COVID-related anthropause highlights the impact of marine traffic but not of tourism on breeding little penguins

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ABSTRACT

The COVID-19 pandemic and its lock-down measures have resulted in periods of reduced human activity, known as anthropause. While this period was expected to be favorable for the marine ecosystem, due to a probable reduction of pollution, shipping traffic, industrial activity and fishing pressure, negative counterparts such as reduced fisheries surveillance could counterbalance these positive effects. Simultaneously, on-land pressure due to human disturbance and tourism should have drastically decreased, potentially benefiting land-breeding marine animals such as seabirds. We analyzed 11 breeding seasons of data on several biological parameters of little penguins from a popular tourist attraction at Phillip Island, Australia. We investigated the impact of anthropogenic activities on penguin behavior during the breeding season measured by (1) distribution at sea, (2) colony attendance, (3) isotopic niche (4) chick meal mass, and (5) offspring investment against shipping traffic and number of tourists. The 2020 lock-downs resulted in a near absence of tourists visiting the Penguin Parade®, which was otherwise visited by 800,000+ visitors on average per breeding season. However, our long-term analysis showed no effect of the presence of visitors on little penguins' activities. Surprisingly, the anthropause did not trigger any changes in maritime traffic intensity and distribution in the region. We found inter- and intra-annual variations for most parameters, we detected a negative effect of marine traffic on the foraging efficiency. Our results suggest that environmental variations have a greater influence on the breeding behavior of little penguins compared to short-term anthropause events. Our long-term dataset was key to test whether changes in anthropogenic activities affected the wildlife during the COVID-19 pandemic.

1. Introduction

With the development of human activities, ecosystems can no longer be considered as undisturbed and independent entities (Mace, 2014), leading to the concept of socio-ecological systems (Everard, 2020; Wei et al., 2018). Because of the numerous interactions at stake, socio-ecological ecosystems are often complex to analyze (Sugihara et al., 2012). The quasi-continuous presence of humans in most, if not all, ecosystems make it challenging to understand the full impact of anthropogenic activities on the environment.

In 2020, the COVID-19 pandemic led to periods of lock-downs that resulted in a major reduction of human activities and movement at both local and global level, a period coined as the "anthropause" (Lamers and Student, 2021; Rutz et al., 2020). The anthropause created an opportunity to quantify the impact of human activities on wildlife. To date, studies found both negative and positive effects of this anthropause on wildlife, through for example, increase of predators presence and disturbance on an iconic seabird colony in the Baltic Sea (Hentati-Sundberg et al., 2021), as well as increased species richness in less-disturbed areas (Manenti et al., 2020). Lock-downs also led to increased illegal hunting and plastic pollution, and reduced conservation efforts with negative effects on wildlife (Bates et al., 2021; Kadykalo et al., 2022). In a comparative study, Bates et al. (Bates et al., 2021) showed that despite the decrease in humans' movement, or industrial activities, the median responses of wildlife to anthropause were centered on 0, because either positive and negative effects balanced

themselves, or for numerous species, no effects were observed.

Moreover, it can be misleading to consider that the anthropause is a phenomenon homogeneously distributed across the globe. The decrease in human activities was not equal across the planet (Bates et al., 2021), calling for more studies across ecosystems.

However, it can be complex to study the dynamics of entire ecosystems, specifically within the context of the COVID lock-downs, considering the difficulties to carry on with species and habitat monitoring activities during these periods. Monitoring "sentinel species" helps tackling this issue. Sentinel species integrate changes happening across ecosystems' levels (Durant et al., 2009), integrate broader processes into rapidly interpretable metrics, are simpler to study, can respond rapidly to environmental changes and cover a large spatial scale (Bost et al., 2008; Durant et al., 2009; Hazen et al., 2019; Siddig et al., 2016). Therefore, long-term dataset on marine predators, especially seabirds, are often used as indicators of ecosystems' changes (Cairns, 1988; Furness and Camphuysen, 1997; Piatt et al., 2007b).

Data collection via continuous monitoring programs allows researchers to compare the pace of parameters responses to global changes and assess effects of human pressure on wild populations (Cairns, 1988; Durant et al., 2009; Einoder, 2009; Ramírez et al., 2017; Tucker et al., 2019, 2018). Techniques used vary depending on the research question and feasibility, comprising of visual observations, counts, nest monitoring, blood sampling and use of bio-logging techniques. In seabirds, chick growth, colony attendance, and individuals' activity budgets vary at different temporal scales and in relation to both environmental and human activities (Cairns, 1988). Depending on the specific effects of the COVID lock-downs and the relative short period these were put in place, some of these traits might show no responses to anthropogenic pressure (Cairns, 1988; Piatt et al., 2007a).

During the breeding season, seabirds are central place foragers exploiting food resources around their breeding colony to which they return due to reproductive requirements (e.g., egg incubation, chick provisioning), hence alternating between nest attendance and foraging trips (Einoder, 2009; Piatt et al., 2007b; Saraux et al., 2011b). Seabirds must cope with constraints of living in two different environments, feeding at sea and breeding on land, making them exposed and vulnerable to threats from both land and sea. The little penguin (*Eudyptula minor*) is the smallest penguin species endemic of Australia and New Zealand (BirdLife International, 2023). Phillip Island, Australia, holds one of the largest little penguin colonies in the world with a population estimated between 28,000 and 32,000 individuals (Sutherland and Dann, 2014). The colony located at the "Penguin Parade®" receives the visit of hundreds of thousands of tourists per breeding season, especially when little penguins return ashore at night (Dann and Chambers, 2013). The "Penguin Parade®" is a nature park, where considerable efforts have been made to manage tourism so as to avoid affecting breeding penguins. Yet, it has been running everyday since 1968, with an average of half a million visitors per year, making it difficult to test whether penguins are disturbed by the presence of humans. At sea, little penguins can also interact with maritime traffic such as commercial shipping, recreational or commercial fishing vessels (Cannell et al., 2020; Crawford et al., 2017). Land introduced predators and starvation are the major causes of little penguins' mortality, but collision with vessels were also reported (Cannell et al., 2020, 2016), even though their foraging range is small (around 30 km for single day trips but can be up to 214 km for multi days trips) (Collins et al., 1999; Poupart et al., 2017; Sánchez et al., 2018).

