysis of parasitological markers (e.g.,  
70 Perdiguero-Alonso et al. 2008). These studies strongly suggest that current ML classifiers are  
71 already well suited to assign fish to stocks and that classification abilities are improved  
72 compared to traditional discriminant analysis.

73 Few studies used ML algorithms and Fourier analysis of otolith shape to discriminate fish  
74 stocks (e.g., Zhang et al. 2016; Mapp et al. 2017). However, these studies did not compare the  
75 ML performance with traditional classifiers like linear discriminant analysis. Only recently  
76 Jones and Checkley (2017) compared random forest with discriminant analysis to identify  
77 otoliths found in sediment cores and showed that the ML approach outperformed the  
78 traditional classifier. However, they applied these algorithms to distinguish between species,

79 i.e. between higher taxonomic groups that naturally show stronger otolith shape differences  
80 than between fish stocks. To the best of our knowledge, no comprehensive comparison of  
81 traditional and modern ML classifiers to assign individuals to fish stocks has been conducted.  
82 Here, we apply six statistical classifiers (two traditional: linear discriminant analysis (LDA),  
83 quadratic discriminant analysis (QDA), and four machine learning classifiers: K-nearest  
84 neighbors (KNN), classification and regression trees (CART), random forest (RF) and support  
85 vector machines (SVM)) to discriminate stocks of two commercially exploited fish species,  
86 where Fourier analysis of otolith shape is required for accurate estimation of mixing ratios for  
87 a proper stock assessment: Atlantic cod (*Gadus morhua*) in the southern Baltic and Atlantic  
88 herring (*Clupea harengus*) in the northeastern Atlantic.

89 This paper aims to i) conduct a systematic review of the available scientific literature focusing  
90 on statistical classifiers associated with Fourier analysis of otolith shape for discrimination  
91 purposes; ii) investigate the otolith shape variability of cod and herring stocks by applying  
92 elliptical Fourier analysis; and iii) assess the performance of traditional and recent ML  
93 classifiers to assign fish individuals to their group of origin based on their otolith shape.

94

## 95 **2. Materials and methods**

### 96 2.1. Literature review of the use of statistical classifiers

97 Peer-reviewed literature was searched in the Web of Science Core Collection database using  
98 the keywords: “otolith\$” and “Fourier”. Only English-language studies on otolith shape that  
99 applied Fourier analysis to discriminate fish groups at different biosystematics levels  
100 (ecotype, stock, population, species) were chosen for further investigation. Selected literature  
101 was reviewed to analyze which statistical classification algorithm was applied to discriminate  
102 different fish groups. Different types of algorithms based on the framework of Fisher  
103 discriminant analysis (Fisher 1936), including parametric and nonparametric extensions, were

104 aggregated as one group ("discriminant analysis"). The list of 106 publications used in the  
105 review process is given in the supplementary materials (Table S1).

106

## 107 2.2. Study species and datasets

### 108 2.2.1. Atlantic cod *Gadus morhua*

109 Atlantic cod is one of the most important commercially exploited fish species across the  
110 North Atlantic Ocean, inhabiting also the brackish waters of the Baltic Sea. Here, Baltic cod  
111 is managed as two separate stocks: one western stock (ICES subdivisions (SDs) 22-24) and  
112 one eastern stock (SDs 24-32, ICES 2019a). The genetically distinct cod stocks coexist in the  
113 Arkona Basin (SD 24, Hemmer-Hansen et al. 2018, Weist et al. 2019), resulting in  
114 uncertainties in the stock assessment. Since the ICES benchmark in 2015, otoliths of cod from  
115 commercial samples from the mixing area are assigned to their respective stock of origin  
116 using Elliptic Fourier descriptors and LDA (ICES 2015, 2019b; Hüseyin et al. 2016). For this  
117 study, we used otolith images of genetically validated Baltic cod samples (N=507, Weist et al.  
118 2019) from the mixing area (SD24, Fig.1) and from adjacent areas (Belt Sea (SD 22),  
119 Øresund (SD 23) and Bornholm Basin (SD 25)). The dataset consists of 52% western Baltic  
120 cod (WBC) and 48% eastern Baltic cod (EBC) (Table 1). For further details refer to Schade et  
121 al. (2019).

122

### 123 2.2.2. Atlantic herring *Clupea harengus*

124 Atlantic herring is a commercially exploited fish species in the northeastern Atlantic that has  
125 been a key species for stock discrimination studies (Geffen 2009). Herring stocks in this  
126 region consist of multiple spawning components. In this study, we analyzed otoliths from four  
127 distinct spawning components (Table 1): Norwegian spring spawners (NSS, 27% of herring  
128 data), coastal Skagerrak spring spawners (CSS, 20%), Greifswald Bay herring (GB, 31%) and

129 central Baltic northern component (CBNC, 22%) (ICES 2018a, 2018b). While NSS is clearly  
130 a separate stock, CSS and GB are managed within the stock of western Baltic spring spawners  
131 (WBSS), whereas CBNC is part of the central Baltic herring (CBH) stock. To ensure that  
132 distinct components were sampled, we only used herring sampled in spawning condition.  
133 Further, only herring of age 5-6 were selected to reduce age effects on shape variability  
134 (Libungan et al. 2015). Herring were mainly collected during scientific surveys, except for  
135 GB and some samples of CSS that were caught by local fishers using gillnets.

136

### 137 2.3. Otolith shape analysis

138 For cod and herring, shape images of clean and unbroken sagittal otoliths were used. Images  
139 of the right otolith were preferred; otherwise, the image of the left otolith was flipped. There  
140 are no differences between left and right otoliths for cod (Campana and Casselman 1993;  
141 Cardinale et al. 2004) and herring (Libungan et al. 2015). High-resolution images were  
142 binarized using the threshold function of the GNU Image Manipulation program (Natterer and  
143 Neumann 2008).

144 For the shape analysis, outlines were automatically obtained from converted images using the  
145 *Momocs* package (Bonhomme et al. 2014) in the R environment (R Core Team 2018).

146 Elliptical Fourier analysis proposed by Kuhl and Giardina (1982) was used to quantify otolith  
147 outlines. This technique decomposes two-dimensional shape with a sum of harmonics, where  
148 each harmonic is described by four coefficients (two for  $x$ -axis and two for  $y$ -axis  
149 coordinates). Precision of approximate reconstruction of shapes increases with the number of  
150 harmonics used, but it is recommended to reduce the number of harmonics for multivariate  
151 analysis. To define the appropriate number of harmonics, 100 otoliths were randomly sampled  
152 from the whole set and the Fourier power ( $PF_n$ ) spectrum and cumulated Fourier power ( $PF_c$ )  
153 was calculated with the following formulas:



$$PF_n = \frac{A_n^2 + B_n^2 + C_n^2 + D_n^2}{2}$$

$$PF_c = \sum_1^n PF_n$$

156 where  $A_n$ ,  $B_n$ ,  $C_n$ ,  $D_n$  are the coefficients of  $n^{\text{th}}$  harmonic (Lord et al. 2012). The number of  
157 harmonics that reaches 99% of cumulated Fourier power of 30 harmonics were chosen to  
158 summarize shapes of otoliths (Stransky et al. 2008b; Vieira et al. 2014). The first three  
159 coefficients were taken as fixed values ( $A_1=1$ ;  $B_1=C_1=0$ ) to normalize otoliths for size,  
160 orientation and starting point (Tracey et al. 2006). Mean otolith shapes of different stock  
161 components were calculated by invert transformation of Fourier coefficients. Overall variance  
162 in the shape of otoliths was assessed with principal component analysis (PCA) integrated with  
163 morphospaces (theoretical shapes were reconstructed based on the PCA scores) (Bonhomme  
164 et al. 2014).

165

#### 166 2.4. Statistical classifiers

167 Among the six selected algorithms, linear discriminant analysis (LDA) and quadratic  
168 discriminant analysis (QDA) were chosen as one of the most popular classifiers, widely  
169 implemented in otolith-based fish stock and species discrimination (e.g., Paul et al. 2013;  
170 Zhang et al. 2013). They are applied to predict the affiliation of observations from two or  
171 more known classes. Both classifiers use the best combination of several characters that  
172 provide the strongest separation of classes by maximizing the standard deviation between  
173 obtained groups and minimizing them within groups (Fisher 1936).

174 K-nearest neighbors (KNN) algorithm is one of the simplest ML classifier that can be applied  
175 both to binary and multiclass problems (Hall et al. 2008). In the first step, it selects the nearest  
176 neighbors and then determines the class of observation using these selected neighbors. One of  
177 the KNN advantages is its higher tolerance of the data structure (Hastie et al. 2009).

178 Similarly, classification and regression trees (CART), a nonparametric procedure, requires no  
179 assumptions about the distribution of the data. These models are obtained by recursively  
180 partitioning the data space and fitting a simple prediction model within each partition. As a  
181 result, the partitioning can be represented graphically as a decision tree (Loh 2011).

