



UK Centre for
Ecology & Hydrology

Hywind Scotland Ornithological Monitoring Programme

Overwintering distributions of common guillemot and razorbill populations in Eastern Scotland

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Contents

| | |
|---|-----|
| Executive summary | 2 |
| 1 Introduction | 4 |
| 1.1 Background | 4 |
| 1.2 Project objectives | 7 |
| 2 Methods | 8 |
| 2.1 Geolocation data collection | 8 |
| 2.2 Geolocation data processing | 100 |
| 2.3 Geolocation data analysis | 111 |
| 2.3.1 Species utilisation distribution (UD) | 111 |
| 2.3.2 Overlap in utilisation distributions | 122 |
| 2.3.3 Minimum adequate sample size of tracked birds | 122 |
| 2.4 Diet studies | 122 |
| 3 Results | 155 |
| 3.1 Species winter distribution..... | 155 |
| 3.1.1 Distribution maps | 155 |
| 3.1.2 Overlap in colony utilisation distributions | 522 |
| 3.1.3 Minimum adequate sample size of tracked birds | 555 |
| 3.2 Diet..... | 688 |
| 3.2.1 Breeding season in 2017 | 688 |
| 3.2.2 Breeding season in 2018 | 688 |
| 3.2.3 Breeding season in 2019 | 69 |
| 4 Discussion | 722 |
| 4.1 Guillemot and razorbill winter distributions | 722 |
| 4.1.1 Within- and between-species variation in winter distributions in relation to colony origin | 722 |
| 4.1.2 Minimum adequate sample size | 733 |
| 4.2 Diet variation among colonies and years..... | 744 |
| 4.3 Conclusions: winter ecology of guillemots and razorbills from the three colonies and implications for offshore renewable developments | 755 |
| 5 Acknowledgements | 777 |
| 6 References | 788 |
| 7 Appendix: Deployment details | 833 |

Executive summary

- The Hywind Scotland Pilot Park located off the east coast of Scotland has been in operation since 2017 and has a total generating capacity of 30 MW. Ornithological monitoring is required as part of its Marine Licence and is carried out in conjunction with a large-scale seabird monitoring and research programme in north-western Europe (SEATRACK) involving the deployment of geolocation loggers to obtain seabird location estimates throughout the winter. Of particular concern are two species of auks (common guillemot *Uria aalge* and razorbill *Alca torda*) known to be highly vulnerable to displacement and barrier effects from offshore renewable developments.
- As part of the Hywind Scotland's ornithological monitoring, this project aims to: 1) quantify year-round distribution and movements of guillemots and razorbills at three major colonies along the east coast of Scotland (East Caithness SPA, Buchan Ness to Collieston Coast SPA and Isle of May National Nature Reserve) over two years (2017-18 and 2018-19); 2) to quantify diet during the breeding season of guillemots and razorbills at the three colonies in three years (2017-19).
- In 2017, a total of 190 geolocation loggers were deployed at the three study colonies. During the 2018 breeding season, a total of 82 loggers were retrieved and 193 new data loggers were deployed. During the 2019 breeding season a total of 102 loggers were retrieved, including several deployed in the first year. We obtained reasonable sample sizes in all cases (range 11-28) except razorbills at Buchan Ness to Collieston Coast, where only five loggers were retrieved each year. The geolocation data were processed using a probabilistic method to obtain two locations per day for each bird throughout the non-breeding period. The geolocation data were analysed to determine utilisation distributions for each species at each colony in 2017-18 and 2018-19. Overlap in distributions was quantified and minimum adequate sample size of tracked birds was examined.
- For both species, the distribution of birds from the three study colonies was similar at a broad spatial scale, with key wintering areas located around the colonies and in the central and southern parts of the North Sea. In guillemots, among-colony spatial similarity in kernel densities was generally high throughout the non-breeding period. In razorbills, spatial similarity among colonies was highest in the post-breeding period and lowest in late winter.
- The analyses of minimum adequate sample size of tracked birds for each species at each of the colonies all showed the typical non-linear decline in rate of increase in the size of area used with increasing sample size. Although the cumulative curves did not level off, their shape, particularly in guillemots, suggested that a larger sample size would not have resulted in a much increased size of core wintering areas.