On land, tourism has been shown to affect various parameters of penguins' ecology such as stress level, reproductive output (Ellenberg et al., 2007) or behavior (Colombelli-Négrel and Katsis, 2021; Ellenberg et al., 2007; French et al., 2019). At sea, vessels can directly (Pichegru et al., 2022) or indirectly (Mattern et al., 2013) affect penguins' foraging through noise pollution and deterioration of the environment, respectively. During the COVID-19 pandemic, Australia underwent a series of rigid lock-downs, drastically reducing anthropogenic activities. During most of that period, the "Penguin Parade®" remained closed to the tourists, providing a good opportunity to understand if the anthropause affected ecology of little penguins. At a moment when ecotourism is often the key to the acceptance of stakeholders for conservation programs, this could provide important answers to the effect of highly-managed tourism.

We investigated whether the anthropause affected metrics linked to little penguin's behavior during the breeding season in 2020 (with lock-downs) by comparing against 10 breeding seasons of population monitoring and movement data (2010–2019) to 2020. The studied colony has been monitored for the past 23 breeding seasons using an automated penguin monitoring system (date, time and weight of penguins recorded when leaving and arriving to the colony), with daily count of penguins arrival at dusk, and with the use of bio-logging techniques (the latter since 2010) (Ramírez et al., 2015). We tested whether reduced anthropogenic activities influenced little penguins (1) at-sea activity by studying their at-sea distribution, overlap with marine traffic, isotopic diet (in terms of prey type and quantity), and (2) on-land activity by studying their colony attendance (departure and arrival time), and meal size given to their chicks. We considered the daily number of tourists at the Penguin Parade® as a proxy of land disturbance, and the number of vessels and their overlap with little penguins' foraging area at sea as a proxy of the at-sea disturbance. We hypothesized that when land disturbance was reduced during the anthropause, due to the absence of tourists in the parks, little penguins would change their colony attendance pattern by coming and leaving more synchronously, as they will not have to avoid tourist disturbance (Klomp and Wooller, 1991; Rodríguez et al., 2016). Moreover, if the anthropause reduced the at-sea disturbance, little penguins would display a higher foraging efficiency as the overall marine environment and its species will be less disturbed by the traffic, through reduction in noise pollution for instance (Pichegru et al., 2017, 2010).

2. Material and methods

2.1. Study site and long-term monitoring of foraging behavior

The study was conducted on the little penguin breeding colony at Penguin Parade®, Phillip Island, Australia (38°21′S, 145°09′E) from 2010 to 2020. The breeding season of little penguins occurs in the austral spring and summer, from September to December.

For the period of our study (11 breeding seasons, 2010–2020), penguins were captured from their nest boxes and equipped with GPS loggers (Axy-Trek, Italy, Mr. Lee, China) recording positions at 120 s interval for incubation and postguard trips and every 20 s for guard trips (Supplementary Table 1). Loggers were attached to their lower backs with Tesa ® tape (Wilson et al., 1997). After returning from their foraging trips, penguins were recaptured at the colony and the logger retrieved. Handling time was kept at <5 min. Details of the logger deployment are described in Pelletier et al. (2014), and Barreau et al. (2021). We combined the information obtained from GPS data for estimating the distribution of little penguins at sea, stable isotopes data to investigate their diet, as well as from the automated monitoring system to track changes in body mass and colony attendance.

Two automated penguin monitoring systems (APMS) are placed on the main pathways between the little penguins' colony and the beach. When walking through APMS, little penguins are individually identified with passive transponders (Allflex, Australia) that had previously been inserted in the back of the penguins, either as chicks or the first time they were encountered in the colony. In addition, APMS record date, time, direction of passage, and the body mass of the individuals (Joly et al., 2022).

This research was conducted under the Phillip Island Nature Parks Animal Experimentation Ethics Committee approval and a research permit issued by the Department of Environment Land, Water and Planning of the state of Victoria, Australia.

2.2. Data manipulation and analysis

Data manipulation and analysis was done in R v4.2.3 (R Core Team, 2023). Unless specified otherwise, results indicate mean and standard error. As well, when more than one variable was considered within a model, all model combinations were tested, and we performed model selection using Akaike's information criterion (AIC) (Bozdogan, 1987). The model with the lowest AIC was considered as best. Normality of residuals, residual autocorrelation and homoscedasticity were checked graphically. We considered *p*-values under 0.05 as significant. Unless stated otherwise, pairwise post-hoc comparisons were performed using Holm p-value correction (Holm, 1979).

2.3. GPS data processing

A foraging trip was defined as a period from the departure to the return to the colony. Because little penguins are visual hunters, foraging activities only occur during daylight (Cannell and Cullen, 1998; Chiaradia et al., 2007). Their foraging trips last typically between 1 and 9 days during incubation (Kato et al., 2008), 1 day during guard (Pelletier et al., 2014) and between 1 and 17 days during post-guard (Saraux et al., 2011a). During guard, parents alternate trips at sea, while in post-guard, chicks are left unattended in the colony (Saraux et al., 2011b). From each foraging trip, and day out at sea, we extracted a "foraging segment", intended as the period between nautical dawn and nautical dusk. Therefore, one-day trips contained only one foraging segment, while multiple-days trips could contain several segments. We removed foraging segments with <3 GPS locations, and segments starting after sunrise or stopping before sunset, from the analysis. Overall, out of 233 foraging trips, a total of 371 foraging segments were extracted and analyzed (range 1–7 segments per individual).

We calculated the distance between each consecutive location on the WGS ellipsoid using the *pointdistance()* function from the "raster" R package (Hijmans, 2022). Swimming speed was then calculated between two consecutive locations as the distance divided by the time interval. Furthermore, we excluded GPS locations with swimming speed higher than 8 km.h^{-1} (i.e., max swimming speed of little penguins, Watanuki et al., 2006), or with a time interval between 2 consecutive GPS locations lower than 7.2 s (i.e., duplicated points). GPS locations can be obtained only when penguins resurface, therefore it is necessary to interpolate raw GPS data and reconstruct their path. For each foraging segment, we regularized the time interval between each location by performing spatial interpolation at 15-min interval using the correlated random walk algorithm within the *crawlWrap()* function from "MomentuHMM" R package (McClintock and Michelot, 2018).