182 Random forest (RF) is an ensemble technique, based on a set of CARTs, where a bootstrap  
183 approach is implemented to select a random set of observations and variables used to  
184 construct each tree in ensemble. Finally, decisions of all trees on object allocation are  
185 aggregated and the majority is used in order to provide final class prediction (Breiman 2001).

186 Support vector machines (SVM) was selected among the broad range of ML approaches,  
187 because of its ability to deal with high-dimensional datasets and its flexibility in modeling  
188 diverse data sources (Ben-Hur et al. 2008). This technique uses kernel functions to project the  
189 predictive variables into feature space with more dimensions than the initial space of the input  
190 data, allowing the construction of linear models (Cortes and Vapnik 1995).

191

## 192 2.5. Statistical analysis

193 All predictors (Fourier coefficients) were examined for normality with graphical tools (Zuur  
194 et al. 2010). None of the variables showed significant deviation from normal distribution. For  
195 each fish species, differences in total fish length between stock components were tested and  
196 found to be significant using one-way ANOVA (TukeyHSD,  $p < 0.001$ ). To test allometric  
197 effects of fish length on shape coefficients, analyses of covariance (ANCOVAs) were  
198 conducted. Information on stock components origin was included in the model as fixed  
199 factors and fish length as covariate. If the interaction between fixed factor and covariate was  
200 significant, the variable was excluded from the dataset, otherwise, shape coefficients with  
201 significant fish length effect were standardized using the common slope for all stock  
202 components (Zhuang et al. 2014).

203 Classification and Regression Training package *caret* (Kuhn 2008) for R was used to compare  
204 performances of selected classifiers. The package allows for different algorithms to be trained  
205 in a consistent environment and to conduct the tuning of the machine learning parameters. All  
206 predictor variables were scaled and centered in a preprocessing stage. Optimal  
207 hyperparameters of KNN ( $k$ ), CART ( $cp$ ), RF ( $mtry$ ) and SVM ( $\sigma$  and  $C$ ) were defined during  
208 preliminary tuning (Fig. S1 and S2). Following Mercier et al. (2011) and Zhang et al. (2016),  
209 a 4-fold cross-validation resampling method was used to provide the data for the assessment  
210 of the performance of each classifier. This validation method is advised as a reasonably stable  
211 and low biased measure of model performance (Hastie et al. 2009), but typically indicates  
212 lower accuracy of the evaluated algorithms than most often applied leave-one-out cross-  
213 validation. Datasets were randomly split into four equal subsets with preservation of class  
214 ratios, where three subsets (75% of observations) were used as training data to classify the  
215 remaining subset (25% of observations). Validation was repeated for each of the four splits.  
216 Additionally, 100 repetitions of the whole process were conducted using a bootstrap approach  
217 with independent resampling (Hastie et al. 2009). Confusion (error) matrices (e.g. Kuhn 2008;  
218 Perdiguero-Alonso et al. 2008) were generated and classification accuracy (the percentage of  
219 fish correctly assigned to their actual class) was calculated as a measure of classifier quality.  
220 In order to assess the influence of the number of Fourier harmonics used for the shape  
221 representation on classification accuracy, each cross-validation procedure (400 repetitions)  
222 was conducted on datasets produced with between 2 to  $n$  harmonics, where  $n$  is the number of  
223 harmonics that reach 99% of cumulated Fourier power. When number of variables was lower  
224 than the specified optimal hyperparameter  $mtry$  for RF, the default  $mtry$  was applied, which  
225 equals the square root of the number of variables. Moreover, in order to assess the influence  
226 of the number of classes on the performance of classifiers, herring dataset was split into two-  
227 class subsets and similar cross-validation was run for each pair of spawning components. The

228 algorithms were developed in parallel, using the same training and test sets. Therefore, paired  
229 t-tests with adjusted p-values to control false discovery rates (Benjamini and Hochberg 1995)  
230 were used to test differences in accuracies of classifiers in relation to the dataset with the  $n$   
231 number of Fourier harmonics. The importance of Fourier descriptors was calculated with the  
232 *varImp* function of the *caret* package and was visualized in decreasing order using mean  
233 importance for all models. All of the models were built using following the R packages: LDA  
234 and QDA with *MASS* (Brian et al. 2015), KNN with *caret* (Kuhn 2008), CART with *rpart*, RF  
235 with *randomForest* (Liaw and Wiener 2002) and SVM based on the radial basis function  
236 (RBF) kernel with *kermlab* (Karatzoglou et al. 2015).

237

### 238 3. Results

#### 239 3.1. Literature review of the use of statistical classifiers

240 Among 106 selected papers published in the period from 1990 to 2018 that incorporate  
241 Fourier analysis as the method for otolith shape description, the framework of Fisher  
242 discriminant analysis (DA) was the most popular statistical approach. Studies that applied  
243 only DA constituted ~92%, while one study (<1%) used DA and RF in parallel (Jones and  
244 Checkley 2017). The remaining ~7% of the publications applied classifiers other than DA to  
245 assign samples to their respective class, e.g., support vector machines or K-nearest neighbors  
246 classifier (Reig-Bolaño et al. 2010b; Benzinou et al. 2013), boundary-based shape  
247 classification (Nasreddine et al. 2009), between-class correspondence analysis (Ponton 2006),  
248 or random forest (e.g., Zhang et al. 2016).

249 Application of more than one classifier in the same analysis was scarce (~8% of papers).

250 Comprehensive comparison of accuracy of nine ML algorithms was done by Mapp et al.  
251 (2017), including naive Bayes, Bayesian networks, logistic regression, HyperPipes, C4.5, RF,  
252 KNN, SVM, and rotation forest. Jones and Checkley (2017) showed that RF algorithms

253 outperformed DA in terms of accuracy. Torres et al. (2000) presented that QDA was superior  
254 to LDA, while Finn et al. (1997) found no differences between LDA and QDA models. SVM  
255 performed better than KNN in terms of correct classification rate, but the second classifier  
256 resulted in more stable performances across the classes and has been chosen for  
257 discrimination of fish based on otolith shape in Benzinou et al. (2013).

### 258 3.2. Otolith shape variability

259 Precision of approximate reconstruction of shapes increased with the number of harmonics  
260 used (Fig. 2). For both species, 13 harmonics were needed to achieve 99% of cumulative  
261 Fourier power summarizing the otolith shapes. Consequently, the first 13 harmonics were  
262 used in further analyses. Due to the significant interaction between stock components and fish  
263 total length in the ANCOVA models ( $p < 0.001$ ), six and 12 Fourier descriptors were excluded  
264 from cod and herring data, respectively. A further 23 (cod) and 29 (herring) descriptors were  
265 corrected for the fish length effect using a common slope.

266 Visual inspection of mean otolith shape identified differences between cod stocks and herring  
267 components (Fig. S3). Among cod stocks, WBC had wider otoliths than EBC. Otoliths of  
268 NSS and CBNC herring were generally wider than those of CSS and GB herring, which mean  
269 otolith shapes were very similar.

270 For cod, the first two PCA axes explained 72.6% of the overall variance in the shape of  
271 otoliths (Fig. 3a). The two cod stocks were mainly separated along the first axis, even though  
272 a strong overlap was observed. For herring, 66.3% of the overall variance was explained by  
273 the first two axes (Fig. 3b).

274

### 275 3.3. Classification accuracy

276 The classification accuracy of cod otoliths increased with increasing number of harmonics but  
277 stayed relatively constant for six and more harmonics (Fig. 4a). One exception is QDA, where

278 the accuracy slightly decreased with a higher number of harmonics. In comparison, the  
279 accuracy continued to increase for herring otoliths with increasing number of harmonics (Fig.  
280 4b).

281 The accuracy differed significantly between classifiers, except for QDA and KNN for cod  
282 otoliths as well as LDA and KNN for herring (Table 2). For both species, SVM resulted in the  
283 highest classification accuracy (Fig. 4), even when herring data were sequentially split into  
284 two-class subsets (Fig. S4). LDA resulted in slightly but significantly lower accuracy for cod  
285 (Fig. 4a, Table 2).

286 The 4-fold cross-validation using SVM (best classifier) and 13 harmonics (accounting for  
287 99% variance of the otolith shape) resulted in an accuracy of 79.54% for cod and 74.13% for  
288 herring (Table 3). For cod, the misclassification rate was equal in both stocks (~10%). For  
289 herring, the highest misclassification occurred between GB and CSS herring (~7%).

290 Misclassification among the other herring components was low (<1%).

291 The relative importance of individual Fourier descriptors was consistent among statistical  
292 classifiers for both species (Fig. 5), except for CART. CART and RF both rely on the  
293 importance of only a few descriptors (~8 or less), while the other classifiers rely on the  
294 importance of a higher number of Fourier descriptors.

