

1 **Seismic survey noise reduces fin whale vocalisations offshore**  
2 **northwestern Spain**

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17  
18 **Abstract**

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20 The ever-increasing level of anthropogenic noise in the ocean is a global concern, yet its impacts  
21 on marine life remain poorly understood. Offshore seismic surveys generate intense and repetitive  
22 sound pulses during the firing of airguns, which potentially affect the behaviour of marine taxa  
23 including marine mammals, but fine-scale behavioural impacts are rarely quantified. We assessed  
24 the impact of a 2013 seismic survey offshore north-western Spain on fin whale vocalisations. Using  
25 a convolutional neural network trained on 50 hours of manually labelled acoustic data (accuracy  
26 85.6%, AUC 0.93), 63 days of continuous ocean-bottom recordings were analysed across four  
27 alternating periods with and without airgun shooting. Across three instruments, vocalisation  
28 detections decreased by 70% on average during shooting, with reductions of 52% after correcting  
29 for worst-case scenario masking. A negative binomial mixed-effects model indicated statistically  
30 significant ( $p = < 0.0001$ ) reductions in calling during shooting compared to quiet periods. Calling  
31 declined rapidly within 1–2 days after shooting began, suggesting reduced vocalisation or  
32 temporary displacement. While the broader ecological implications cannot be determined from  
33 this dataset, these findings indicate short-term behavioural responses to seismic activity and  
34 underscore the need for further research into potential consequences for communication,  
35 foraging, and habitat use.  
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## 45 Introduction

46  
47 The level of anthropogenic noise in the ocean is rising continuously as a consequence of the  
48 ongoing expansion and intensification of human activities [1–3]. A systematic review of the relevant  
49 research available from 1973-2020 regarding the impact of anthropogenic noise on marine animals  
50 found that 81–94 % of studies reported significant negative effects of anthropogenic noise across a  
51 range of marine taxa [4]. Marine mammals are especially vulnerable to these impacts due to their  
52 reliance on sound for essential life functions such as communication, navigation, and foraging [5–  
53 7]. Documented effects of noise on marine mammals include acoustic masking, avoidance or  
54 evasive movement, altered vocalisation patterns and elevated stress levels [5,8,9].

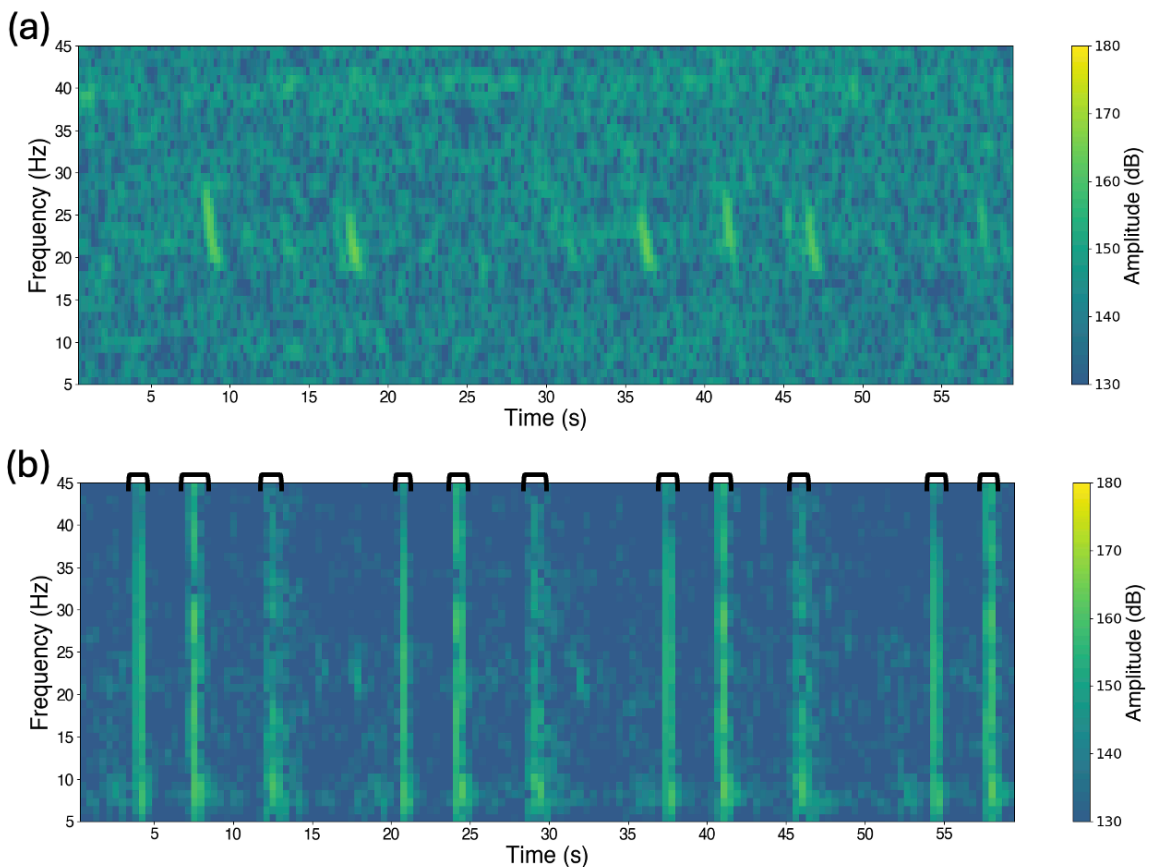
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56 Among anthropogenic sound sources, seismic surveys are of particular concern. This geophysical  
57 exploration technique is widely employed for the purpose of seabed mapping and hydrocarbon  
58 exploration, typically carried out by the deployment of vessel-towed airgun arrays to produce high-  
59 resolution images of the Earth’s subsurface [10]. The repeated firing of these airguns is commonly  
60 referred to as ‘shooting’. During shooting, the rapid expansion and contraction of the air bubble  
61 formed during the airguns’ high-pressure release generates sound waves that propagate into the  
62 surrounding water column and subsurface beneath [2]. The returning sound signals are detected  
63 by receivers in the ocean and processed to image the subsurface structure. Airguns produce  
64 intense and impulsive sounds, typically recurring every 10–20 seconds, with low-frequency peak  
65 spectral energy at 5–300 Hz and source levels of 200–250 dB [2,11]. These airgun pulses can  
66 propagate over long distances, having been recorded more than 3,000 km from their source [12],  
67 while their dominant frequencies overlap with the vocalisation range of baleen whales (10 Hz to 1  
68 kHz) [13]. Given that sound is fundamental for marine mammal communication and survival, this  
69 overlap raises particular concerns about the impacts of seismic airgun activity on these  
70 populations [4,14–16].

71  
72 Baleen whales are often adversely affected by anthropogenic noise, with 85% of relevant studies  
73 reporting negative impacts [4]. In response to seismic survey noise, research to date has reported a  
74 variety of behavioural responses in baleen whales. For example, bowhead whales (*Balaena*  
75 *mysticetus*) avoided areas within 20 km of active airgun operations [17], while gray whales  
76 (*Eschrichtius robustus*) altered their movement and respiration in response to increasing sound  
77 exposure from seismic survey vessels [18]. Other studies have observed a range of acoustic  
78 changes in baleen whales exposed to seismic surveys, including alterations in vocalisation  
79 patterns and call rates in bowhead whales (*Balaena mysticetus*) [19], fin whales (*Balaenoptera*  
80 *physalus*) [14], blue whales (*Balaenoptera musculus*) [15] and humpback whales (*Megaptera*  
81 *novaeangliae*) [20,21]. However, some studies have found no significant correlations between  
82 seismic activity and whale behaviour [22,23].