Interpolated foraging segments were projected into the GDA94/Australian Albers projection. For each breeding season, we then used the *kernelUD()* function from the "adehabitatHR" R package (Calenge, 2006) to calculate 50 % (core area), and 95 % (home range) kernel utilization distribution (UD). The smoothing parameter *h* was calculated using the ad hoc method (Seaman et al., 1998).

2.4. Stable isotope data processing

To describe the isotopic niches of little penguins and examine differences between 2020 and previous breeding seasons (2010–2019), we analyzed δ15N and δ13C stable isotopes from 842 blood samples (*n* = 196 in incubation, *n* = 367 in guard, *n* = 279 in post-guard). Values in δ15N increase with prey trophic level, while δ13C values are higher inshore than offshore (Hobson et al., 1994). We followed the protocol described in Chiaradia et al., 2016. Whole blood was freeze-dried and then powdered. As mass C/N ratios were all below 3.5, there was no need for correction of lipid contents in whole blood (Post et al., 2007). Isotopic analysis was then performed by means of a Robo-Prep elemental analyzer coupled to a Europe 20:20 continuous-flow isotope ratio mass spectrometer. Based on replicate measurements of within-run standards, measurement error was estimated to be ±0.3 and ±0.1 ‰ for δ15N and δ13C measurements, respectively.

2.5. Automated penguin monitoring system

We evaluated two measures of body mass variation. First, we calculated the mass gained after a foraging trip, which we considered to be an estimate of foraging efficiency (Saraux et al., 2011a). Two body masses were considered belonging to the same foraging trip when their records were consecutive in date and time for a given transponder number and the trip duration was not longer than 1 d in guard and 17 d in incubation and post-guard (Salton et al., 2015). Then, for post-guard only, we calculated the overnight mass variation after returning from a foraging trip, which we considered to be an estimate of chick provisioning during chick-rearing. During this stage, little penguins stay only a few hours in the colony, so we assumed that all body mass loss was due to chick provisioning. Body mass gained at sea was only considered when ranging from 700 to 1700 g and body mass change from −75 to 500 during incubation and 0 to 600 g during guard and post-guard (Joly et al., 2022; Saraux et al., 2011a).

Using APMS, we also calculated penguins' attendance at the colony. When a penguin crosses the weighbridge, it registers the timestamp, and transponder number of the penguin, allowing us to know departure and arrival times of each foraging trip. We calculated departure and arrival times relative to nautical dawn and dusk, respectively, to account for variation in day length (Rodríguez et al., 2016).

2.6. Proxies of anthropogenic activities

Given that little penguins breed on land and forage at sea, we defined both on land and at-sea indicators of anthropogenic activities. The number of tourists present each night at the Penguin Parade® was used as an index of human activity on land, as it is the only moment where penguins are exposed to tourists. At the Penguin Parade®, visitors are allowed only at sunset (when penguins come ashore, Rodríguez et al., 2016) for 50 min and confined to boardwalks where they cannot move around the penguins. After, the colony is returned to darkness. On average, around 2000 visitors attend the parade daily. Due to this setup, visitors never directly interact with penguins. Number of tourists at the Penguin Parade® was counted daily between 2010 and 2020. Over the studied period, artificial lighting (orange halogen lights, 3 lx) was used to enhance visibility of penguins for tourists. These lights were turned on from sunset to 1.5 h after the arrival of the first penguins (Rodríguez et al., 2016). During the COVID lock-downs, these lights were still in place but without the presence of tourists. The COVID lock-downs did not affect the monitoring protocol.

For the activity at sea, we used the number of vessels (fishing, commercial and leisure) within the little penguin foraging area (longitude 145 to 146°E, latitude 38.5 to 39.5°S) during their breeding season (September to December). We used the open-source dataset from the Australian Marine Safety Authority (https://www.operations.amsa.gov.au/Spatial/DataServices/DigitalData) and for each vessel we obtained its ID, latitude, longitude, type and timestamp. As vessels transmit their locations at different time interval (from one per 15 min to once a day), we built daily indices by keeping only one location per vessel and day: the closest to noon available. Data were available only between 2014 and 2020. Information earlier than 2014 was not considered because of the lower time resolution compared to later data, and data for November 2019 was missing. Hereafter, we refer to the number of vessels within little penguin foraging grounds as the marine traffic intensity. We calculated marine traffic UD using the same method described before for the little penguins.
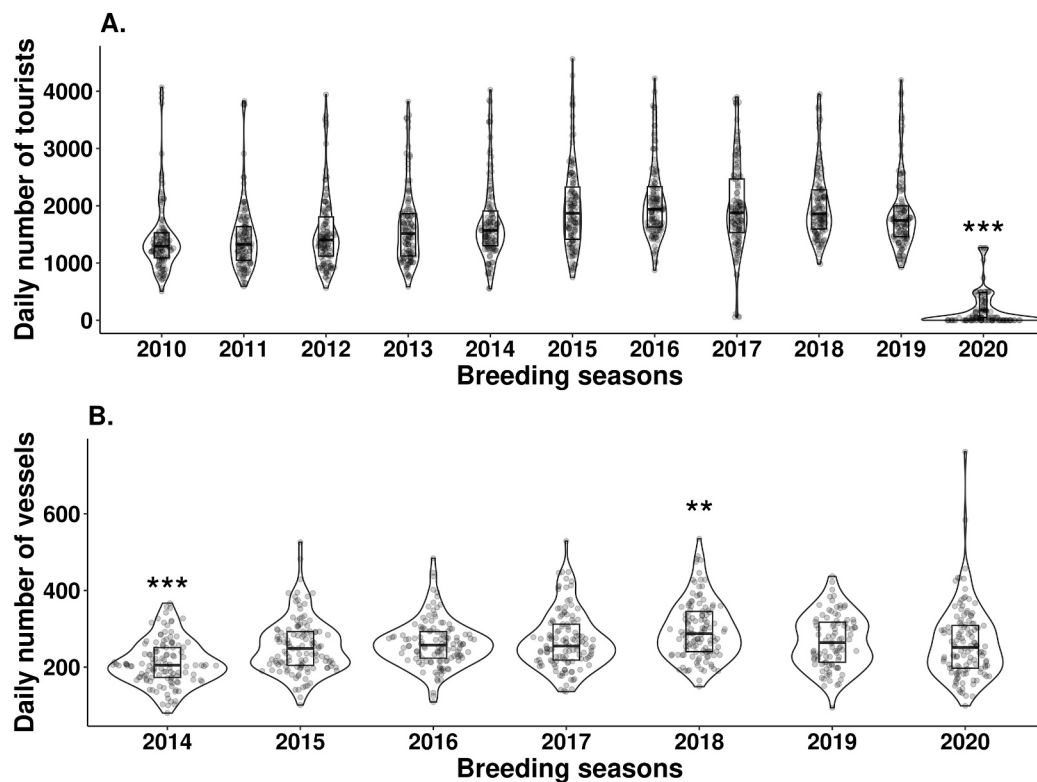
Fig. 1. Evolution of anthropogenic activities in the studied area. (A) Daily number of tourists at the Penguin Parade® between 2010 and 2020. (B) Daily number of vessels at sea in the foraging area of little penguins between 2014 and 2020. Asterisks represent statistical significance of post-hoc comparisons between 2020 (lockdown season) and the others (2010–2019, ** = $p < 0.01$, *** = $p < 0.001$).