295

#### 296 **4. Discussion**

297 Presented review of the literature showed that the application and comparison of alternative  
298 classifiers to discriminate fish groups based on their otolith shape is limited. In this study,  
299 stock-specific differences in otolith shapes for cod and herring could be detected, which  
300 enables the assignment of individual fish to its respective stock of origin. Moreover, a  
301 comparison of different statistical classifiers suggested that ML algorithms, in particular

302 SVM, can improve the accuracy in stock discrimination approaches using the shape of  
303 otoliths.

304

#### 305 4.1 Literature review of the use of statistical classifiers

306 The literature review emphasized that traditional DA was used in most of the studies for the  
307 classification of fish groups based on the elliptical Fourier descriptors of otolith shape, while  
308 application of alternative classifiers was less common. For example, Zhang et al. (2016) used  
309 random forest to discriminate stocks of the Japanese Spanish mackerel (*Scomberomorus*  
310 *niphonius*) based on Fourier descriptors of otolith shapes, but no comparison with other  
311 classifiers was reported. Mapp et al. (2017) used nine ML algorithms for fish stock separation  
312 of two clupeid species using otolith shapes. However, the study of Mapp et al. (2017) was not  
313 focused on the absolute classification accuracy, but on the applicability of morphometric  
314 approaches that incorporate size information. No comparison with traditional classifiers, like  
315 linear discriminant analysis, was made in Mapp et al. (2017), while Jones and Checkley  
316 (2017) showed that RF algorithms were superior to DA during classification of fish  
317 individuals into different taxonomic groups based on the morphological descriptors and  
318 elemental compositions of otoliths.

319 Studies comparing more than one statistical classification algorithm indicated that the success  
320 of fish classification can be significantly improved by alternative classifiers (Torres et al.  
321 2000). These findings stress the need for the comparison of different classifiers, i.e., different  
322 approaches should be explored so that the best method is used in order to achieve the best  
323 possible assignment. More accurate assignment of individual fish allows for more robust  
324 estimation of the contribution of different fish stocks within the mixing areas (i.e., a mixed  
325 stock scenario, Hüseyin et al. 2016). Accurate estimates of mixing levels can help to understand

326 how movement and mixing affect stock dynamics and provide the quantitative basis for  
327 annual stock assessments and scientific advice (Horbowy 2005; Taylor et al. 2011).

328

#### 329 4.2 Otolith shape variability

330 Our results support the previous studies showing that Baltic cod stocks can be successfully  
331 discriminated based on the elliptical Fourier analysis of otolith outlines (Paul et al. 2013;  
332 Hüsey et al. 2016). Significant differences in otolith shape were also reported for other stocks  
333 and spawning populations of cod, e.g., the northeast Arctic and Norwegian coastal cod  
334 (Stransky et al. 2008a), Faroe Plateau cod (Cardinale et al. 2004) or Icelandic cod  
335 (Petursdottir et al. 2006). Mean shapes reconstructed on the calculated Fourier descriptors  
336 indicated that the otolith outline of WBC and EBC differ in the large-scale shape  
337 characteristics (mainly length–width relationship), where otoliths from the western stock are  
338 wider and rounder than those from the eastern stock, which is in line with previous  
339 observations (Paul et al. 2013; Hüsey et al. 2016). Differences in circularity and rectangularity  
340 of otoliths were also reported in other cod stocks (Campana and Casselman 1993; Cardinale et  
341 al. 2004).

342 Similarly, discrimination methods based on the analysis of otolith outlines were applied to  
343 separate populations of herring in the Northern Atlantic (e.g., Burke et al. 2008; Libungan et  
344 al. 2015). Our study revealed differences in otolith shape between herring components. Most  
345 of the differences were based on the relationships between the length and width of the whole  
346 otolith. NSS and CBNC have wider otoliths, but the rostrum of NSS herring otoliths is clearly  
347 longer. Confusion matrices of the cross-validated models (Table 3) indicated that a relatively  
348 large number of individuals from the CSS and GB were mis-assigned, suggesting similarity in  
349 otolith shape. This result supports the current assessment approach, where both spawning  
350 components are considered as one stock (WBSS) because of the high level of overlap (ICES



2018b). Although selected herring spawning components were discriminated with a high level of accuracy, further studies need to include other stock components in this region, such as the autumn spawners and the southern component of CBH (ICES 2018a).

The differences in the shape of fish otoliths, for both fish species, may be associated with a combination of environmental and genetic drivers (Cardinale et al. 2004; Vignon and Morat 2010). To explore how these factors influence otolith shape, further analyses are needed, including experimental and laboratory studies with appropriate control of the potentially confounding variables (e.g., Berg et al. 2018). However, even without the mechanistic understanding of the sources of shape variability, these results support the applicability of Fourier analysis of otolith shape in stock discrimination routines and assessment of fish stocks (Cadrin et al. 2014). The use of otoliths as indicator of stock identity has been previously advocated because otoliths are routinely collected for aging in traditional fish monitoring, providing a robust and cost-effective method for stock discrimination (Campana and Casselman 1993; Cardinale et al. 2004).

#### 4.3 Assessment of statistical classifiers

There were significant differences in accuracy between the six statistical classifiers tested. The highest accuracy of fish classification was achieved by SVM, one of the rapidly developing ML classifiers. Accuracy of the SVM model trained on cod data was only 0.9% higher than of the second best performing classifier (LDA), but differences were significant. However, the accuracy of the SVM trained on herring data was 7% to 20% higher than the other classifiers. Good performance of the SVM algorithm, as well as other ML algorithms, has been previously shown in discrimination studies of stocks, species or higher taxonomic levels of fishes based on their otolith shapes (Reig-Bolaño et al. 2010a; Benzinou et al. 2013; Zhang et al. 2016; Mapp et al. 2017).

376 These findings suggest that ML algorithms are a good alternative to traditional classifiers and  
377 can help to improve the accuracy of routine fish stock discrimination using the shape of the  
378 otolith. Although SVM achieved the highest accuracy in this study, we strongly advise to test  
379 a range of statistical classifiers in discrimination studies, because the selection of the best  
380 performing algorithm can be case-specific, and depends e.g., on the number of classes,  
381 similarity between groups, or type and number of variables in the dataset (Fernández-Delgado  
382 et al. 2014).

383 Caution is however warranted. The proposed benchmark of different statistical classifiers  
384 should be conducted only in systems with well-defined units. The ability of ML classifiers to  
385 find structures and clusters in the data needs to be considered with caution. Application of the  
386 ML algorithms for the discrimination of fish groups, where training baselines are not  
387 validated (e.g., by genetics or by sampling spawning individuals in their respective spawning  
388 area), may potentially lead to confusing results and recognition of subgroups, which may not  
389 represent the real biological or management units. The practical problems of managing natural  
390 resources with poorly defined units continue to be an important issue (Geffen 2009). For these  
391 reasons, the definition of robust baselines for the training of classification algorithms is a  
392 crucial point in the development of operational discrimination systems (Cadrin et al. 2014;  
393 Hüsey et al. 2016; Schade et al. 2019).

394

#### 395 4.4 Study limitations and future implications

396 In this study, a simple approach was applied, using only Fourier descriptors of otolith shapes  
397 as predictors of fish stock affiliation. The focus was exclusively on the differences of  
398 statistical classifier accuracies on the length-normalized descriptors of otolith shape (Hüsey et  
399 al. 2016). However, incorporation of other potentially informative variables, such as shape  
400 indices or routinely collected information on length-at-age, and sex of individual fish can

401 further improve the predictive abilities of classification algorithms (Burke et al. 2008; Mapp  
402 et al. 2017). Further, alternatives to reconstruct the otolith shape like wavelet transformation  
403 or curvature scale space representation should be reconsidered. Fourier descriptors focusing  
404 on periodic phenomena (Harbitz and Albert 2015) might be more suited for cod otoliths that  
405 are almost elliptical. For more complex otolith shapes with very localized landmarks, like  
406 herring otoliths, wavelet transformation could be better-suited (Sadighzadeh et al. 2014).

407 Besides otolith shapes, ML algorithms were already used successfully in other stock  
408 discrimination fields, e.g., population genetics (Guinand et al. 2002), otolith microchemistry  
409 (Mercier et al. 2011), hydroacoustics (Robotham et al. 2010) or parasitology (Perdiguero-  
410 Alonso et al. 2008), even though the application is still rare.

411 In our study, the analysis of Fourier power spectrum indicated that 13 harmonics were needed  
412 to explain 99% of the variance in the otolith shape both for cod and herring. Interestingly,  
413 high accuracy for the cod assignment was already obtained with only 5 to 6 harmonics,  
414 suggesting that additional higher-frequency harmonics do not incorporate much information  
415 for the discrimination of these stocks. These results are in line with the analysis of variable  
416 importance which showed that lower-rank descriptors (D5, D1 - describing a global form of  
417 otoliths) were the most powerful predictors in all models. The broadly applied practice to  
418 include only a certain subset of harmonics (e.g., first  $N$  harmonics needed to describe 99% of  
419 shape variance) may not be optimal in the context of classification model performance. For  
420 fish species with simple otolith shapes, a reduced number of Fourier harmonics may be  
421 advantageous. Conversely, the inclusion of a larger number of harmonics in classification  
422 systems developed for species with more complex otolith structures, like herring, can help to  
423 achieve a better quality of classification models. In our study, a steady improvement of model  
424 accuracy with increasing number of harmonics was observed for SVM and RF, trained on the  
425 herring dataset. In the case of increasing dimensionality, the ML algorithms clearly

426 outperform traditional classifiers due to their ability to integrate information from many  
427 variables without the high risk of overfitting (Breiman 2001; Ben-Hur et al. 2008).  
428 Improvement of the ML models accuracy can also be obtained by the elimination of non-  
429 informative variables during the model building (e.g., Smoliński 2019). Furthermore,  
430 heterogeneous ensemble techniques combining predictions of different model types could also  
431 be applied to improve the classification of fish stocks. Such an approach could help to  
432 minimize model-specific errors in class predictions and to obtain a more robust assignment of  
433 the fish origin.