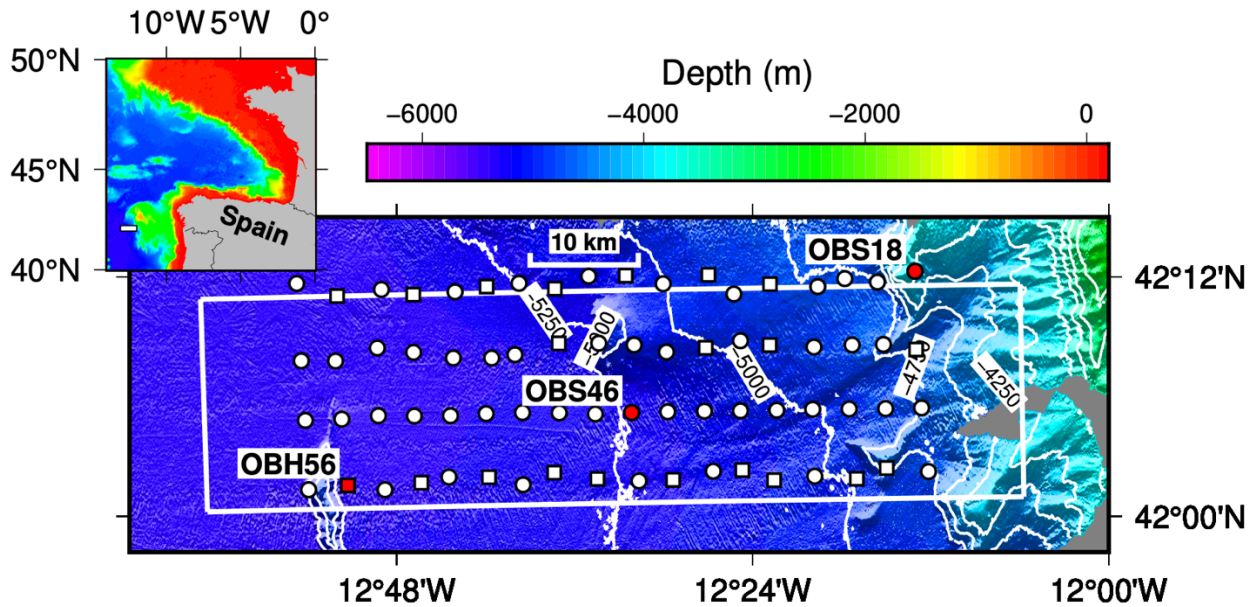
83  
84 Fin whales are widely distributed throughout the world’s oceans and are classified as Vulnerable  
85 by the IUCN Red List, due to ongoing concerns about their conservation status [24]. They use low-  
86 frequency vocalisations, the most commonly observed being the ‘20 Hz pulse’ [25,26], that are the  
87 focus of this study (Fig. 1). Although the precise function of this call remains uncertain, repetitive  
88 and stereotyped sequences of 20 Hz pulses have been suggested to represent reproductive  
89 displays [25] produced by males [27], whilst isolated or irregular pulses may occur across different  
90 demographic groups and appear linked to social or other behavioural contexts [28–30]. The overlap  
91 between these vocalisations and the dominant energy of seismic airgun pulses raises concerns of  
92 potential acoustic masking, behavioural disruption and/or spatial displacement effects on the

93 species. Despite such concerns, very few studies have examined the responses of fin whales to  
94 seismic activity, with the limited evidence indicating changes in vocal behaviour and potential  
95 large-scale spatiotemporal avoidance of survey areas [14].

96  
97 Located offshore northwestern Spain, the Deep Galicia Margin lies along known fin whale migratory  
98 routes [31]. In 2013, a large-scale 3D multichannel seismic reflection and wide-angle seismic  
99 survey was conducted in this region, including the deployment of 72 ocean-bottom recording  
100 instruments across an area of approximately 65 × 25 km (Fig. 2) [32,33]. Two 3,300 cubic inch  
101 airgun arrays were fired alternately from *RV Marcus Langseth* at ~16-second intervals along 50  
102 parallel survey lines, with a quiet interlude during vessel repairs in Vigo port providing a natural  
103 contrast between periods with and without shooting. This study used these contrasting periods to  
104 investigate potential impacts of seismic survey activity on fin whale acoustic behaviour. Hereafter,  
105 we refer to these periods as Shooting 1 (Julian Day (JD) 156-173), Quiet 1 (JD 174-196), Shooting 2  
106 (197-212) & Quiet 2 (JD 213-218). We used a Convolutional Neural Network (CNN) model to detect  
107 fin whale 20 Hz pulses within the passive acoustic recordings. By comparing vocal activity across  
108 periods of active airgun firing and relative quiet, we evaluated whether seismic operations  
109 influenced fin whale calling behaviour. This approach provided novel insights into how high  
110 amplitude, temporally persistent, and spatially pervasive anthropogenic noise can alter calling  
111 behaviour of fin whales along a key migratory corridor.  
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114 Figure 1  
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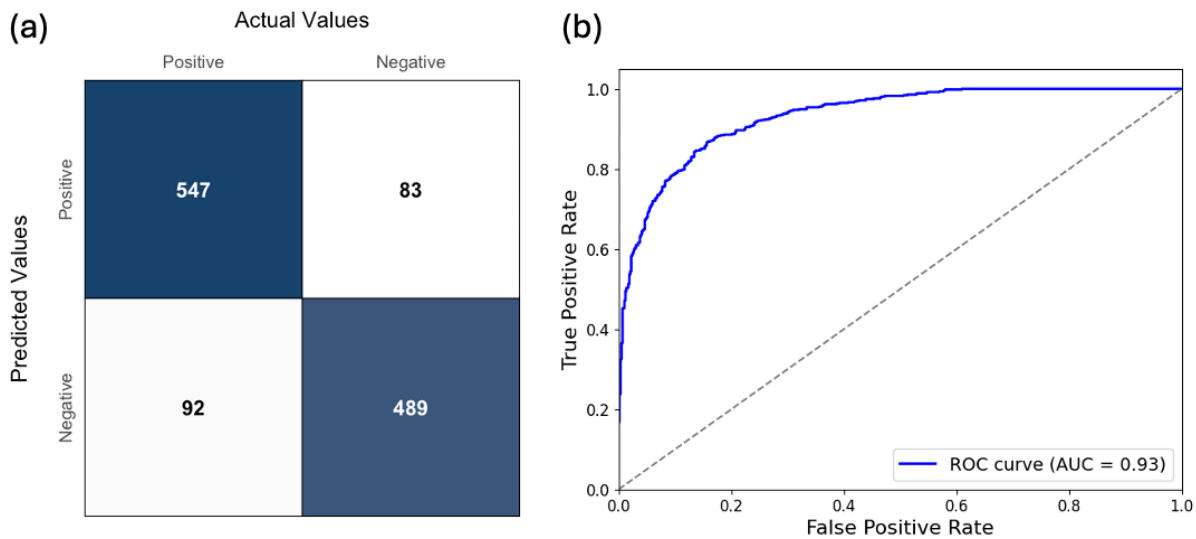
116  
117 Figure 2

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120 **Results**

121  
122 **Machine learning model.** The final CNN achieved an overall accuracy of 85.6% in detecting fin  
123 whale 20 Hz pulses on the validation dataset. Precision and recall were 86.8% and 85.6%,  
124 respectively, resulting in an F1 score of 86.2%. The confusion matrix for the CNN’s output on the  
125 validation dataset (Fig. 3a) was produced with a threshold of 0.5 applied to the CNN’s output  
126 predicted probabilities. The Receiver Operating Characteristic (ROC) curve, which computes true  
127 positive rates and false positive rates across thresholds from 0 to 1, had an Area Under the Curve  
128 (AUC) of 0.93 (Fig. 3b). These results showed that the CNN could reliably detect fin whale pulses in  
129 the analysed acoustic dataset.

130  
131 **Fin whale pulse detections.** Acoustic recordings were analysed from two ocean-bottom  
132 seismometers (OBSs) and one ocean-bottom hydrophone (OBH). Across the entire survey period  
133 (JD156–218), the CNN detector identified the highest average hourly number of 30-second periods  
134 containing pulses (“positive frames”) on OBH56 (mean = 46.9, range = 1–119), followed closely by  
135 OBS18 (mean = 45.3, range = 1–116). OBS46 recorded the fewest detections, with an average of  
136 36.1 positive frames per hour (range = 1–103). OBH56 and OBS18 had similar proportions of  
137 positive frames (39.1% and 37.7%, respectively), while OBS46 had a lower proportion (30.1%).  
138 Notably, these hourly trends were consistent across the survey, and preliminary analysis indicated  
139 no significant diel cycle in fin whale vocalisations, justifying our focus on hourly mean counts.