2.7. Statistical analysis

2.7.1. Variation of anthropogenic activities

Using linear models, we investigated the variation of the number of vessels in little penguins foraging area and the number of tourists at the Penguin Parade® between months and breeding seasons. Then, using pairwise post-hoc comparison, we tested the difference between the COVID breeding season (2020) and the previous ones.

2.7.2. Spatial variation of at-sea distribution and overlap with marine traffic

Overlap analyses were performed using the Utilization Distribution Overlap Index (UDOI, Fieberg and Kochanny, 2005) which quantifies the pattern of space-use as a function of the product of the overlapping UDs. UDOI is equal to 0 when two UDs do not overlap and to 1 if the UDs are completely overlapping and uniformly distributed. Values higher than 1 indicate higher normal overlap relative to uniform space-use. UDOI were calculated using the *kerneloverlap()* function from the "adehabitatHR" R package (Calenge, 2006).

We calculated the UD overlap at 95 % of the distribution of penguins of breeding season A with the little penguin distributions of all the other breeding seasons, generating a distribution of UDOIs for breeding season A. We then assessed whether the observed distribution in 2020 was different compared to the previous breeding seasons. We also calculated the UDOI between little penguins and marine traffic. Again, we calculated UDOIs of breeding season A (little penguin) with all the marine traffic data between 2014 and 2020. This was done because marine traffic spatial distribution was stable throughout the studied period. We tested differences between breeding seasons using generalized linear models (GLMs) with a gamma distribution. We then performed post-hoc pairwise comparison to assess the significance of differences observed between 2020 and the other breeding seasons.

2.7.3. Effect of number of tourists on little penguins attendance and foraging efficiency

Linear models (LMs) were used to test the effect of lock-down on (a) average departure and arrival times of little penguin relative to nautical dawn and nautical dusk, respectively and (b) average mass variation per day over a foraging trip, and overnight. Models were built using the 'nlme' R package (Pinheiro et al., 2021). For both models (a and b) we considered number of tourists and number of vessels as explanatory variables, and breeding stage and season as fixed effects. To assess the effect sizes of both vessels and tourists counts, we standardized these data.

2.7.4. Quantification of isotopic niche

We computed standard ellipse area corrected for small sample size and extreme values (SEA_C) to estimate isotopic niche width and overlap among the different breeding seasons and stages. SEA_C represents the isotopic niche width of 40 % of typical individuals within the groups, based on bivariate normal distribution. The overlap in SEA_C was calculated for all pairs of breeding seasons within a breeding stage following (Catry et al., 2016) where isotopic niche overlap was expressed as a proportion of the area of overlap between two SEA_C to its own SEA_C. We also computed Bayesian Standard ellipse area (SEA_B) ($n = 20,000$ iterations) to obtain credible intervals (99 %, 95 % and 50 %) for the calculated ellipses. We considered non-overlapping 95 % CI around SEA_B as an indicator of statistically significant difference between niches width. For all this analysis, we followed the method described in (Jackson et al., 2011) and the 'SIBER' R package.