434 The ability of SVM and other ML algorithms to model complex and non-linear patterns  
435 without any assumptions is of great importance in many biological applications (Noble 2006).  
436 Therefore, the variable transformations are not needed for the application of these algorithms,  
437 which make the pre-processing more straightforward and faster. Moreover, variables with  
438 non-normal distribution (typically required for the traditional parametric models) do not need  
439 to be excluded after an unsuccessful transformation, preventing from the loss of information  
440 potentially valuable for the discrimination of fish groups (Mercier et al. 2011).

441 Future operationalization of developing stock discrimination methods needs profound  
442 analyses of the level of temporal variability of within- and between-group differences,  
443 particularly in otolith shapes. The presented results are based on the samples collected within  
444 a short period of time, limiting the influence of the year-classes and long-term environmental  
445 effects on otolith shape. However, if the temporally stable character of fish otolith shapes can  
446 be confirmed for particular stocks, it may enable continuous enlargement of databases. In  
447 consequence, better performance of ML algorithms can be achieved, because their  
448 classification accuracy typically boosts with increasing size of training datasets.

449

## 450 **4.5 Conclusions**

451 Our study emphasized the potential for applying novel ML algorithms to improve the  
452 accuracy of classification systems based on the otolith shape of fish. We recommend  
453 conducting comparisons of different statistical classifiers in systems of well-identified stock  
454 structures using validated baselines. When temporal mixing of different fish stocks or stock  
455 components occurs, as with Baltic cod and herring in the Northeast Atlantic, possible  
456 improvements of stock discrimination processes by modern classifiers may be of great  
457 importance. More accurate assignment of fish individuals may help to more precisely estimate  
458 the contribution of different fish stocks within the mixing areas and in consequence, provide a  
459 more reliable quantitative basis for annual stock assessments and scientific advice.

460

#### 461 **Acknowledgments**

462 We thank Thomas Naatz (MS “JULE”) for providing cod samples from SD 23 and all staff  
463 members involved in sampling during research and monitoring cruises. We are grateful also  
464 to the technical staff at the Thuenen Institute of Baltic Sea Fisheries for photographing cod  
465 otoliths. We thank Tomas Gröhsler for providing herring otoliths from Greifswald Bay.  
466 Institute of Marine Research technicians are thanked for their contribution in collecting and  
467 photographing otoliths of the NSS and CSS herring components. We also acknowledge  
468 Audrey J. Geffen, Uwe Krumme, Richard D. M. Nash and two anonymous reviewers for the  
469 input and comments on this manuscript. FMS was partly funded by the European Maritime  
470 and Fisheries Fund (EMFF) of the European Union (EU) under the Data Collection  
471 Framework (DCF, Regulation 2017/1004 of the European Parliament and of the Council). FB  
472 was funded by the Research Council Norway project 254774 (GENSINC).

473

#### 474 **References**

475 Begg, G.A., Friedland, K.D., and Pearce, J.B. 1999. Stock identification and its role in stock

- 476 assessment and fisheries management: An overview. *Fish. Res.* **43**: 1–8.  
477 doi:10.1016/S0165-7836(99)00062-4.
- 478 Ben-Hur, A., Ong, C.S., Sonnenburg, S., Schölkopf, B., and Rätsch, G. 2008. Support vector  
479 machines and kernels for computational biology. *PLoS Comput. Biol.* **4**: e1000173.  
480 doi:10.1371/journal.pcbi.1000173.
- 481 Benjamini, Y., and Hochberg, Y. 1995. Controlling the false discovery rate: a practical and  
482 powerful approach to multiple testing. *J. R. Stat. Soc. B* **57**(1): 289–300.  
483 doi:10.2307/2346101.
- 484 Benzinou, A., Carbini, S., Nasreddine, K., Elleboode, R., and Mahé, K. 2013. Discriminating  
485 stocks of striped red mullet (*Mullus surmuletus*) in the Northwest European seas using  
486 three automatic shape classification methods. *Fish. Res.* **143**: 153–160.  
487 doi:10.1016/j.fishres.2013.01.015.
- 488 Berg, F., Almeland, O.W., Skadal, J., Slotte, A., Andersson, L., and Folkvord, A. 2018.  
489 Genetic factors have a major effect on growth, number of vertebrae and otolith shape in  
490 Atlantic herring (*Clupea harengus*). *PLoS One* **13**(1): 1–16.  
491 doi:10.1371/journal.pone.0190995.
- 492 Bonhomme, V., Picq, S., Gaucherel, C., and Claude, J. 2014. Momocs: Outline analysis using  
493 R. *JSS J. Stat. Softw.* **56**(13).
- 494 Breiman, L. 2001. Random forests. *Mach. Learn.* **45**: 5–32. doi:10.1023/A:1010933404324.
- 495 Brian, R., Venables, B., Bates, D.M., Firth, D., and Ripley, M.B. 2015. Package MASS.
- 496 Burke, N., Brophy, D., and King, P.A. 2008. Shape analysis of otolith annuli in Atlantic  
497 herring (*Clupea harengus*); a new method for tracking fish populations. *Fish. Res.* **91**:  
498 133–143. doi:10.1016/j.fishres.2007.11.013.
- 499 Cadrin, S.X., Kerr, L.A., and Mariani, S. 2014. Stock identification methods: Applications in  
500 fishery science. *In* 2nd edition. Elsevier. doi:10.1016/B978-0-12-397003-9.01001-8.

- 501 Campana, S.E., and Casselman, J.M. 1993. Stock discrimination using otolith shape analysis.  
502 Can. J. Fish. Aquat. Sci. **50**(5): 1062–1083. doi:10.1139/f93-123.
- 503 Cardinale, M., Doering-Arjes, P., Kastowsky, M., and Mosegaard, H. 2004. Effects of sex,  
504 stock, and environment on the shape of known-age Atlantic cod (*Gadus morhua*)  
505 otoliths. Can. J. Fish. Aquat. Sci. **61**(2): 158–167. doi:10.1139/F03-151.
- 506 Cortes, C., and Vapnik, V. 1995. Support-vector networks. Mach. Learn. **20**(3): 273–297.  
507 doi:10.1007/BF00994018.
- 508 Fernández-Delgado, M., Cernadas, E., Barro, S., and Amorim, D. 2014. Do we need hundreds  
509 of classifiers to solve real world classification problems? J. Mach. Learn. Res. **15**: 3133–  
510 3181.
- 511 Finn, J.E., Burger, C. V., and Holland-Bartels, L. 1997. Discrimination among populations of  
512 sockeye salmon fry with Fourier analysis of otolith banding patterns formed during  
513 incubation. Trans. Am. Fish. Soc. **126**: 559–578.
- 514 Fisher, R. 1936. The use of multiple measurements in taxonomic problems. Ann. Eugen. **7**(2):  
515 179–188.
- 516 Geffen, A. 2009. Advances in herring biology: from simple to complex, coping with plasticity  
517 and adaptability. ICES J. Mar. Sci. **66**(8): 1688–1695.
- 518 Guinand, B., Topchy, A., Page, K., Burnham-Curtis, M., Punch, W., and Scribner, K. 2002.  
519 Comparisons of likelihood and machine learning methods of individual classification. J.  
520 Hered. **93**(4): 260–269. doi:10.1093/jhered/93.4.260.
- 521 Guisan, A., and Zimmermann, N.E. 2000. Predictive habitat distribution models in ecology.  
522 Ecol. Modell. **135**: 147–186. doi:10.1016/S0304-3800(00)00354-9.
- 523 Haines, J., and Crampton, J.S. 2000. Improvements to the method of Fourier shape analysis as  
524 applied in morphometric studies. Paleontology **43**(4): 765–783.
- 525 Hall, P., Park, B.U., and Samworth, R.J. 2008. Choice of neighbor order in nearest-neighbor