140



141  
142 Figure 3

143  
144 Hourly counts showed clear temporal patterns across the four survey periods (Fig. 4) that we  
145 named as Shooting 1 (JD156–173, 18 days), Quiet 1 (JD 174–196, 23 days), Shooting 2 (JD 197–212,  
146 16 days), and Quiet 2 (JD 213–218, 6 days). All instruments showed a sharp increase in detections  
147 at the onset of quiet periods, followed by a rapid decline when shooting resumed (Fig. 5). OBS18  
148 detections were consistently higher during quiet periods, with Quiet 1 and Quiet 2 averaging 74.4  
149 (range 15–116) and 83.0 (range 26–113) positive frames per hour, respectively. Counts declined  
150 during shooting periods, with Shooting 1 and Shooting 2 averaging 18.7 (range 1–102) and 14.0  
151 (range 1–79) frames per hour. OBS46 exhibited a similar temporal pattern. Quiet periods yielded  
152 higher hourly counts (Quiet 1: 60.2, range 3–101; Quiet 2: 51.4, range 1–97) compared to shooting  
153 periods (Shooting 1: 18.4, range 1–103; Shooting 2: 13.7, range 1–65). For OBH56, the general  
154 pattern was similar, although counts during shooting periods were slightly higher than the other  
155 instruments (Shooting 1: 30.2, range 1–83; Shooting 2: 24.3, range 4–107), while quiet periods  
156 remained the periods of highest activity (Quiet 1: 69.9, range 1–119; Quiet 2: 60.2, range 1–119).  
157 Calling activity dropped markedly between quiet and shooting periods, with mean detections  
158 reduced by 78.2% on OBS18, 72.3% on OBS46, and 60.8% on OBH56 (Table 1).

159  
160 Detection counts of fin whale calls varied strongly with survey period. A negative binomial mixed-  
161 effects model indicated strong reductions in detections during shooting relative to quiet periods,  
162 after accounting for instrument-level and day-level random effects (Supplementary Table S4a).  
163 Both shooting periods showed significant decreases in detections (Shooting 1:  $z = -15.1$  mean  
164 frames/hr,  $p < 0.001$ ; Shooting 2:  $z = -17.1$  mean frames/hr,  $p < 0.001$ ). Comparisons between quiet  
165 periods showed no significant overall change ( $z = -1.07$  mean frames/hr,  $p = 0.29$ ). Random effects  
166 revealed modest variation among instruments ( $SD = 0.12$ ) and stronger day-specific variation  
167 within instruments ( $SD = 0.43$ ), while the dispersion parameter ( $\theta = 6.22$ ) confirmed that the  
168 negative binomial distribution adequately accounted for overdispersion. Overall, detections were  
169 consistently higher during quiet periods than during shooting.

170  
171 Model-derived Estimated Marginal Means (EMMs) supported these patterns: predicted hourly  
172 detections were highest during Quiet 1 (64.6 mean frames/hr, 95% CI: 54.5–76.6) and Quiet 2 (57.1

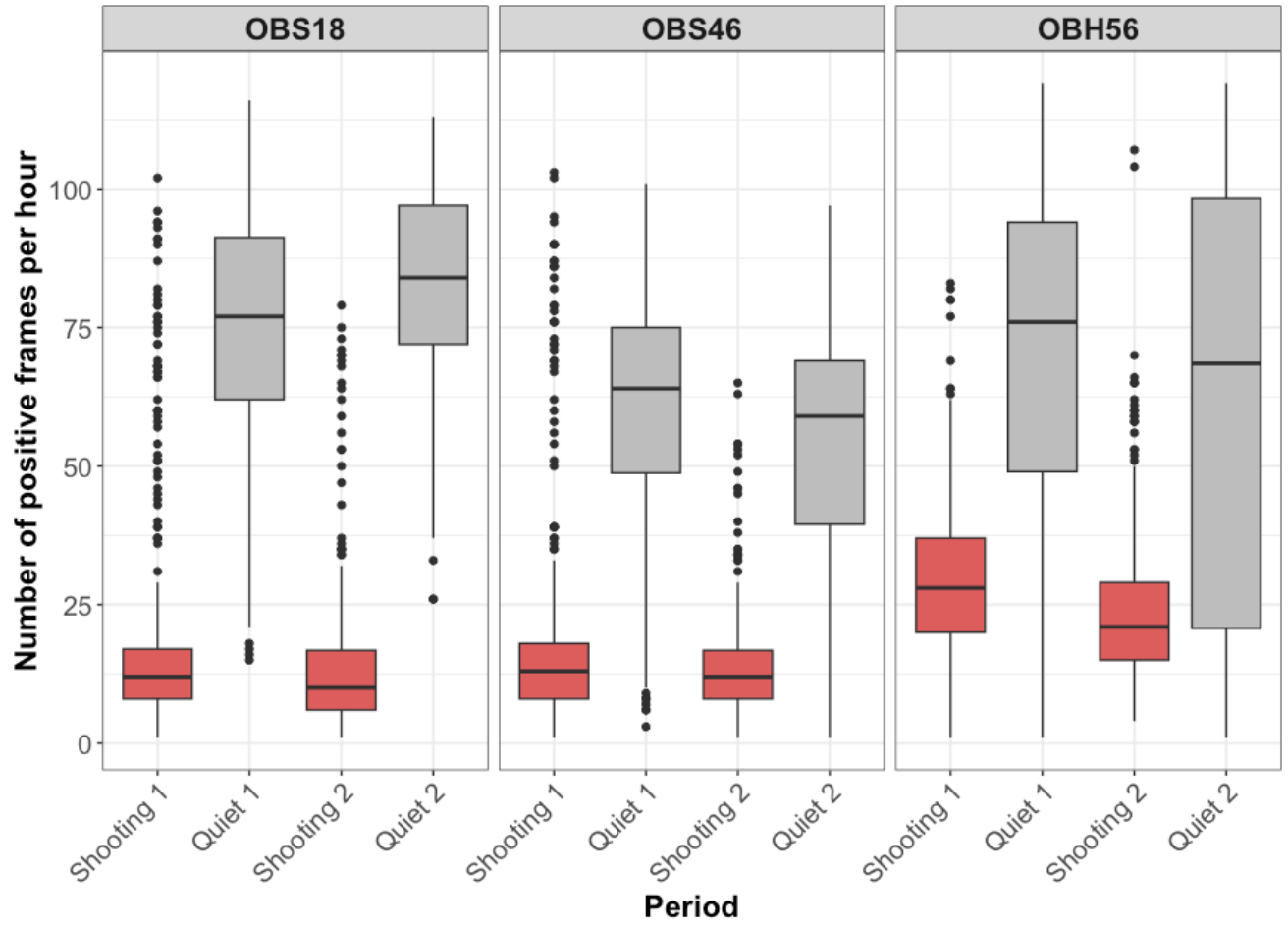
173 mean frames/hr, 95% CI: 44.7–72.8), and markedly lower during Shooting 1 (18.5 mean frames/hr,  
 174 95% CI:15.4–22.2) and Shooting 2 (15.4 mean frames/hr, 95% CI:12.8–18.5). Pairwise contrasts  
 175 showed significant differences for all quiet–shooting comparisons ( $p < 0.001$ ), while the contrasts  
 176 between the two quiet periods and the two shooting periods were not significant. Full EMMs and  
 177 pairwise comparisons of hourly detection counts are provided in Supplementary S4b and S4c.

178  
 179 Table 1

Instrument	Quiet mean (positive frames hr <sup>-1</sup> )	Shooting mean (frames hr <sup>-1</sup> )	Percentage drop (observed) (%)	Corrected shooting mean (frames hr <sup>-1</sup> , worst case)	Percentage drop (worst case) (%)
<b>OBS18</b>	76.2	16.6	78.2	23.2	69.6
<b>OBS46</b>	58.4	16.2	72.3	22.7	61.1
<b>OBH56</b>	67.8	26.6	60.8	37.2	45.1
<b>Mean</b>			<b>70.4</b>		<b>52.0</b>

180  
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 182  
 183 **Maximum masking estimation.** Application of the worst-case correction confirmed that seismic  
 184 airgun activity had a strong suppressive effect on fin whale detections, even under the highly  
 185 conservative assumption that every masked interval overlapped a fin whale call. Corrected mean  
 186 hourly detection counts during shooting remained consistently lower than during quiet periods  
 187 across all instruments (Table 1). The percentage decrease in calling activity when corrected for  
 188 masking was 45.1 – 69.6%.