- Diet was quantified from observations of prey carried in the bills of breeding adults returning to the colony to feed their young. At all three colonies, the diet of guillemots was dominated by clupeids. On the Isle of May, guillemots brought back fewer 1+ group sandeels to their young compared to birds at the other two colonies. 0 group sandeels were unimportant at all three colonies. The diet of razorbills showed marked among-colony variation, with 0 group sandeels dominating at East Caithness and Buchan Ness to Collieston Coast and clupeids dominating on the Isle of May in 2017 and 2018. However, patterns were different in 2019 with 0 group sandeels the most important prey at all three colonies. Few 1+ group sandeels were observed in the diet of razorbills.
- *Conclusions.* Our results indicate that the three populations of guillemots and razorbills had a similar overall non-breeding distribution, with extensive use of the central and southern North Sea and areas around the breeding colonies, and this was broadly consistent between years. However, in both species there were important differences among colonies in the location of hotspots during the non-breeding period. The data provide important insights into the year-round space use of these two key species at three major colonies on the east coast of Scotland and their potential interaction with offshore renewable developments.

1 Introduction

1.1 Background

In 2015, Equinor ASA (formerly Statoil ASA) received a Marine Licence to develop the Hywind Scotland Pilot Park Project (“Hywind Scotland”). During the summer of 2017 five floating wind turbines (FWTs) with a total generating capacity of 30 MW were installed at Buchan Deep, approximately 25 km off the coast of Peterhead. The wind farm has been in full production since October 2017.

Ornithological monitoring is required as part of Hywind Scotland’s Marine Licence (Licence Number 05515/17/0) and is described in the Project Environmental Monitoring Programme (PEMP). Equinor is already involved in an ornithological monitoring and research programme which includes different seabird species and locations in north-western Europe (the SEATRACK programme). Equinor, as the operator of the Hywind Scotland project, agreed with Marine Scotland that Hywind Scotland’s ornithological monitoring programme builds on this ongoing work and extends the programme to include additional locations and species relevant to the project. Accordingly, this project delivers the ornithological programme as described in Hywind Scotland’s PEMP and agreed with the Licencing Authority (Marine Scotland Licencing Operations Team).

The focus of the current project is on two species of auks (common guillemot *Uria aalge* and razorbill *Alca torda*) considered vulnerable to displacement and barrier effects from offshore renewable developments (Furness et al. 2013, Dierschke et al. 2016). The interest in these species arose also because at-sea surveys carried out as part of the PEMP indicated that they are present in the Hywind Scotland project area in large numbers throughout the year (Natural Research Projects Ltd 2015). Of particular concern were the high densities recorded during August, a period which coincides with the post-breeding moult (when individuals are flightless) and with extended parental care (in the case of males), both of which can increase vulnerability to marine developments due to increased energy expenditure and reduced mobility.

Recent technological advances in animal-borne instrumentation have enabled the over-wintering ecology of seabirds to be studied in unprecedented detail (e.g. Phillips et al. 2007; Harris et al. 2010). Studies have deployed geolocation (“GLS”) loggers on a range of species across the globe, obtaining daily estimates of location, timing of movements and activity throughout the year. Important within-species variation in distribution, movements and behaviour outside the breeding season, both among individuals breeding at different colonies across the range, and among years has been shown (Frederiksen et al. 2012; Daunt et al. 2014; Bogdanova et al. 2017; Fayet et al. 2017). This new knowledge has been a critical step forward since most mortality of seabirds occurs in winter associated with reduced food abundance and poor weather conditions. Furthermore, individuals that are constrained by prevailing environmental conditions may be more susceptible to perturbations associated with anthropogenic activities, such as the effects of offshore renewable developments. As such, studies of over-wintering ecology from multiple colonies are key to establishing the drivers of change in seabird populations.

The SEATRACK project is a collaborative tracking project involving countries bordering the north-east Atlantic and North Sea whose objective is to understand over-wintering ecology across multiple colonies of a range of seabird species (<http://www.seapop.no/en/seatrack/>). The multi-colony approach enables space use and activity to be compared among populations in the north-east Atlantic. SEATRACK is ambitious and innovative, and will lead to important new insights in the year-round ecology of seabirds in the region. UKCEH is a partner in SEATRACK and currently deploys geolocation (GLS) loggers on four seabird species on the Isle of May National Nature Reserve: common guillemot (hereafter “guillemot”), Atlantic puffin *Fratercula arctica*, black-legged kittiwake *Rissa tridactyla* and European shag *Phalacrocorax aristotelis*.