3. Results

3.1. Variation of anthropogenic activities

In 2020, during the COVID lock-downs, Phillip Island Nature Parks

UD levels (%) ■ Marine traffic 50% UD ■ Marine traffic 95% UD □ Little penguins 50% UD □ Little penguins 95% UD
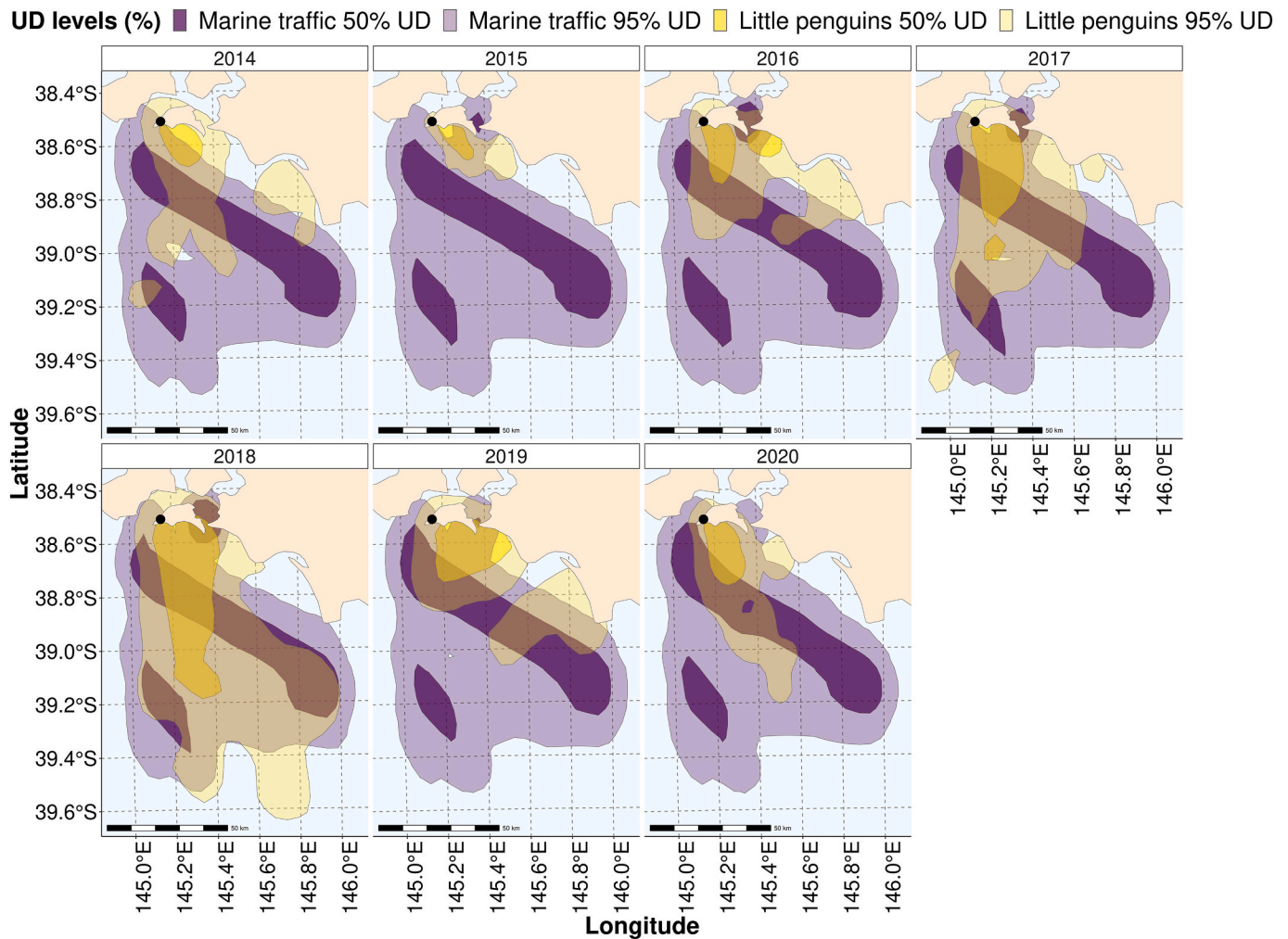

Fig. 2. Spatial distribution of little penguins and marine traffic between 2014 and 2020. The black dot represents the studied colony on Phillip Island.

remained closed for most of the breeding season, resulting in an average number of tourists 10 times lower than usual (180.4 ± 27.9 tourists per day in 2020 vs. 1770.0 ± 20.5 on average in 2010–2019, all $p < 0.001$, Fig. 1, Supplementary Table 2.A).

In 2020, the daily average number of vessels recorded at sea was 262 ± 8.51. This was significantly higher than the one recorded for 2014 of 212 ± 5.35 (estimate = 50.607, $t = 5.562$, $p < 0.001$), and lower than 2018 with 297 ± 6.91 vessels (estimate = −34.279, $t = −3.767$, $p = 0.002$, Fig. 1, Supplementary Table 2.B).

3.2. Spatial variation of at-sea distribution and overlap with marine traffic

While spatial distribution of marine traffic remained similar across seasons, little penguins core (50 % UD) and home ranges (95 % UD) showed great inter-annual variation across the studied period (Fig. 2). We compared the overlaps of penguins distribution in 2020 (average UDOI of 0.86 ± 0.07) to all the other breeding seasons (average UDOI ranging from 0.58 ± 0.09 in 2015 to 1.09 ± 0.07 in 2014) at 95 % UD. We did not find any significant difference in the overlap distributions in 2020 vs any other season (Supplementary Table 2.C).

Regarding overlap with marine traffic, model selection pointed at the model with the effect of the breeding season as explanatory variable as best (Supplementary Table 2.D). We found variation in the overlap between marine traffic and little penguins' distributions (from 2014 to 2020). In 2020, the overlap between little penguins and marine traffic

was significantly lower (0.184 ± 0.009) than in 2018 (average = 0.649 ± 0.144, estimate = 3.879, $p < 0.001$) and 2017 (average = 0.358 ± 0.128, estimate = 2.627, $p < 0.001$), but higher than the one of 2015 (average = 0.021 ± 0.003, estimate = −41.868, $p < 0.001$, Supplementary Table 3.D).

3.3. Effect of number of tourists and marine traffic on colony attendance

Little penguins left the colony on average 52.9 ± 0.5 min ($n = 11,116$) before nautical dawn and there was no difference across seasons. The best model testing the effect of anthropogenic activities on the time of departure relative to nautical dawn retained only the effect of breeding stage as explanatory variable (Supplementary Table 2.E, p-value < 0.01), with therefore no significant inter-annual variations (Supplementary Table 3.E) and no effect of the number of tourists or boats. During incubation, penguins left 31.8 min (95 % CI [24.5; 39.0]) before nautical dawn, compared to 74.9 min (95 % CI [67.7;82.2]) during guard and 47.1 min (95 % CI [39.9,54.4]) during post-guard (Fig. 3A).