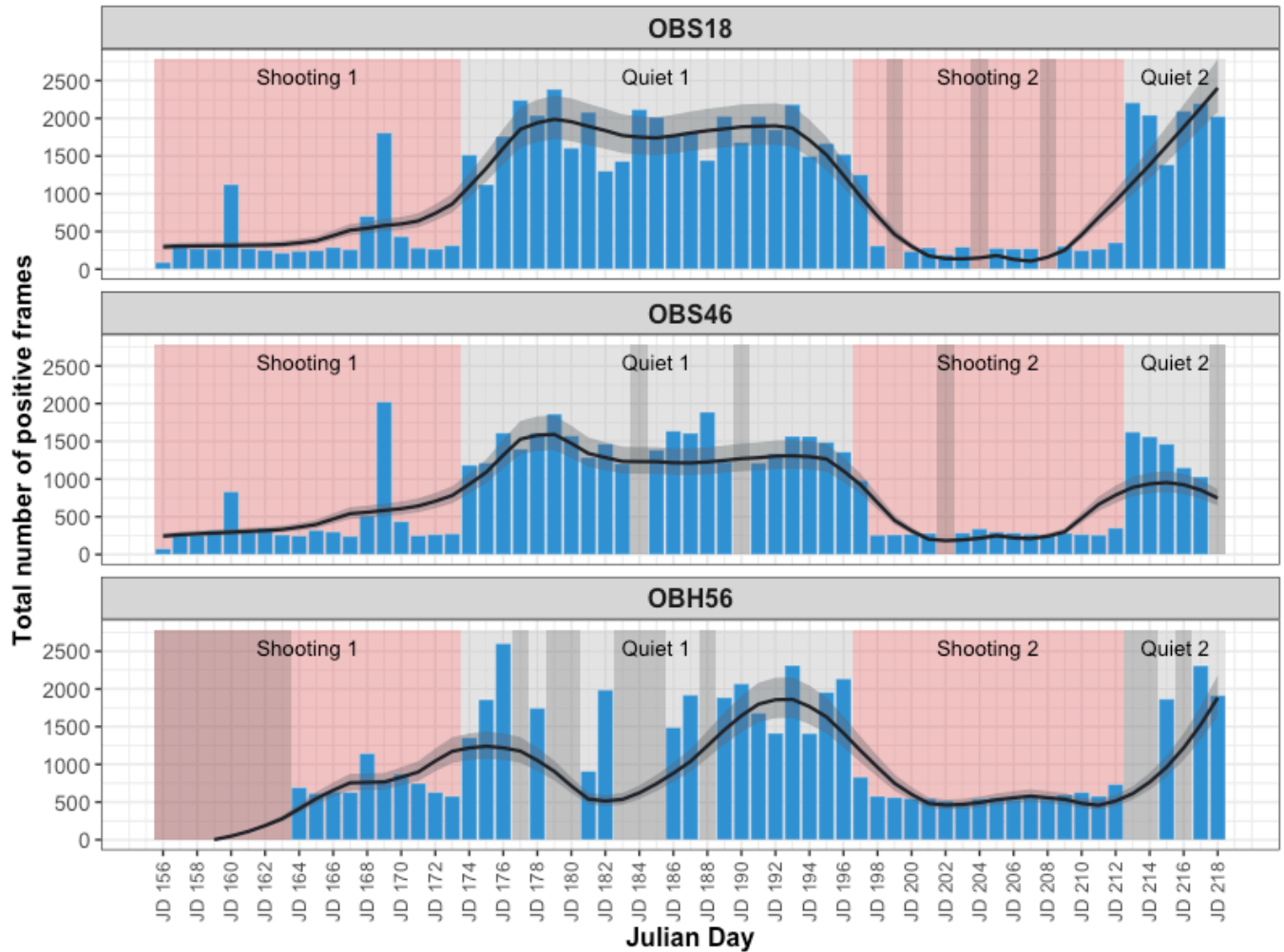
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192 Figure 4

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196 Figure 5

## 197 Discussion

198 The results of this study demonstrated a clear association between seismic airgun activity and  
199 reduced fin whale vocalisation rates across the survey period. Across all three instruments,  
200 periods of active airgun shooting were characterised by marked declines in hourly detections of fin  
201 whale 20-Hz pulses compared to quiet periods. On average, vocal activity decreased by 70.4%  
202 when airguns were firing (Table 1), highlighting the scale of disruption caused by seismic survey  
203 noise. This decrease was supported by the mixed-effects model, which found statistically  
204 significant drops in detections during shooting relative to quiet periods (Supplementary S4a).  
205

206 This observation contributes to a growing body of evidence that anthropogenic ocean noise has  
207 widespread and often adverse impacts on marine mammals [4,6,34]. However, the specific effects

208 of seismic surveys on fin whales remain underexplored, largely due to the logistical challenges of  
209 conducting research in offshore environments and the complexity of possible behavioural  
210 responses to noise [14,35]. Interpretation of reduced call activity is further complicated by the  
211 context-dependent and still partially unresolved function of fin whale 20-Hz pulses [29,36]. While  
212 this uncertainty limits our ability to infer precise ecological consequences, it does not alter the  
213 clear behavioural changes in our data: fin whales reduce their vocal activity during periods of active  
214 shooting. The present study therefore fills an important gap by documenting repeated and  
215 prolonged reduction of fin whale calling behaviour during airgun use in the Northeast Atlantic.  
216

217 The temporal patterns observed suggested that the reduction in detections is not random but  
218 reflects behavioural change in response to seismic noise. Periods of quiet were marked by  
219 immediate increases in detections, often within one to two days of airgun cessation. During the  
220 shooting periods, lower detections persisted throughout the periods, with Shooting 2 showing a  
221 rapid decline at onset (Fig. 5). Model-derived estimated marginal means showed the same pattern,  
222 with higher predicted detections during quiet periods than during either shooting period  
223 (Supplementary S4b and S4c). The elevated detections observed on JD160 and JD168–170 (Fig. 5)  
224 coincided with short operational pauses in airgun use reported in the cruise report, which occurred  
225 due to adverse weather conditions and equipment repairs [37]. While the underlying behavioural  
226 response cannot be determined, the temporal link indicates that these peaks were likely  
227 influenced by reduced noise exposure. The broader temporal patterns suggest that fin whales  
228 either reduced their calling rates, moved away from the survey area or exhibited a combination of  
229 these responses. While disentangling these mechanisms is beyond the scope of this study, both  
230 reduced vocal activity and displacement have been reported as common whale responses to  
231 seismic noise [14,20].  
232

233 In our dataset, fin whales maintained their responses across successive airgun shooting periods  
234 (Fig. 5), showing no signs of habituation, which would be characterised by diminished responses  
235 over time and continued calling during shooting [38]. We did not detect direct evidence for  
236 cumulative impacts but repeated responses could potentially contribute to such effects [14]. There  
237 is broader evidence that repeated or prolonged exposures to stressors, including noise, can be  
238 problematic for marine mammals, potentially leading to issues such as suppression of  
239 reproduction and accelerated aging [39]. Additionally, behavioural changes due to noise can lead  
240 to displacement from preferred habitats and negatively influence vital rates, as indicated by  
241 movement modelling [40]. If fin whales follow this broader pattern, and successive shooting  
242 periods are not fully isolated events, then repeated or prolonged exposure could exacerbate  
243 impacts on fin whale behaviour and habitat use.  
244

245 These observed findings align with the outcomes of previous studies, albeit from a very limited  
246 literature base on fin whales. Altered song structure and reduced detections have been  
247 documented during seismic surveys, with vocal activity remaining suppressed for two weeks after  
248 survey completion along with prompt decreases in fin whale pulses within the first 72 hours of  
249 active shooting [14]. Additionally, the tracking of individual fin whales has shown movements away  
250 from survey vessels during the onset of airgun use [41]. At a broader taxonomic scale, declines in  
251 humpback whale singing have been associated with seismic survey noise, interpreted as both a  
252 reduction in calling behaviour and possible avoidance of noisy areas [20,21]. In contrast, a study  
253 using the same OBS dataset did not detect a clear correlation between seismic activity and fin  
254 whale vocal activity, likely due to the limited sample size and use of data predominantly within

255 Quiet 1 period [23]. Overall, these studies have parallels with our results, strongly suggesting that  
256 the effects observed here are part of a wider pattern of seismic survey impacts on baleen whales.

257  
258 Masking of fin whale pulses by seismic airgun activity represents an unavoidable limitation in this  
259 study. To assess its potential impact, we applied a conservative worst-case period-based  
260 correction, accounting for the number of airgun shots that could have obscured detections. Even  
261 under this extreme assumption, detections during shooting periods remained substantially lower  
262 than during quiet periods, with minimum guaranteed reductions of 45.1% (OBH56) and maximum  
263 of 69.6% (OBS18) (Table 1). These results indicate that observed decreases in fin whale vocal  
264 activity are not solely an artefact of masking. The estimates likely overstate the true effect of  
265 masking, as it is improbable that every airgun shot obscured a vocalisation. Incorporating this  
266 conservative analysis provides robustness to our findings and reinforces confidence that seismic  
267 activity reduces calling behaviour, rather than merely obscuring it.

268  
269 The CNN demonstrated strong discrimination between fin whale pulses and background noise,  
270 including airgun pulses (Fig. 3). To account for potential limits in detection, we incorporated error  
271 margins from the CNN's false positive and false negative rates into the detection counts and  
272 corresponding percentage reductions. Even with these adjustments and the worst-case masking  
273 correction, reductions in detected pulses remained substantial, thus, reinforcing that the observed  
274 declines reflect genuine behavioural responses rather than artefacts of detection or masking  
275 limitations (also see Supplementary Fig. S5).