- 526 classification. *Ann. Stat.* **36**(5): 2135–2152. doi:10.1214/07-AOS537.
- 527 Harbitz, A., and Albert, O.T. 2015. Pitfalls in stock discrimination by shape analysis of otolith  
528 contours. *ICES J. Mar. Sci.* **72**(7): 2090–2097. doi:10.1093/icesjms/fsv048.
- 529 Hastie, T., Tibshirani, R., and Friedman, J. 2009. *The elements of statistical learning - Data*  
530 *mining, inference.* Springer Verlag. doi:10.1007/b94608.
- 531 Hemmer-Hansen, J., Hüsey, K., Baktoft, H., Huwer, B., Bekkevold, D., Haslob, H.,  
532 Herrmann, J., Hinrichsen, H., Krumme, U., Mosegaard, H., Nielsen, E.E., Reusch,  
533 T.B.H., Storr-Paulsen, M., Velasco, A., von Dewitz, B., Dierking, J., and Eero, M. 2018.  
534 Genetic analyses reveal complex dynamics within a marine fish management area. *Evol.*  
535 *Appl.* **12**: 830–844. doi:10.1111/eva.12760.
- 536 Horbowy, J. 2005. Assessing Baltic herring stocks with a model that incorporates migration.  
537 *Fish. Res.* **76**: 266–276. doi:10.1016/j.fishres.2005.06.013.
- 538 Hüsey, K., Mosegaard, H., Albertsen, C.M., Nielsen, E.E., Hemmer-Hansen, J., and Eero, M.  
539 2016. Evaluation of otolith shape as a tool for stock discrimination in marine fishes using  
540 Baltic Sea cod as a case study. *Fish. Res.* **174**: 210–218.  
541 doi:10.1016/j.fishres.2015.10.010.
- 542 ICES. 2015. Report of the benchmark workshop on Baltic cod stocks (WKBALTCOD). 2-6  
543 March 2015, Rostock, Germany.
- 544 ICES. 2018a. Report of the workshop on mixing of western and central Baltic herring stocks  
545 (WKMixHER). 11-13 September, Gdynia, Poland.
- 546 ICES. 2018b. Report of the herring assessment working group for the area south of 62°N  
547 (HAWG), 29-31 January 2018 and 12-20 March 2018, ICES HQ, Copenhagen,  
548 Denmark.
- 549 ICES. 2019a. Baltic fisheries assessment working group (WGBFAS). *ICES Scientific Reports*  
550 **1**:20. 651 pp.



- 551 ICES. 2019b. Benchmark workshop on Baltic cod stocks (WKBALTCOD2). ICES Scientific  
552 Reports. 1:9. 310 pp. doi:http://doi.org/10.17895/ices.pub.4984.
- 553 ICES. 2019c. ICES Spatial Facility. Available from <http://gis.ices.dk> [accessed 29 May 2019].
- 554 Jones, W.A., and Checkley, D.M. 2017. Classification of otoliths of fishes common in the  
555 Santa Barbara Basin based on morphology and chemical composition. *Can. J. Fish.*  
556 *Aquat. Sci.* **74**(8): 1195–1207. doi:10.1139/cjfas-2015-0566.
- 557 Karatzoglou, A., Smola, A., and Hornik, K. 2015. Package ‘ kernlab .’
- 558 Kell, L.T., Crozier, W.W., and Legault, C.M. 2004. Mixed and multi-stock fisheries:  
559 Introduction. *ICES J. Mar. Sci.* **61**(8): 1330. doi:10.1016/j.icesjms.2004.09.002.
- 560 Kerr, L.A., Hintzen, N.T., Cadrin, S.X., Clausen, L.W., Dickey-Collas, M., Goethel, D.R.,  
561 Hatfield, E.M.C., Kritzer, J.P., and Nash, R.D.M. 2017. Lessons learned from practical  
562 approaches to reconcile mismatches between biological population structure and stock  
563 units of marine fish. *ICES J. Mar. Sci.* **74**(6): 1708–1722. doi:10.1093/icesjms/fsw188.
- 564 Kuhl, F.P., and Giardina, C.R. 1982. Elliptic Fourier features of closed contour. *Comput.*  
565 *Graph. Image Process.* **18**: 236–258.
- 566 Kuhn, M. 2008. Building predictive models in R using the caret package. *J. Stat. Softw.*  
567 **28**(5): 1–26. doi:10.1053/j.sodo.2009.03.002.
- 568 Liaw, A., and Wiener, M. 2002. Classification and regression by randomForest. *R news* **2**:  
569 18–22. doi:10.1177/154405910408300516.
- 570 Libungan, L.A., Óskarsson, G.J., Slotte, A., Jacobsen, J.A., and Pálsson, S. 2015. Otolith  
571 shape: a population marker for Atlantic herring *Clupea harengus*. *J. Fish Biol.* **86**: 1377–  
572 1395. doi:10.1111/jfb.12647.
- 573 Loh, W.Y. 2011. Classification and regression trees. *Wiley Interdisciplinary Rev. Data Min.*  
574 *Knowl. Discov.* **1**: 14–23. doi:10.1201/9781315139470.
- 575 Lord, C., Morat, F., Lecomte-Finiger, R., and Keith, P. 2012. Otolith shape analysis for three

- 576 Sicyopterus (Teleostei: Gobioidae: Sicydiinae) species from New Caledonia and  
577 Vanuatu. *Environ. Biol. Fishes* **93**: 209–222. doi:10.1007/s10641-011-9907-y.
- 578 Mapp, J., Hunter, E., Van Der Kooij, J., Songer, S., and Fisher, M. 2017. Otolith shape and  
579 size: The importance of age when determining indices for fish-stock separation. *Fish.*  
580 *Res.* **190**: 43–52. doi:10.1016/j.fishres.2017.01.017.
- 581 Mercier, L., Darnaude, A.M., Bruguier, O., Vasconcelos, R.P., Cabral, H.N., Costa, M.J.,  
582 Lara, M., Jones, D.L., and Mouillot, D. 2011. Selecting statistical models and variable  
583 combinations for optimal classification using otolith microchemistry. *Ecol. Appl.* **21**(4):  
584 1352–1364. doi:10.1890/09-1887.1.
- 585 Nasreddine, K., Benzinou, A., and Fablet, R. 2009. Shape geodesics for the classification of  
586 calcified structures: Beyond Fourier shape descriptors. *Fish. Res.* **98**: 8–15.  
587 doi:10.1016/j.fishres.2009.03.008.
- 588 Natterer, M., and Neumann, S. 2008. GNU image manipulation program. The GIMP Team.
- 589 Noble, W.S. 2006. What is a support vector machine? *Nat. Biotechnol.* **24**(12): 1565–1567.  
590 doi:10.1038/nbt1206-1565.
- 591 Olden, J.D., Lawler, J.J., and Poff, N.L. 2008. Machine learning methods without tears: a  
592 primer for ecologists. *Q. Rev. Biol.* **83**(2): 171–193. doi:10.1086/587826.
- 593 Parisi-Baradad, V., Lombarte, A., Garcia-Ladona, E., Cabestany, J., Piera, J., and Chic, O.  
594 2005. Otolith shape contour analysis using affine transformation invariant wavelet  
595 transforms and curvature scale space representation. *Mar. Freshw. Res.* **56**: 795–804.  
596 doi:10.1071/MF04162.
- 597 Paul, K., Oeberst, R., and Hammer, C. 2013. Evaluation of otolith shape analysis as a tool for  
598 discriminating adults of Baltic cod stocks. *J. Appl. Ichthyol.* **29**: 743–750.  
599 doi:10.1111/jai.12145.
- 600 Perdiguero-Alonso, D., Montero, F.E., Kostadinova, A., Raga, J.A., and Barrett, J. 2008.

- 601 Random forests, a novel approach for discrimination of fish populations using parasites  
602 as biological tags. *Int. J. Parasitol.* **38**(12): 1425–1434. doi:10.1016/j.ijpara.2008.04.007.
- 603 Petursdottir, G., Begg, G.A., and Marteinsdottir, G. 2006. Discrimination between Icelandic  
604 cod (*Gadus morhua* L.) populations from adjacent spawning areas based on otolith  
605 growth and shape. *Fish. Res.* **80**: 182–189. doi:10.1016/j.fishres.2006.05.002.
- 606 Ponton, D. 2006. Is geometric morphometrics efficient for comparing otolith shape of  
607 different fish species? *J. Morphol.* **267**: 750–757. doi:10.1002/jmor.10439.
- 608 R Core Team. 2018. R: A language and environment for statistical computing. Vienna,  
609 Austria.
- 610 Reig-Bolaño, R., Marti-Puig, P., Lombarte, A., Soria, J.A., and Parisi-Baradad, V. 2010a. A  
611 new otolith image contour descriptor based on partial reflection. *Environ. Biol. Fishes*  
612 **89**: 579–590. doi:10.1007/s10641-010-9700-3.
- 613 Reig-Bolaño, R., Marti-Puig, P., Rodriguez, S., Bajo, J., Parisi-Baradad, V., and Lombarte, A.  
614 2010b. Otoliths identifiers using image contours EFD. *Adv. Intell. Soft Comput.*  
615 doi:10.1007/978-3-642-14883-5\_2.
- 616 Robotham, H., Bosch, P., Gutiérrez-Estrada, J.C., Castillo, J., and Pulido-Calvo, I. 2010.  
617 Acoustic identification of small pelagic fish species in Chile using support vector  
618 machines and neural networks. *Fish. Res.* **102**: 115–122.  
619 doi:10.1016/j.fishres.2009.10.015.
- 620 Sadighzadeh, Z., Valinassab, T., Vosugi, G., Motallebi, A.A., Fatemi, M.R., Lombarte, A.,  
621 and Tuset, V.M. 2014. Use of otolith shape for stock identification of John's snapper,  
622 *Lutjanus johnii* (Pisces: Lutjanidae), from the Persian Gulf and the Oman Sea. *Fish. Res.*  
623 **155**: 59–63. doi:10.1016/j.fishres.2014.02.024.
- 624 Schade, F., Weist, P., and Krumme, U. 2019. Evaluation of four stock discrimination methods  
625 to assign individuals from mixed-stock fisheries using genetically validated baseline