Model selection for the models testing the effect of anthropogenic activities on the time of arrival relative to nautical dusk pointed at the null model as best (Supplementary table 2.F), indicating an absence of effect of tourists presence and marine traffic on colony attendance. Penguins showed highly synchronized arrival time regardless of season or breeding stage, arriving at the colony on average 8.2 ± 0.4 min after nautical dusk ($n = 11,087$, Fig. 3B, Supplementary Table 3.F).
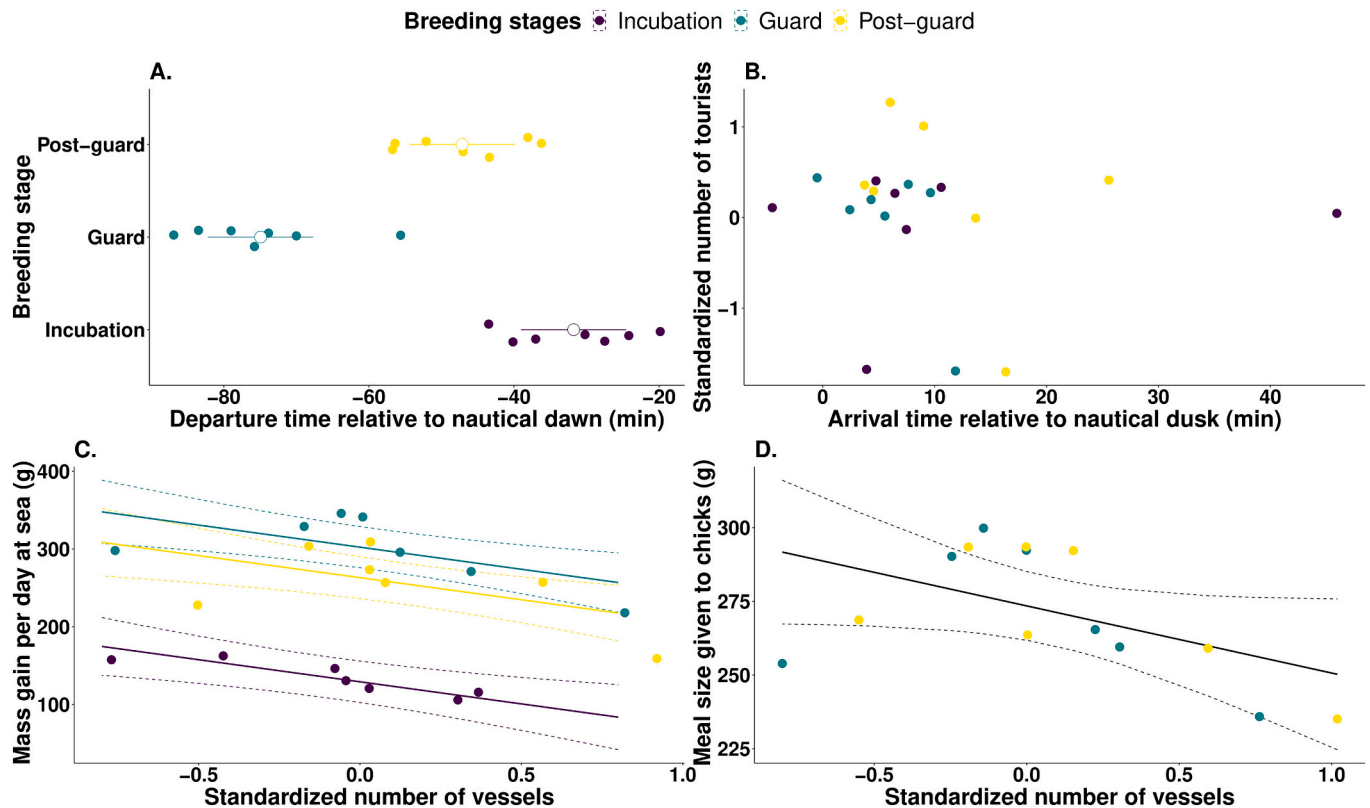
Fig. 3. Anthropogenic activities effect on APMS-derived parameters. (A) Departure time relatively to nautical dawn, and (B) Arrival time relatively to nautical dusk of little penguins at the colony depending on the number of tourists. Effect of the number of vessels on the (C) Mass gain per day at sea and (D) Meal mass given to the chicks at the colony. Colored points represent a season average, and white points the overall mean with its SE. Dashed lines represent the 95 % CI around model predictions.

3.4. Effect of number of tourists and marine traffic on foraging efficiency

Over a foraging trip, penguins gained on average 258.65 ± 1.75 g per day ($n = 6617$). The best model testing the effect of anthropogenic activities and temporal variations on mass gained per day at sea retained breeding stage and daily average number of vessels at sea as explanatory variables (Fig. 3, Supplementary Table 2.G), indicating an effect of marine traffic intensity but not of anthropause on foraging efficiency. Higher number of vessels was associated with lower mass gain at sea for little penguins (estimate $= -56.9 \pm 17.3$ g, $F = 10.8$, $p = 0.004$, Supplementary Table 3.G). Predicted breeding stage mass gain were all significantly different from one another ($F = 46.6$, all $p < 0.05$). During incubation, penguins gained 127.4 g (95 % CI [100.6;154.3]) per day, compared to 300.7 g (95 % CI [274.2;327.1]) in guard, and 261.4 g (95 % CI [234.6;288.2]) in post-guard.

The average overnight mass change during post-guard, i.e. chick meal size, was of 278.6 ± 0.1 g ($n = 1794$). Though the best model was the one with the average number of vessels at sea (Supplementary Table 2.H), its effect was not significant on meal size given to the chicks (estimate $= -22.8 \pm 11.2$, $F = 4.2$, $p = 0.06$, Fig. 3D, Supplementary Table 3.H).

3.5. Quantification of isotopic niche

A total of 842 blood samples were collected from little penguins across the different breeding stages (Supplementary Table 4). We observed variations in the isotopic niche values and areas at different breeding stages over 10 breeding seasons (Fig. 4). During incubation, the SEA_B mode of 2020 was 0.79 ‰^{2} with 95 % CI [0.48;1.25] and was significantly higher than 2 others breeding seasons, 2011 (0.19 ‰^{2} [0.13;0.31]) and 2015 (0.21 ‰^{2} [0.14;0.34]) (Fig. 5). During the guard

stage, SEA_B was higher for 2020 (0.98 ‰^{2} [0.69; 1.39] than 2011 again (0.24 ‰^{2} [0.17;0.33]), and 2010 (0.46 ‰^{2} [0.33;0.65]). Finally, during the post-guard, the SEA_B of 2020 decreased (0.60 ‰^{2} [0.45;0.87]). It was still significantly higher than the SEA_B of 2011 (0.29 ‰^{2} [0.21;0.42]), but also significantly lower than the one of 2014 (1.51 ‰^{2} [0.88; 2.72]). These inter annual variations led to low overlap between the isotopic niches of little penguins between 2010 and 2020.