276  
277 A primary limitation of this study is the absence of baseline data prior to the initiation of airgun  
278 shooting. Without pre-shooting measurements, it is not possible to determine the “normal”  
279 vocalisation activity or natural variability of the fin whales in the absence of anthropogenic  
280 disturbance. Given the clear impact of seismic surveys on fin whale vocalisations observed here,  
281 future survey designs could incorporate a secondary aim of assessing these effects. Deploying  
282 passive acoustic recorders, perhaps from a different vessel, for a period before the start of  
283 shooting would provide a baseline of typical calling behaviour, allowing a more robust assessment  
284 of how survey activity alters vocal activity. Additional limitations arise from the spatial  
285 configuration of the array and the vessel track. While multiple instruments were deployed across  
286 the survey area, behavioural responses may have varied spatially depending on proximity to the  
287 airgun source as well as habitat features [19,22]. Future work could expand analysis across all of  
288 the instruments recovered with useful data to enable quantification of response variability across  
289 the population. Finally, combining visual or tagging data alongside acoustic monitoring could help  
290 distinguish between reduced vocalisation versus actual displacement from the area [19,42].

291  
292 The consistent and substantial reduction in pulse-positive frames across all instruments,  
293 averaging 70.4% during active airgun shooting and 52.0% when accounting for a worst-case  
294 masking scenario, underscores the significant influence of anthropogenic noise on fin whale  
295 vocalising behaviour. These results suggest that seismic survey activity can induce an immediate  
296 response and may create progressive behavioural responses, likely reflecting a combination of  
297 decreased calling and temporary avoidance of noisy areas. Given the widespread use of seismic  
298 surveys and their capacity to propagate sound over long distances [11–13], these findings raise  
299 concerns about potential long-term impacts on fin whale communication, energy expenditure, and  
300 habitat use [38]. From a conservation and management perspective, these results highlight the  
301 necessity of considering the timing, intensity, and spatial coverage of seismic surveys in areas  
302 important for baleen whales, especially given that the reach of airgun noise extends far beyond the

303 immediate survey zone [12]. Mitigation measures, such as exclusion zones, or seasonal  
304 restrictions, may help reduce behavioural disruptions [18,22]. Alternative source technologies  
305 could also provide a quieter approach to seismic exploration. For example, marine vibroseis  
306 systems use a vibrating plate or shell to generate longer-duration, lower-amplitude signals that are  
307 variable in frequency to deliver the acoustic energy required for seismic imaging, and these are  
308 considered a less environmentally disruptive alternative to the impulsive signals of conventional  
309 airgun sources [43]. More broadly, this study highlights the importance of integrating acoustic  
310 monitoring into activities that produce anthropogenic noise to better understand the impacts of  
311 permitted sound-generating operations on marine mammals and inform responsible management  
312 of such activities.  
313

## 314 **Methods**

315

### 316 **OBH and OBS acoustic data.**

317 The acoustic data used in this study were collected during the Galicia3D seismic survey in 2013  
318 and are openly available through PANGAEA [44]. A total of 72 ocean-bottom instruments were  
319 deployed to continuously record the ocean's soundscape throughout the survey period,  
320 comprising of systems from the British Ocean Bottom Instrumentation Consortium (OBIC) [45] and  
321 GEOMAR (Papenberg and Klaeschen, 2014).  
322

323 To assess the impact of the survey on whale vocalisations, it was not necessary to analyse the  
324 whole dataset, which would have introduced significant redundancy since the same sets of  
325 vocalisations were detected on multiple instruments [23]. We analysed data from two ocean-  
326 bottom seismometer (OBS) instruments (OBS18 and OBS46) that recorded at 250 Hz sample rate  
327 and one ocean-bottom hydrophone (OBH) instrument (OBH56) that recorded at 200 Hz (Fig. 2),  
328 selected to provide both broad spatial coverage and variation in recording systems for training and  
329 evaluating the CNN. These instruments had a spacing of 30-40 km and are likely to have recorded  
330 independent sets of vocalisations. Data were taken from the hydrophones of OBS18 and OBH56  
331 and the vertical geophone of OBS46. Further details of these recording instruments are outlined in  
332 supplementary material (Supplementary Table S1).  
333

334 The data analysed for this study spanned from 5<sup>th</sup> June (Julian Day (JD)156) to 6<sup>th</sup> August (JD 218).  
335 This timeframe included two periods of active seismic shooting operations (Shooting 1: JD156-173;  
336 Shooting 2: JD 197-212) and two quiet periods with no airgun activity (Quiet 1: JD 174-196 during  
337 vessel repairs; Quiet 2: JD 213-218 after survey completion). For the purposes of analysis, any day  
338 during which airgun shooting occurred at any time was classified as a shooting day. These four  
339 periods provided the framework for evaluating fin whale acoustic activity in relation to shooting  
340 operations.  
341

### 342 **Manually labelled training dataset.**

343 To train and assess the performance of the CNN algorithm for its automated detection of the fin  
344 whale pulses, a manually labelled subset of data served as training and validation datasets. This  
345 manual signal labelling process was carried out in HydroSeek, a custom signal labelling  
346 application for passive acoustic monitoring purposes [47]. This software converted the acoustic  
347 data into 30-second spectrogram frames allowing the user to manually label signals present in the  
348 time-frequency domain. Each processed frame was assigned a binary score, corresponding to the  
349 presence or absence of signals. Within HydroSeek, label classifications were made using three

350 spectrogram views of varying input parameters for multi-scale analysis of the acoustic signals  
351 present. The three spectrogram configurations were: FFT 256 points, 78 % overlap, 1-99 Hz; FFT  
352 256 points, 86 % overlap, 1-80 Hz; and FFT 300 points, 83 % overlap, 18-30 Hz. The dynamic range  
353 parameter was set between -30 and 10 dB, but due to differences in sample rates, these dynamic  
354 range values required occasional minor adjustment for the clearest spectrograms.

355 To minimise subjectivity and maintain consistency in the manual labelling process, clear criteria  
356 were established for each label classification (Supplementary Table S2). Despite these criteria,  
357 some subjectivity is unavoidable [48]. To maintain consistency, the labelling process was limited  
358 to two trained analysts. A shared blind review of 10% of the dataset indicated a 3.75% inter-analyst  
359 disagreement rate. In all cases, the annotation of the original labelling was retained for analyses.

360  
361 During this process, 30 minutes of acoustic data were manually labelled in 2-hour intervals for the  
362 for the first complete day of the Shooting 1 (JD156), Quiet 1 (JD175) and Shooting 2 (JD198) periods  
363 from OBS18, OBS46 and OBH56. The range of instruments and days ensured the dataset contained  
364 a comprehensive repertoire of the signals present (Table S3).  
365

## 366 **Machine learning**

367  
368 **Model structure.** To detect fin whale 20 Hz calls, we used a Convolutional Neural Network (CNN)  
369 classifier by adapting an established image-recognition architecture and training it on spectrogram  
370 representations of the acoustic data. We implemented the CNN using the EfficientNet [49] model  
371 family's baseline architecture, EfficientNetB0, pre-trained on the ImageNet dataset [50]. Global  
372 average pooling [51] was applied to the output of the EfficientNetB0 feature extractor before  
373 feeding it into the classification part. This classification part consists of two fully connected layers  
374 with 512 and 256 neurons, each with a rectified linear unit activation function and followed by  
375 dropout layers [52] with the rates of 0.5 and 0.2, respectively. The final output layer uses a softmax  
376 activation with two units for binary classification (i.e., presence vs. absence of fin whale calls). The  
377 dense layers in the classification part were initialized with Glorot uniform [53] for the weights and  
378 zeros for the bias vectors and the EfficientNetB0 feature extractor with the pre-trained parameters.  
379 The network was trained with full fine-tuning, meaning that the entire network was trained on the  
380 target dataset, after parameter initialisation. The same network architecture has been employed in  
381 several underwater acoustics classification studies [54–56].  
382

383 **Input representation.** Training inputs consisted of 30 s segments, downsampled to 200Hz and  
384 transformed into spectrograms using FFT 256 points and 75% overlap with frequency band-limiting  
385 applied between 4 and 80 Hz. Per-channel energy normalisation (PCEN) was used to enhance  
386 signal detection under noisy conditions by suppressing background energy and emphasising  
387 transient low-frequency calls [57]. PCEN hyperparameters were set following previous work on  
388 tuning them empirically, which demonstrated their efficacy compared with alternative input  
389 representations [54]. To avoid bias from temporal sequence, each spectrogram was treated  
390 independently of its chronological context. The final input dimensions were 97 time frames by 90  
391 frequency bins with red-green-blue conversion applied, 97 x 90 x 3.  
392

393 **Training dataset.** The model was trained using manually labelled data from OBS18, OBS46, and  
394 OBH56 across JDs 156, 175, and 198, covering both survey and quiet periods. Positive samples  
395 contained  $\geq 1$  fin whale pulses, while negative samples were drawn from a range of acoustic

396 backgrounds, including airgun pulses and ambient noise. To minimise class imbalance,  
397 approximately equal numbers of positive and negative frames were used for training (3,099  
398 positive, 2,892 negative frames), totalling approximately 50 hours of labelled data. A full  
399 breakdown of the training datasets is provided in the supplementary material (Supplementary  
400 Table S3).