- 626 samples. *Mar. Ecol. Prog. Ser.*: Accepted.
- 627 Smoliński, S. 2019. Incorporation of optimal environmental signals in the prediction of fish  
628 recruitment using random forest algorithms. *Can. J. Fish. Aquat. Sci.* **76**: 15–27.  
629 doi:<https://doi.org/10.1139/cjfas-2017-0554>.
- 630 Stransky, C. 2014. Morphometric outlines. *In* Stock identification methods: Applications in  
631 fishery science, 2nd edition. Elsevier Academic Press. pp. 129–140.
- 632 Stransky, C., Baumann, H., Fevolden, S.E., Harbitz, A., Høie, H., Nedreaas, K.H., Salberg,  
633 A.B., and Skarstein, T.H. 2008a. Separation of Norwegian coastal cod and Northeast  
634 Arctic cod by outer otolith shape analysis. *Fish. Res.* **90**: 26–35.  
635 doi:[10.1016/j.fishres.2007.09.009](https://doi.org/10.1016/j.fishres.2007.09.009).
- 636 Stransky, C., Murta, A.G., Schlickeisen, J., and Zimmermann, C. 2008b. Otolith shape  
637 analysis as a tool for stock separation of horse mackerel (*Trachurus trachurus*) in the  
638 Northeast Atlantic and Mediterranean. *Fish. Res.* **89**(2): 159–166.  
639 doi:[10.1016/j.fishres.2007.09.017](https://doi.org/10.1016/j.fishres.2007.09.017).
- 640 Taylor, N.G., McAllister, M.K., Lawson, G.L., Carruthers, T., and Block, B.A. 2011. Atlantic  
641 bluefin tuna: A novel multistock spatial model for assessing population biomass. *PLoS*  
642 *One* **6**(12): e27693. doi:[10.1371/journal.pone.0027693](https://doi.org/10.1371/journal.pone.0027693).
- 643 Torres, G.J., Lombarte, A., and Morales-Nin, B. 2000. Sagittal otolith size and shape  
644 variability to identify geographical intraspecific differences in three species of the genus  
645 *Merluccius*. *J. Mar. Biol. Assoc. UK* **80**(02): 333–342.  
646 doi:[10.1017/S0025315499001915](https://doi.org/10.1017/S0025315499001915).
- 647 Tracey, S.R., Lyle, J.M., and Duhamel, G. 2006. Application of elliptical Fourier analysis of  
648 otolith form as a tool for stock identification. *Fish. Res.* **77**: 138–147.  
649 doi:[10.1016/j.fishres.2005.10.013](https://doi.org/10.1016/j.fishres.2005.10.013).
- 650 Vieira, A.R., Neves, A., Sequeira, V., Paiva, R.B., and Gordo, L.S. 2014. Otolith shape

- 651 analysis as a tool for stock discrimination of forkbeard (*Phycis phycis*) in the Northeast  
652 Atlantic. *Hydrobiologia* **728**: 103–110. doi:10.1007/s10750-014-1809-5.
- 653 Vignon, M., and Morat, F. 2010. Environmental and genetic determinant of otolith shape  
654 revealed by a non-indigenous tropical fish. *Mar. Ecol. Prog. Ser.* **411**: 231–241.  
655 doi:10.3354/meps08651.
- 656 Weist, P., Schade, F.M., Damerau, M., Barth, J.M.I., Dierking, J., André, C., Petereit, C.,  
657 Reusch, T., Jentoft, S., Hanel, R., and Krumme, U. 2019. Assessing SNP-markers to  
658 study population mixing and ecological adaptation in Baltic cod. *PLoS One* **14**(6):  
659 e0218127. doi:10.1371/journal.pone.0218127.
- 660 Zhang, C., Ye, Z., Li, Z., Wan, R., Ren, Y., and Dou, S. 2016. Population structure of  
661 Japanese Spanish mackerel *Scomberomorus niphonius* in the Bohai Sea, the Yellow Sea  
662 and the East China Sea: evidence from random forests based on otolith features. *Fish.*  
663 *Sci.* **82**(2): 251–256. doi:10.1007/s12562-016-0968-x.
- 664 Zhang, C., Ye, Z., Panhwar, S.K., and Shen, W. 2013. Stock discrimination of the Japanese  
665 Spanish mackerel (*Scomberomorus niphonius*) based on the otolith shape analysis in the  
666 Yellow Sea and Bohai Sea. *J. Appl. Ichthyol.* **29**(2): 368–373. doi:10.1111/jai.12084.
- 667 Zhuang, L., Ye, Z., and Zhang, C. 2014. Application of otolith shape analysis to species  
668 separation in *Sebastes* spp. from the Bohai Sea and the Yellow Sea, northwest Pacific.  
669 *Environ. Biol. Fishes* **98**(2): 547–558. doi:10.1007/s10641-014-0286-z.
- 670 Zuur, A.F., Ieno, E.N., and Elphick, C.S. 2010. A protocol for data exploration to avoid  
671 common statistical problems. *Methods Ecol. Evol.* **1**: 3–14. doi:10.1111/j.2041-  
672 210X.2009.00001.x.

673 **Figures**

674

675 Fig. 1. Distribution of sampling locations of cod and herring. The shape and color of the  
676 points indicate the fish species and stock component, respectively. Size of the point shows the  
677 number of fish analyzed from the given location. The map was created based on the layer of  
678 ICES statistical areas (ICES 2019c).

679

680 Fig. 2. Cumulative Fourier power ( $PF_c$ ) calculated for cod and herring showing examples of  
681 reconstructions of otolith outline with different numbers of harmonics. The box represents the  
682 interquartile range (IQR) with the median (midline) and the first and third quantiles at the  
683 bottom and top of the box, respectively. Lower and upper whiskers are restricted to 1.5 x IQR  
684 and black dots represent outliers.

685

686 Fig. 3. Principal component analysis (PCA) conducted on the Fourier coefficients of otolith  
687 shape for cod (a) and herring (b). The levels of variance explained by the first PCA axes are  
688 shown on the axes. The morphospace plotted over the observations represents theoretical  
689 shapes reconstructed based on the PCA scores.

690

691 Fig. 4. Classification accuracy of different statistical models based on different numbers of  
692 Fourier harmonics of otolith shapes. Lines represent median accuracy, shades 10<sup>th</sup> and 90<sup>th</sup>  
693 percentile. Models in the legend were arranged according to the median accuracy of  
694 classification on the dataset with highest number of harmonics.

695

Fig. 5. Variable (Fourier descriptors) relative importance obtained for cod (a) and herring (b)  
from otolith shape classification models.

## Tables

Table 1. Summary of analyzed samples including fish species, stocks, components, capture years, sample size (N), mean total fish length (TL)  $\pm$  standard deviation (SD), mean and range of age. \*Due to age reading difficulties of eastern Baltic cod (EBC), age was only determined for western Baltic cod (WBC) captured in SD22 and SD23. NA= not available.

Species	Stock	Component	Years	N	Mean TL $\pm$ SD [cm]	Mean age	Age range
Cod	EBC		2015, 2016	243	43.11 $\pm$ 5.24	NA	NA
Cod	WBC		2015, 2016	264	47.71 $\pm$ 9.86	2.89*	1-6*
Herring	WBSS	CSS	2006, 2012, 2017	157	29.25 $\pm$ 1.40	5.40	5-6
Herring	NSS	NSS	2018	207	31.08 $\pm$ 1.63	5.20	5-6
Herring	CBH	CBNC	2017	170	19.39 $\pm$ 1.81	5.51	5-6
Herring	WBSS	GB	2018	238	27.63 $\pm$ 1.32	5.28	5-6

Table 2. Comparison of algorithm accuracies for cod (a) and herring (b). Dataset for the comparison was built on 13 harmonics (accounting for 99% variance of the otolith shape). Estimates of the difference (% accuracy) are reported in the upper diagonals, while p-values (with Bonferroni adjustment) for the hypothesis of no difference are reported in the lower diagonals.

a) cod	LDA	QDA	KNN	CART	RF	SVM
LDA		6.63	7.29	12.18	3.50	-0.90
QDA	< 0.001		0.66	5.55	-3.13	-7.53
KNN	< 0.001	0.0689		4.88	-3.79	-8.20
CART	< 0.001	< 0.001	< 0.001		-8.67	-13.08
RF	< 0.001	< 0.001	< 0.001	< 0.001		-4.41
SVM	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	
b) herring	LDA	QDA	KNN	CART	RF	SVM
LDA		-4.69	-0.48	8.80	-2.96	-11.20
QDA	< 0.001		4.21	13.49	1.73	-6.50
KNN	0.204	< 0.001		9.28	-2.48	-10.71
CART	< 0.001	< 0.001	< 0.001		-11.76	-19.99
RF	< 0.001	< 0.001	< 0.001	< 0.001		-8.24
SVM	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	



Table 3. Cross-validated (4-fold, repeated 100 times) confusion matrix obtained for cod and herring stocks with the best classifier (support vector machines) on Fourier descriptors of otolith shape. Entries are percentual average cell counts across resamples. Average accuracy (sum of diagonal cells): cod = 79.54; herring = 74.13.