4. Discussion

Humans have increasingly altered natural habitats, triggering changes in movements, habitat use and population dynamics in wild species (Holles et al., 2013; Margalida et al., 2014). The anthropause period caused by the COVID-19 pandemic set an unprecedented opportunity to study the effects of reduced human activities on the biology and ecology of a range of species (Rutz et al., 2020). During the anthropause, human activity on land decreased massively in our studied area, with a reduction almost by a factor 10 in the number of tourists of the Penguin Parade®, Phillip Island Nature Parks, Australia. Contrary to the expected (Bates et al., 2021), the lock-down policy did not seem to affect the marine traffic, neither spatially nor quantitatively within the penguin foraging zone within our study site in the Bass Strait. This specific setup allowed us to study the effect of on land activity through a stable at-sea potential pressure throughout the study period. Despite the important inter-annual variability in at-sea distribution and diet of little penguins over the period 2010–2020, no effect of the anthropause was found. Still, we found anthropogenic effect not linked with the anthropause. Despite the marine traffic intensity stability over the studied period, thanks to our long-term dataset, we were able to identify a negative relationship between marine traffic intensity and mass gained at sea per day by little penguins.
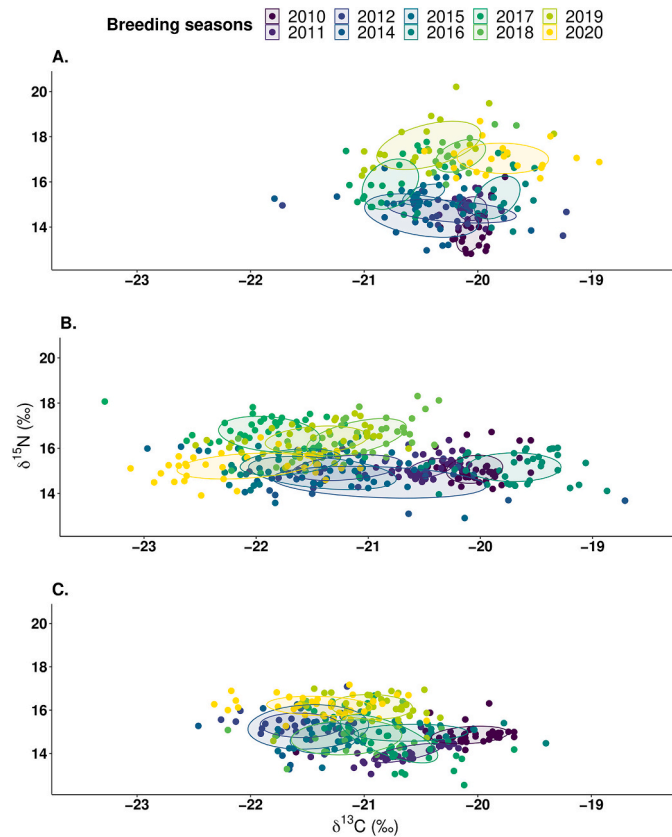
Fig. 4. Isotopic niche of little penguins between 2010 and 2020, across different breeding stages: A. Incubation, B. Guard and C. Post-guard. Ellipses represent the corrected standard ellipses of each niche (40 % of the individuals).

Fig. 5. Standard ellipses' area of little penguin's isotopic niches between 2010 and 2020 during different breeding stages: A. Incubation, B. Guard and C. Post-guard. Black dots represent the mode of the Bayesian standard ellipse area, and error bars the confidence intervals at 50, 95 and 99 %. Circles represent the corrected standard ellipse areas.

Human activities are known to affect seabirds' physiology and behavior. Previous studies showed the negative effects of anthropogenic noise (Pichegru et al., 2017), human presence (Ellenberg et al., 2013), domestic animal (Ratcliffe et al., 2010), food waste (Grémillet et al., 2008) and marine pollution (Trathan et al., 2015) on seabirds. A similar study (i.e., seabird in parks or area without tourists due to lock-downs) found that the absence of tourists could, counter-intuitively, lead to more disturbance for the seabirds, which translated in a later laying date and more egg predation (Hentati-Sundberg et al., 2021), underlining the protective role tourists can have for some species. We did not find such an effect in the parameters studied in this paper. On land, predators of little penguins are mainly goannas, snakes and cats (Colombelli-Négrel and Katsis, 2021). However, these predators are a not a threat on Phillip Island, thanks to the conservation program in place (BirdLife International, 2023).