401  
402 **Model training.** Training was conducted by using a categorical cross-entropy loss function. We use  
403 an adaptive moment estimation optimizer with exponential learning rate decay [58]. The  
404 exponential decay schedule used an initial rate of 0.001 and a decay rate of 0.75 at every 90 steps  
405 with a staircase function.

406  
407 **Validation.** An unseen dataset of 30 s spectrogram samples containing a similar number of  
408 positive (n = 639) and negative (n = 572) frames was used for validation (Supplementary Table S3).  
409 Model performance was quantified using multiple metrics. Overall accuracy provided a baseline  
410 measure of correct classifications. Precision was defined as the proportion of CNN detections that  
411 were true positives, whereas recall represented the proportion of true fin whale pulses correctly  
412 detected by the model. The F1 score, the harmonic mean of precision and recall, was included as a  
413 summary metric that balances the trade-off between false positives and false negatives. In  
414 addition, Receiver Operating Characteristic (ROC) analysis was performed to evaluate model  
415 performance across all decision thresholds, with the Area Under the Curve (AUC) used to  
416 summarise the model's overall discriminative ability independent of the choice of threshold.

417  
418 **Large-scale application.** The final CNN was applied to the full acoustic dataset from JD 156–218,  
419 spanning 63 days of continuous recordings across all three instruments. Each 30 s segment was  
420 independently classified, producing >180,000 outputs per instrument. A probability threshold of  
421 0.5 was used to convert model outputs into binary classifications. This optimal ROC-derived  
422 threshold value was 0.51, and we therefore used the rounded value of 0.5, consistent with  
423 standard practice in binary classification [59]. The same threshold was used for evaluating the  
424 model and for generating daily detection counts. Processing was carried out on Google Colab  
425 using a T4 High-RAM runtime. In total 25 days contained corrupt data, likely due to data conversion  
426 or transcription issues. These data were excluded from further analysis.

## 427 **Statistical analysis.**

428 Detection count data generated by the CNN were analysed to evaluate fin whale acoustic activity in  
429 relation to seismic shooting operations. Each positive 'detection' corresponded to a 30-second  
430 frame containing at least one fin whale pulse. Daily detection counts were summarised separately  
431 for each instrument and partitioned into the four survey periods to observe phase changes.

432 CNN-derived detections were first aggregated to hourly means to reduce fine-scale variability and  
433 to standardise comparisons across the full survey period. Periods with corrupted data files were  
434 treated as missing (NA) and excluded from all analyses. To assess differences in hourly mean  
435 detection rates among survey periods, we fitted a negative binomial Generalised Linear Mixed-  
436 effect Model (GLMM) using the glmmTMB package in R [60]. The model specified Period as a fixed  
437 effect and incorporated two random-effect terms: (1| Instrument) to account for variation among  
438 recorders, and (1| Instrument: JD) to account for day-level temporal autocorrelation within each  
439 instrument. A negative binomial error structure was selected after the initial Poisson models  
440 showed substantial overdispersion. Model assumptions were evaluated using the DHARMA

441 package [61], including dispersion tests, zero-inflation tests and simulation-based residual  
442 diagnostics.

443 Post hoc comparisons between survey periods were conducted using Estimated Marginal Means  
444 (EMMs) using the emmeans package in R [62]. Pairwise contrasts were adjusted using Tukey  
445 correction, and estimates were reported on the response scale (mean detections per hour) with  
446 95% confidence intervals. This approach allowed direct comparison of call occurrence rates  
447 between active shooting and quiet conditions for each instrument and location independently.  
448 Model residuals were visually inspected to confirm distributional assumptions and goodness of fit.  
449 All analyses were conducted in R (v4.4.1)[63]

#### 450 **Maximum masking estimation.**

451 To assess the inherent limitation presented by the masking of fin whale pulses by airgun shot  
452 signals within the data, we estimated the fraction of spectrogram frames likely to be obscured  
453 during active shooting. Airgun pulses occurred on average every 16 s, with each pulse producing a  
454 masking duration of 5.5 s (Mean = 5.50, range = 2.73–7.73 seconds). Pulse characteristics were  
455 obtained through manual review of 20 representative pulses sampled across both legs (Fig. 1b).  
456

457 We used the recorded number of airgun shots per shooting period to estimate actual masked time.  
458 For each period we multiplied the shot count by the mean masking duration (5.5 s) to obtain total  
459 masked seconds and then converted to hours. For Shooting 1 (n = 66,898 shots), this produced 102  
460 hrs masked out of 432 hrs of surveying. For Shooting 2 (n = 84,417 shots), this produced 129 hrs  
461 masked out of 384 hrs.  
462

463 To incorporate the effect of this masked time into the detection analysis, we derived a single  
464 average masking correction factor across both shooting periods. The proportion of shooting time  
465 that was masked was calculated (23.5% in Shooting 1; 33.6% in Shooting 2), producing an overall  
466 mean masked fraction of 28.7%. Assuming detections are equally likely to occur within masked  
467 and unmasked intervals, the expected reduction in detectability is proportional to the masked  
468 time. Therefore, we applied a correction factor, calculated as  $1 / (1 - 0.287) = 1.40$ , to the observed  
469 mean hourly detections during shooting to account for calls potentially lost due to masking. This  
470 approach allowed us to assess the robustness of observed decreases in calling during active  
471 airgun activity.  
472

473  
474  
475

#### 476 **Data availability**

477 The acoustic data used in this study were collected during the Galicia3D seismic survey in 2013 and are  
478 openly available via PANGAEA (Bayrakci et al., 2022): <https://doi.org/10.1594/PANGAEA.940656>. All code  
479 used for data processing, statistical analysis, and the CNN model is available from the corresponding author  
480 upon reasonable request.  
481

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488

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493

494

#### 495 **Author contributions**

496 TM conceived the study and supervised the project. EE and AF carried out the analyses and wrote  
497 the manuscript. EE, AF and TM prepared the figures. AO contributed to data analysis and model  
498 development. All authors discussed the results and reviewed the manuscript. EE and AF  
499 contributed equally to this work.

500

#### 501 **Competing interests**

502 The author(s) declare no competing interests.

#### 503 **References**

- 504 1. Erbe, C. *et al.* Sources of Underwater Noise. in *Marine Mammal Acoustics in a Noisy*  
505 *Ocean* (eds Erbe, C., Houser, D., Bowles, A. & Porter, M. B.) 85–178 (Springer Nature  
506 Switzerland, Cham, 2025). doi:10.1007/978-3-031-77022-7\_2.
- 507 2. Hildebrand, J. Anthropogenic and natural sources of ambient noise in the ocean. *Mar.*  
508 *Ecol. Prog. Ser.* **395**, 5–20 (2009).
- 509 3. Studds, G. E. & Wright, A. J. A Brief Review of Anthropogenic Sound in the Oceans. *Int. J.*  
510 *Comp. Psychol.* **20**, (2007).
- 511 4. Duarte, C. M. *et al.* The soundscape of the Anthropocene ocean. *Science* **371**,  
512 eaba4658 (2021).
- 513 5. Clark, C. *et al.* Acoustic masking in marine ecosystems: intuitions, analysis, and  
514 implication. *Mar. Ecol. Prog. Ser.* **395**, 201–222 (2009).
- 515 6. Erbe, C. *et al.* The Effects of Ship Noise on Marine Mammals—A Review. *Front. Mar.*  
516 *Sci.* **6**, 606 (2019).