Prediction	Reference					
	EBC	WBC	CSS	NSS	CBNC	GB
EBC	<b>38.0</b>	10.5				
WBC	9.9	<b>41.6</b>				
CSS			<b>10.7</b>	0.9	0.7	6.3
NSS			0.9	<b>23.1</b>	2.2	0.8
CBNC			0.4	2.3	<b>17.7</b>	1.1
GB			8.4	0.4	1.4	<b>22.6</b>

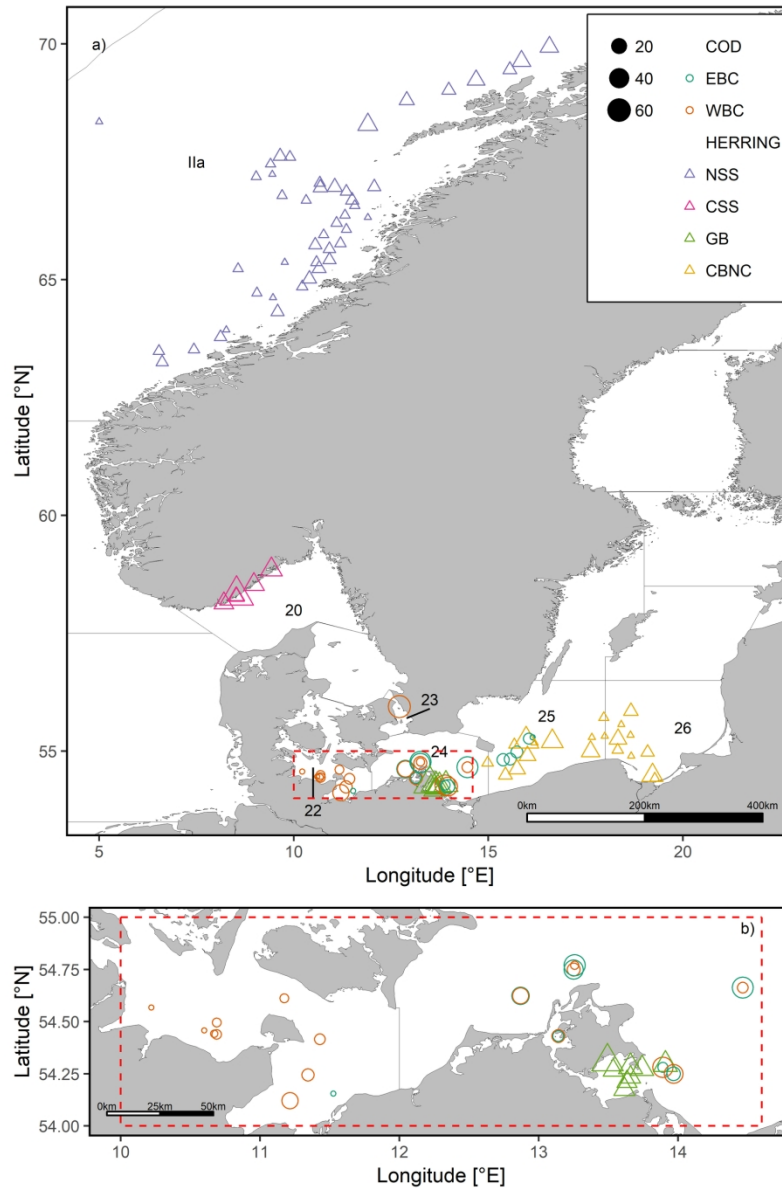


Fig. 1. Distribution of sampling locations of cod and herring. The shape and color of the points indicate the fish species and stock component, respectively. Size of the point shows the number of fish analyzed from the given location. The map was created based on the layer of ICES statistical areas (ICES 2019c).

152x228mm (300 x 300 DPI)

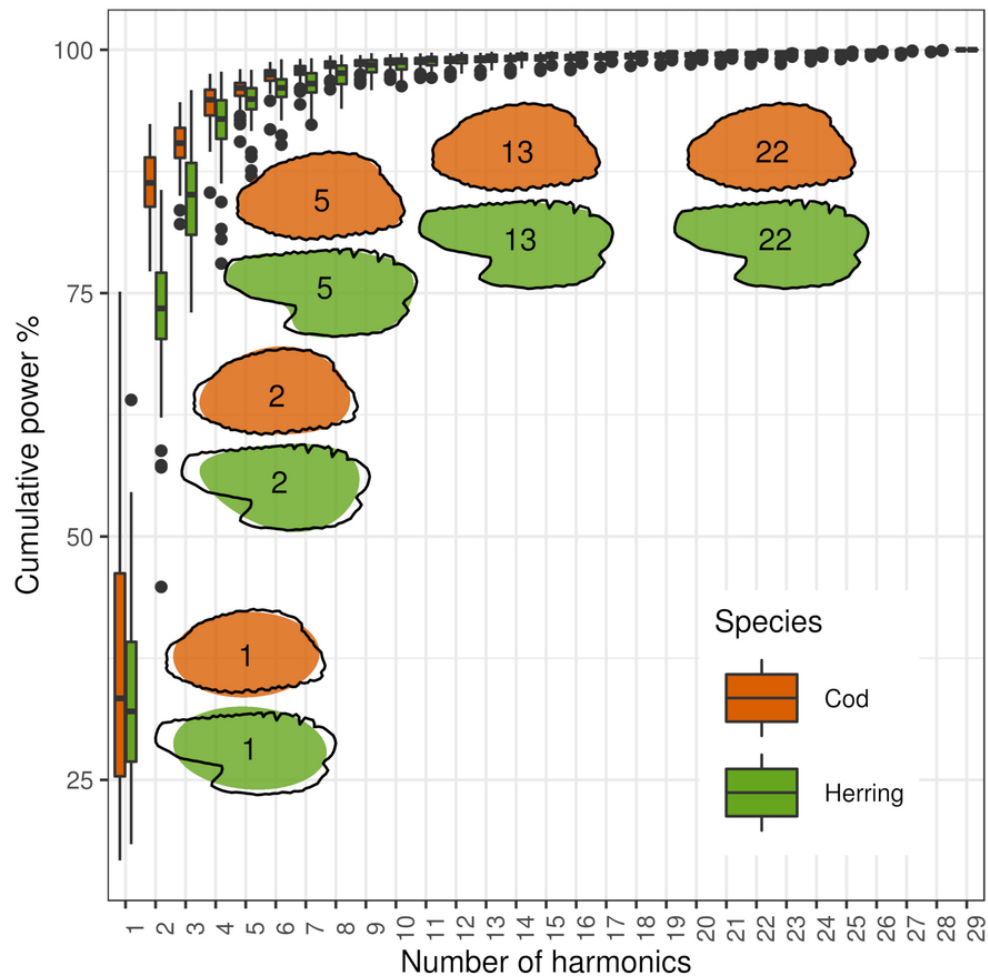


Fig. 2. Cumulative Fourier power ( $PF_C$ ) calculated for cod and herring showing examples of reconstructions of otolith outline with different numbers of harmonics. The box represents the interquartile range (IQR) with the median (midline) and the first and third quartiles at the bottom and top of the box, respectively. Lower and upper whiskers are restricted to 1.5 x IQR and black dots represent outliers.