Still, little penguins are known to be sensitive to anthropogenic activities like human presence around the nest (Colombelli-Négrel and Katsis, 2021). A recent study identified the negative effect of white light sources, at night on little penguins (Costello and Colombelli-Négrel, 2023), but evidences are mixed since another study found the opposite (Rodríguez et al., 2018). Multiple hypotheses could explain the absence of response during the anthropause in our study. One could argue that the duration and/or magnitude of the anthropause was negligible to trigger a response in the foraging behavior of little penguins. Plasticity being species dependent (Crawford et al., 2017), more studies on little penguins would be necessary to assess the extent of their plasticity in response to anthropogenic activities, and the potential different threshold that could trigger a response in the studied parameters (Cairns, 1988). Long-term exposure to tourists at the Penguin Parade®, could have habituated little penguins to anthropogenic disturbance
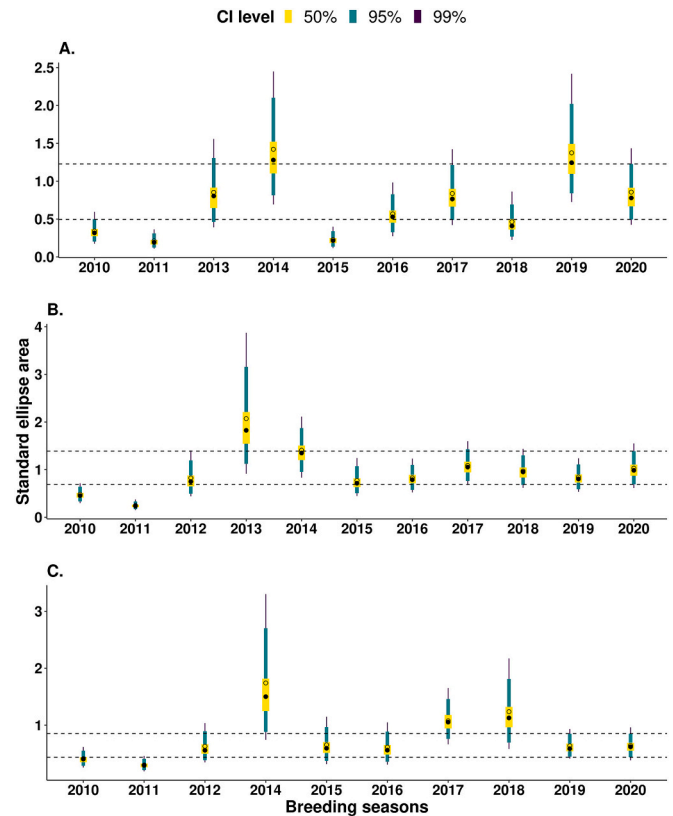
(Rodríguez et al., 2016). Our results underline the positive outcome of management and conservation actions put in place to mitigate human-wildlife co-existence. Similarly, several successful habitat restoration or predator removal programs have been carried out to aid conservation on both marine and terrestrial species (Moor et al., 2022; Jones et al., 2008).

To our knowledge, this paper provides the first empirical assessment of the negative effect of marine traffic on little penguins' foraging success during their breeding period. Our study underlines that the at-sea disturbances are more important than those on land when it comes to affect little penguins foraging. Our spatial analysis revealed an overlap between little penguins and marine traffic in the Bass strait. However, in the Bass strait, fisheries represent only a small proportion (<1 %) of the marine traffic in comparison with cargo (50–60 %), tanker (10–20 %), and passenger vessels (5–8 % Supplementary Fig. 1). It is therefore unlikely that the observed effect is due to a competition with fisheries for food. Collisions with leisure boats, for example, can represent a high mortality source for little penguins (Cannell et al., 2016). Increased traffic, especially of big cargo ships, leads to noise pollution affecting the behavior and reproduction of marine predators (Pichegru et al., 2022; Pirotta et al., 2022) and the distribution of preys (Ivanova et al., 2020). Another sentinel species of the Bass Strait, the Australian fur seal (*Arctocephalus pusillus doriferus*), showed behavioral changes (more alertness behavior) when in the presence of vessels (Speakman et al., 2020). More investigations are needed to fully understand how marine traffic impacts little penguins to be implemented on marine spatial planning.

Using quantitative information about human activity, like the number of tourists, rather than qualitative one (e.g., comparing lockdown season vs past observed trends) is key to be able to compare study results and assess the shape of the response of wildlife to

anthropogenic activities. Indeed, many studies fail to properly quantify anthropogenic activities, and only compared "COVID breeding season" with other breeding seasons before and/or after (Gordo et al., 2021; Hentati-Sundberg et al., 2021). While informative, this approach does not allow to properly disentangle anthropogenic pressure from seasonal and environmental variations. Incorporating a quantification of anthropogenic pressure in our models (i.e., number of tourists and vessels) allowed us to disentangle the natural inter- and intra-annual variations from anthropogenic pressure. Our study highlighted that long-term monitoring studies are key to be able to disentangle such effects. Occupying high trophic levels, sentinel species inform us about changes happening within ecosystems (Hazen et al., 2019). Therefore, long-term monitoring of such species, like the little penguin, can allow us to identify such changes at longer time scales and implement conservation actions (Duffy, 2002). For instance, the conservation actions associated with the return of wolves (*Canis lupus*) in the Yellowstone National Park, triggered consequences all along the trophic chain of this ecosystem (Berger and Smith, 2005).

The effect of the anthropauses caused by the COVID related lock-downs on little penguins' ecology during the breeding might be negligible compared to those induced by long-term environmental variations and global changes (Joly et al., 2022). Other significant effects found in our study are mostly related to intra and inter-annual variations. Thanks to long-term monitoring and online data availability, we were able to have a detailed picture of the impact of anthropogenic activities over the 10 breeding seasons. Species showing high plasticity and therefore quickly responding to reduced pressures during anthropauses are likely to use that same plasticity in the other way when anthropogenic activities increase again. Such punctual changes could be buffered by phenotypic plasticity and unlikely to change population trends compared to long-term variations (Gordo et al., 2021).

In conclusion, we did not detect any positive or negative effect of COVID-19 lock-downs on the little penguin breeding ecology. Behavioral variation during penguins' breeding cycle was mostly due to the inter- and intra-annual variation. Increased marine traffic can affect foraging efficiency of little penguins. Still, as seabirds live at the interface between sea and land, more information needs to be gathered on the mechanisms behind the effects of marine activities on little penguins' foraging. Our paper is of example of how conservation actions can be implemented to manage human-wildlife co-existence. Given the variability in the responses to anthropogenic activities and the fast changes of the marine environment, maintaining and developing long-term monitoring sites and studies are keys for guiding conservation policies. This will help scientists and stakeholders to better distinguish between environmental and anthropogenic effects on wild species.

CRediT authorship contribution statement

Conceptualization: AC, YRC, AK.
Data curation: BD, MC, NJ, AK.
Analysis: BD.
Writing: BD, MC.
Review of draft: all authors.
Supervision: MC, AK, AC.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

All R codes used to manipulate and analyze the data used are available at https://github.com/bendps/COVIDxLP. Data used in this paper are available at https://data.indores.fr/dataset.xhtml?

persistentId=doi:10.48579/PRO/6IQN7I.

Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.biocon.2023.110323.

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