- 517 7. Nowacek, D. P., Christiansen, F., Bejder, L., Goldbogen, J. A. & Friedlaender, A. S.  
518 Studying cetacean behaviour: new technological approaches and conservation  
519 applications. *Anim. Behav.* **120**, 235–244 (2016).
- 520 8. Richardson, W. J., Miller, G. W. & Greene, C. R. Displacement of migrating bowhead  
521 whales by sounds from seismic surveys in shallow waters of the Beaufort Sea. *J.*  
522 *Acoust. Soc. Am.* **106**, 2281–2281 (1999).
- 523 9. Rolland, R. M. *et al.* Evidence that ship noise increases stress in right whales. *Proc. R.*  
524 *Soc. B Biol. Sci.* **279**, 2363–2368 (2012).
- 525 10. Carroll, A. G., Przeslawski, R., Duncan, A., Gunning, M. & Bruce, B. A critical review of  
526 the potential impacts of marine seismic surveys on fish & invertebrates. *Mar. Pollut.*  
527 *Bull.* **114**, 9–24 (2017).
- 528 11. Gillespie, A. Noise Pollution, the Oceans, and the Limits of International Law. *Yearb.*  
529 *Int. Environ. Law* **21**, 114–139 (2010).
- 530 12. Nieuwkirk, S. L., Stafford, K. M., Mellinger, D. K., Dziak, R. P. & Fox, C. G. Low-frequency  
531 whale and seismic airgun sounds recorded in the mid-Atlantic Ocean. *J. Acoust. Soc.*  
532 *Am.* **115**, 1832–1843 (2004).
- 533 13. Kavanagh, A. S., Nykänen, M., Hunt, W., Richardson, N. & Jessopp, M. J. Seismic  
534 surveys reduce cetacean sightings across a large marine ecosystem. *Sci. Rep.* **9**, 19164  
535 (2019).
- 536 14. Castellote, M., Clark, C. W. & Lammers, M. O. Acoustic and behavioural changes by fin  
537 whales (*Balaenoptera physalus*) in response to shipping and airgun noise. *Biol.*  
538 *Conserv.* **147**, 115–122 (2012).

- 539 15. Di Iorio, L. & Clark, C. W. Exposure to seismic survey alters blue whale acoustic  
540 communication. *Biol. Lett.* **6**, 51–54 (2010).
- 541 16. Gordon, J. *et al.* A Review of the Effects of Seismic Surveys on Marine Mammals. *Mar.*  
542 *Technol. Soc. J.* **37**, 16–34 (2003).
- 543 17. Greene, C. R. & Richardson, W. J. Characteristics of marine seismic survey sounds in  
544 the Beaufort Sea. *J. Acoust. Soc. Am.* **83**, 2246–2254 (1988).
- 545 18. Gailey, G. *et al.* Western gray whale behavioral response to seismic surveys during their  
546 foraging season. *Environ. Monit. Assess.* **194**, 740 (2022).
- 547 19. Blackwell, S. B. *et al.* Effects of airgun sounds on bowhead whale calling rates in the  
548 Alaskan Beaufort Sea. *Mar. Mammal Sci.* **29**, (2013).
- 549 20. Cerchio, S., Strindberg, S., Collins, T., Bennett, C. & Rosenbaum, H. Seismic Surveys  
550 Negatively Affect Humpback Whale Singing Activity off Northern Angola. *PLoS ONE* **9**,  
551 e86464 (2014).
- 552 21. Maciel, I. *et al.* Cetaceans Change Their Acoustic Behavior During the Airgun Noise of  
553 Seismic Surveys. *J. Mar. Sci. Eng.* **14**, 181 (2026).
- 554 22. Dunlop, R. A. *et al.* The behavioural response of migrating humpback whales to a full  
555 seismic airgun array. *Proc. R. Soc. B Biol. Sci.* **284**, 20171901 (2017).
- 556 23. Fisher, J., Minshull, T. A., White, P. R., Tian, B. & Bayrakci, G. Tracking and Monitoring  
557 Fin Whales Offshore Northwest Spain Using Passive Acoustic Methods. in *Geophysical*  
558 *Monograph Series* (eds Bayrakci, G. & Klingelhoefer, F.) 209–221 (Wiley, 2024).  
559 doi:10.1002/9781119750925.ch12.

- 560 24. Cooke, J. G. *Balaenoptera physalus*: The IUCN Red List of Threatened Species 2018:  
561 e.T2478A50349982. IUCN <https://doi.org/10.2305/iucn.uk.2018->  
562 [2.rlts.t2478a50349982.en](https://doi.org/10.2305/iucn.uk.2018-2.rlts.t2478a50349982.en) (2018).
- 563 25. Watkins, W. A., Tyack, P., Moore, K. E. & Bird, J. E. The 20-Hz signals of finback whales (  
564 *Balaenoptera physalus*). *J. Acoust. Soc. Am.* **82**, 1901–1912 (1987).
- 565 26. Guazzo, R. A., Stevenson, D. L., Edell, M. K., Gagnon, G. J. & Helble, T. A. A decade of  
566 change and stability for fin whale song in the North Atlantic. *Front. Mar. Sci.* **11**,  
567 1278068 (2024).
- 568 27. Croll, D. A. *et al.* Only male fin whales sing loud songs. *Nature* **417**, 809–809 (2002).
- 569 28. Aulich, M. G. *et al.* Diel patterns of fin whale 20 Hz acoustic presence in Eastern  
570 Antarctic waters. *R. Soc. Open Sci.* **10**, (2023).
- 571 29. McDonald, M. A., Hildebrand, J. A. & Webb, S. C. Blue and fin whales observed on a  
572 seafloor array in the Northeast Pacific. *J. Acoust. Soc. Am.* **98**, 712–721 (1995).
- 573 30. Watkins, W. A. Activities and underwater sounds of fin whales. *Sci Rep Wales Res.*  
574 **Inst.**, (1981).
- 575 31. Johnson, C. M. *et al.* *Protecting Blue Corridors - Challenges and Solutions for Migratory*  
576 *Whales Navigating National and International Seas.*  
577 <https://zenodo.org/record/6196130> (2022) doi:10.5281/ZENODO.6196130.
- 578 32. Bayrakci, G. *et al.* Fault-controlled hydration of the upper mantle during continental  
579 rifting. *Nat. Geosci.* **9**, 384–388 (2016).

- 580 33. Davy, R. G. *et al.* Continental hyperextension, mantle exhumation, and thin oceanic  
581 crust at the continent-ocean transition, West Iberia: New insights from wide-angle  
582 seismic. *J. Geophys. Res. Solid Earth* **121**, 3177–3199 (2016).
- 583 34. Williams, R. *et al.* Impacts of anthropogenic noise on marine life: Publication patterns,  
584 new discoveries, and future directions in research and management. *Ocean Coast.*  
585 *Manag.* **115**, 17–24 (2015).
- 586 35. Heide-Jørgensen, M. P. *et al.* Behavioral Response Study on Seismic Airgun and Vessel  
587 Exposures in Narwhals. *Front. Mar. Sci.* **8**, 658173 (2021).
- 588 36. Watkins, W. A., Tyack, P., Moore, K. E. & Bird, J. E. The 20-Hz signals of finback whales  
589 (*Balaenoptera physalus*). *J. Acoust. Soc. Am.* **82**, 1901–1912 (1987).
- 590 37. Sawyer, D. S., Morgan, J. K. & Shillington, D. J. *MGL1307 Cruise Report.* (2013).
- 591 38. Senigaglia, V., Houser, D., Jolliffe, C. & Erbe, C. Biological Significance of Responses to  
592 Noise. in *Marine Mammal Acoustics in a Noisy Ocean* (eds Erbe, C., Houser, D.,  
593 Bowles, A. & Porter, M. B.) 699–729 (Springer Nature Switzerland, Cham, 2025).  
594 doi:10.1007/978-3-031-77022-7\_11.
- 595 39. Wright, A. J. *et al.* Do Marine Mammals Experience Stress Related to Anthropogenic  
596 Noise? *Int. J. Comp. Psychol.* **20**, (2007).
- 597 40. Joy, R. *et al.* A fine-scale marine mammal movement model for assessing long-term  
598 aggregate noise exposure. *Ecol. Model.* **464**, 109798 (2022).
- 599 41. Caruso, F. *et al.* Detection and tracking of fin whales during seismic exploration in the  
600 Gulf of California. in 070021 (Dublin, Ireland, 2016). doi:10.1121/2.0000424.