76x76mm (300 x 300 DPI)

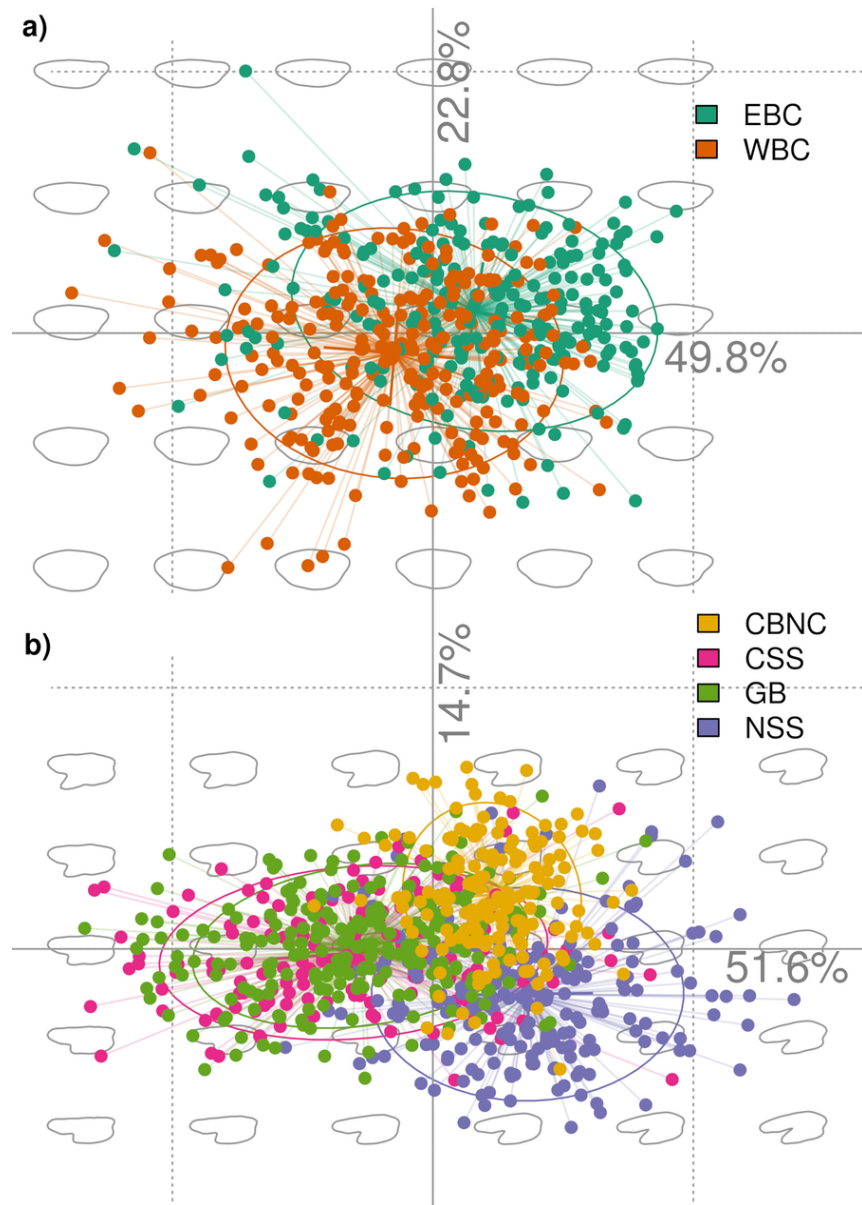


Fig. 3. Principal component analysis (PCA) conducted on the Fourier coefficients of otolith shape for cod (a) and herring (b). The levels of variance explained by the first PCA axes are shown on the axes. The morphospace plotted over the observations represents theoretical shapes reconstructed based on the PCA scores.

76x106mm (300 x 300 DPI)

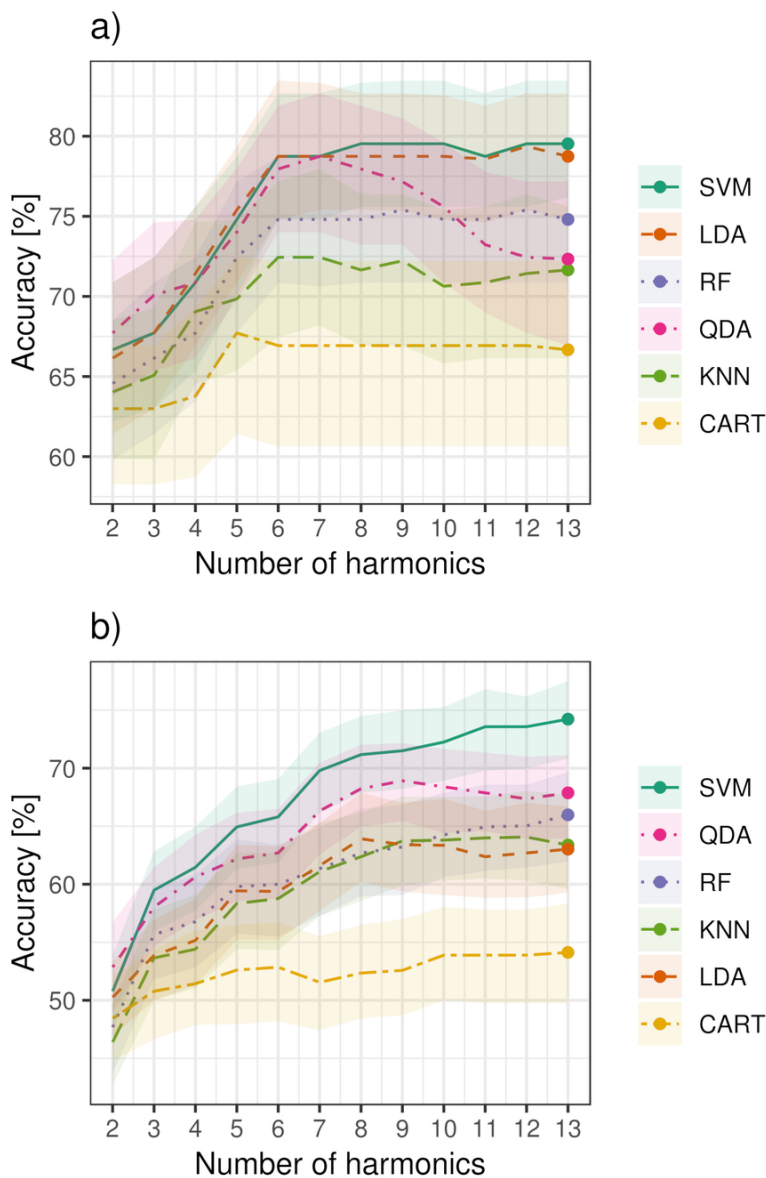


Fig. 4. Classification accuracy of different statistical models based on different numbers of Fourier harmonics of otolith shapes. Lines represent median accuracy, shades 10<sup>th</sup> and 90<sup>th</sup> percentile. Models in the legend were arranged according to the median accuracy of classification on the dataset with highest number of harmonics.

76x114mm (300 x 300 DPI)

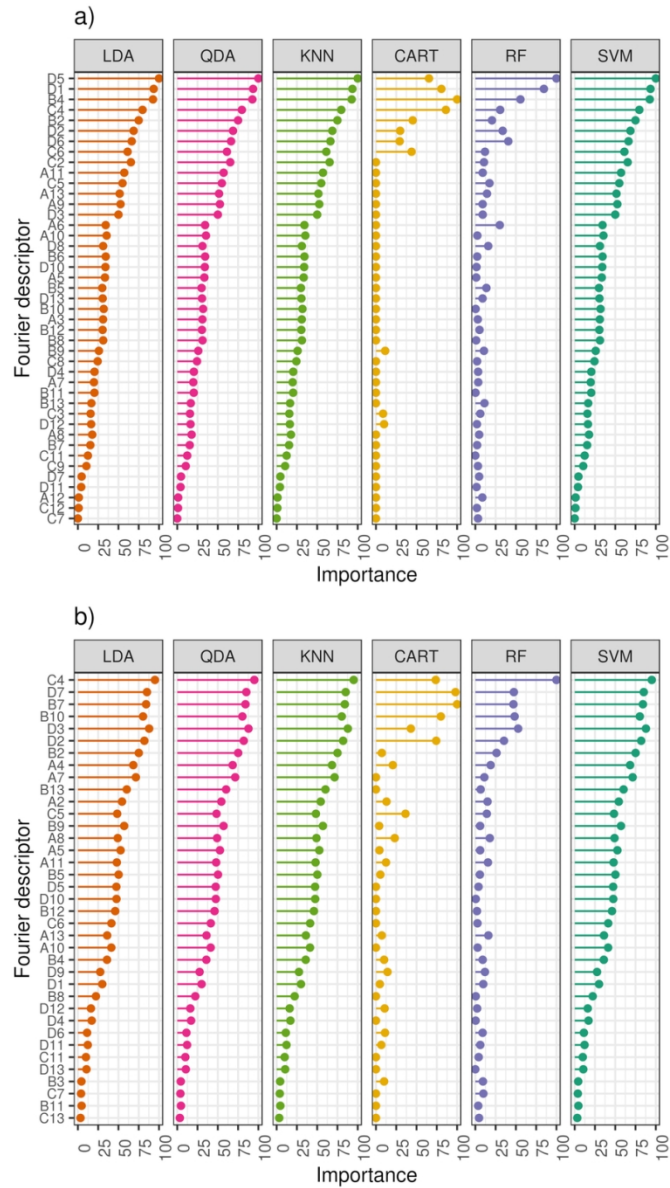


Fig. 5. Variable (Fourier descriptors) relative importance obtained for cod (a) and herring (b) from otolith shape classification models.

76x137mm (300 x 300 DPI)