- 601 42. Madsen, P. T. *et al.* Quantitative measures of air-gun pulses recorded on sperm whales  
602 (*Physeter macrocephalus*) using acoustic tags during controlled exposure  
603 experiments. *J. Acoust. Soc. Am.* **120**, 2366–2379 (2006).
- 604 43. Duncan, A. J., Weilgart, L. S., Leaper, R., Jasny, M. & Livermore, S. A modelling  
605 comparison between received sound levels produced by a marine Vibroseis array and  
606 those from an airgun array for some typical seismic survey scenarios. *Mar. Pollut. Bull.*  
607 **119**, 277–288 (2017).
- 608 44. Bayrakci, G. *et al.* Ocean bottom seismic data from the continent-ocean transition in  
609 the Deep Galicia Margin, offshore west Iberia. 3 datasets Preprint at  
610 <https://doi.org/10.1594/PANGAEA.940656> (2022).
- 611 45. Minshull, T. A., Sinha, M. C. & Peirce, C. Multi-disciplinary, sub-seabed geophysical  
612 imaging - a new pool of 28 seafloor instruments in use by the United Kingdom Ocean  
613 Bottom Instrument Consortium. *Sea Technol.* **46**, 27–31 (2005).
- 614 46. Papenberg, C. & Klaeschen, D. *RV POSEIDON Fahrtbericht / Cruise Report POS 453 &*  
615 *458 'COMM3D': Crustal Structure and Ocean Mixing Observed with 3D Seismic*  
616 *Measurements, 20.05. - 12.06.2013 (POS453) Galway, Ireland - Vigo, Portugal ; 24.09. -*  
617 *17.10.2013 (POS458) Vigo, Portugal - Vigo, Portugal.*  
618 <https://oceanrep.geomar.de/id/eprint/25620/> (2014).
- 619 47. White, E. Hydroseek. (2025).
- 620 48. Leroy, E. C., Thomisch, K., Royer, J.-Y., Boebel, O. & Van Opzeeland, I. On the reliability  
621 of acoustic annotations and automatic detections of Antarctic blue whale calls under  
622 different acoustic conditions. *J. Acoust. Soc. Am.* **144**, 740–754 (2018).

- 623 49. Tan, M. & Le, Q. V. EfficientNet: Rethinking Model Scaling for Convolutional Neural  
624 Networks. <https://doi.org/10.48550/ARXIV.1905.11946> (2019)  
625 doi:10.48550/ARXIV.1905.11946.
- 626 50. Deng, J. *et al.* ImageNet: A large-scale hierarchical image database. in *2009 IEEE*  
627 *Conference on Computer Vision and Pattern Recognition* 248–255 (IEEE, Miami, FL,  
628 2009). doi:10.1109/CVPR.2009.5206848.
- 629 51. Lin, M., Chen, Q. & Yan, S. Network In Network. Preprint at  
630 <https://doi.org/10.48550/ARXIV.1312.4400> (2013).
- 631 52. Srivastava, N., Hinton, G., Krizhevsky, A., Sutskever, I. & Salakhutdinov, R. Dropout: a  
632 simple way to prevent neural networks from overfitting. *J. Mach. Learn. Res.* **15**, 1929–  
633 1958 (2014).
- 634 53. Glorot, X. & Bengio, Y. Understanding the difficulty of training deep feedforward neural  
635 networks. in *Proceedings of the thirteenth international conference on artificial*  
636 *intelligence and statistics* (eds Teh, Y. W. & Titterton, M.) vol. 9 249–256 (PMLR, Chia  
637 Laguna Resort, Sardinia, Italy, 2010).
- 638 54. Olcay, A. *et al.* Sounds of the deep: How input representation, model choice, and  
639 dataset size influence underwater sound classification performance. *J. Acoust. Soc.*  
640 *Am.* **157**, 3017–3032 (2025).
- 641 55. White, E. L., Klinck, H., Bull, J. M., White, P. R. & Risch, D. One size fits all? Adaptation  
642 of trained CNNs to new marine acoustic environments. *Ecol. Inform.* **78**, 102363  
643 (2023).

- 644 56. White, E. L. *et al.* More than a whistle: Automated detection of marine sound sources  
645 with a convolutional neural network. *Front. Mar. Sci.* **9**, 879145 (2022).
- 646 57. Lostanlen, V. *et al.* Per-Channel Energy Normalization: Why and How. *IEEE Signal*  
647 *Process. Lett.* **26**, 39–43 (2019).
- 648 58. You, K., Long, M., Wang, J. & Jordan, M. I. How Does Learning Rate Decay Help Modern  
649 Neural Networks? Preprint at <https://doi.org/10.48550/ARXIV.1908.01878> (2019).
- 650 59. Freeman, E. A. & Moisen, G. G. A comparison of the performance of threshold criteria  
651 for binary classification in terms of predicted prevalence and kappa. *Ecol. Model.* **217**,  
652 48–58 (2008).
- 653 60. Brooks, M., E. *et al.* glmmTMB Balances Speed and Flexibility Among Packages for  
654 Zero-inflated Generalized Linear Mixed Modeling. *R J.* **9**, 378 (2017).
- 655 61. Hartig, F. *DHARMA: Residual Diagnostics for Hierarchical (Multi-Level / Mixed)*  
656 *Regression Models*. <https://CRAN.R-project.org/package=DHARMA> (2024).
- 657 62. Lenth, R. V. emmeans: Estimated Marginal Means, aka Least-Squares Means. 1.11.2-8  
658 <https://doi.org/10.32614/CRAN.package.emmeans> (2017).
- 659 63. R Core Team. R: A Language and Environment for Statistical Computing. R Foundation  
660 for Statistical Computing (2024).

661

## 662 **Figure legends**

663

664 **Figure 1.** Spectrograms of fin whale pulses recorded by ocean-bottom seismometer (OBS)  
665 instruments. (a) Example of the low frequency, downsweep structure of the fin whale pulses from  
666 Julian Day 175, 2013 (OBS18). (b) A segment from Julian Day 198, 2013 (OBS46), recorded during  
667 active shooting period, showing fin whale calls at approximately 17 and 32 seconds occurring  
668 between airgun pulses. The black brackets above the plot indicate the portions used to estimate  
669 the average masking duration of individual airgun pulses (mean = 5.5 s, range = 2.7–7.7 s;  
670 calculated from 20 representative pulses across both survey legs). The spectrograms were

671 generated using a 256-point FFT, 80% overlap, a Hanning window, and a frequency range of 5–45  
672 Hz.

673 **Figure 2.** Bathymetry of survey area offshore Galicia Bank, with the grey colour representing areas  
674 without swath bathymetric coverage. White symbols mark seafloor instruments deployed, with  
675 circles marking ocean-bottom seismometers (OBSs) and squares marking ocean-bottom  
676 hydrophones (OBHs). Red symbols represent instruments from which data for this study were  
677 obtained. White box marks area of 3D seismic survey. Sail lines were parallel to the long side of the  
678 box and at 400 m spacing (see Fig. S2 of Bayrakci et al., 2016). Inset shows regional location of map  
679 area offshore Iberia (small white box).

680  
681 **Figure 3** (a) Confusion matrix summarising the model’s detections for 30-second periods from the  
682 validation dataset. Numbers represent the count of frames correctly or incorrectly classified,  
683 predicted values shown on the y-axis and actual values on the x-axis. (b) Receiver Operating  
684 Characteristic (ROC) curve showing the trade-off between true positive rate and false positive rate  
685 for the classification model. The Area Under the Curve (AUC) quantifies overall model performance,  
686 with values closer to 1 indicating better discrimination.

687 **Table 1.** Mean hourly detections of fin whale pulse-positive frames per hour during quiet and  
688 shooting periods, with observed and worst-case (frame-based) masking corrections applied. Quiet  
689 means were averaged across Quiet 1 and Quiet 2; shooting means were averaged across Shooting  
690 1 and Shooting 2. Corrected shooting means (positive frames  $hr^{-1}$ ) were calculated using the  
691 correction factor, assuming that every airgun shot overlapped a fin whale pulse. Percentage drops  
692 are relative to quiet means.

693  
694 **Figure 4.** Positive frame counts per hour across different periods for each instrument. Each box  
695 represents the interquartile range, the horizontal line marks the median, whiskers extend to 1.5x  
696 the interquartile range, and the black circles represent outliers. Red boxes represent Shooting  
697 periods, while grey boxes represent Quiet periods.

698  
699 **Figure 5.** Daily total positive frame counts (bars) and Locally Estimated Scatterplot Smoothing  
700 (LOESS) smoothed trend lines for each instrument across Julian Days (JD). Shaded backgrounds  
701 indicate Quiet and Shooting (red) periods. The smoothed line was generated using a LOESS  
702 regression (span = 0.3) and is plotted with an error ribbon (semi-transparent band) that accounts  
703 for detection uncertainty, calculated by scaling the smoothed counts with instrument false positive  
704 (FPR = 0.158) and false negative (FNR = 0.132) rates. Days with corrupt data (grey vertical bands)  
705 were excluded from the analysis and treated as NAs.