This project addresses two limitations of SEATRACK that existed in the context of the over-wintering ecology of seabirds breeding in north-east Scotland:

- The Isle of May was the only UK colony represented in SEATRACK, where work is being carried out on the above four species. The addition of two other nearby colonies

(Buchan Ness to Collieston Coast and East Caithness SPAs) will result in considerable advances in our understanding of the extent of migration scheduling and aggregation (or segregation) during late summer moult and mid-winter of guillemots from the same region. This also allows us to test whether results from the Isle of May (Harris et al. 2015) are representative of other colonies along the east coast of Scotland.

- SEATRACK does not include the razorbill, a species of high conservation importance in the UK that shows strongly contrasting foraging and population ecology to its sister species, the guillemot (Thaxter et al. 2010; Glew et al. 2018; Dunn et al. 2019).

Working at additional colonies to the Isle of May in this project enabled us to propose a secondary goal of collecting diet data for both species. The Seabird Monitoring Programme (SMP) is a national programme of monitoring of the demography and diet of UK seabirds. It is the UK's equivalent to Norway's SEAPOP programme. The SMP monitors many sites around the UK, with varying intensity. There is current interest in the colonies on the east coast of Scotland because of proposed offshore wind farms in the region, including Hywind. For Buchan Ness to Collieston Coast and East Caithness SPAs, the quality of count data is moderate to good but there is a paucity of diet data (Anderson et al. 2014). Accordingly, diet data collection was planned for each study season. Given the regional significance of these colonies, it is strategically important to increase the intensity of diet monitoring so that any assessments of population change can be undertaken with a better understanding of predator-prey relationships, to complement data collected on the Isle of May as part of UKCEH's Isle of May long-term study (IMLOTS) that are being contributed to this project.

Furthermore, there is increasing evidence of a link between breeding conditions in summer and wintering ecology in seabirds (Daunt et al. 2006, 2014; Bogdanova et al. 2011; 2017). These so-called "carry-over" effects arise when conditions in one season have downstream consequences in subsequent seasons (Harrison et al. 2011). Impacts of summer breeding on winter ecology can occur because of the costs of reproduction; furthermore, winter ecology can affect condition in late winter and early spring, impacting on breeding scheduling and ultimately breeding success. By quantifying winter distribution and activity and summer diet, we will gain insights on seasonal interactions at the three study colonies.

1.2 Project objectives

This project has two principal objectives

- To quantify year-round distribution, movements and activity of guillemots and razorbills at East Caithness SPA and Buchan Ness to Collieston Coast SPA, and of razorbills on the Isle of May National Nature Reserve.
- To quantify diet during the breeding season of guillemots and razorbills at all the study colonies.

These data are compared with year-round distribution, movements and activity of Isle of May guillemots from the SEATRACK project to complete the structure of data collection on two ecological elements (winter ecology and summer diet) in two species at three colonies.

This report details all work carried out in the project between the 2017 to 2019 breeding seasons. The work comprised the deployment of data loggers and collection of diet data in 2017 (Daunt et al. 2017), retrieval of data loggers, deployment of new data loggers and collection of diet data in 2018 (Bogdanova et al. 2019) and the retrieval of data loggers and diet data collection in 2019. The report presents results on: 1) distribution of the two species at each study colony over the winters of 2017-18 and 2018-19; 2) similarity in the distribution of birds from the three colonies throughout each winter; 3) minimum adequate sample size of tracked birds required for a robust estimation of distribution (to ascertain whether we had enough data on each species at each colony) and 4) diet of each species at each colony in 2017, 2018 and 2019.

2 Methods

2.1 Geolocation data collection

During the 2017 breeding season, guillemots and razorbills breeding at the three colonies along the east coast of Scotland (Fig. 1) were captured at their nest sites with a noose or crook on the end of a long pole. Each bird was fitted with a geolocation logger (model MK3006, Biotrack, UK) attached to a plastic leg ring. A BTO metal ring was placed on the other leg, and the bird released. Deployment methods and protocols were the same at all colonies and handling time was less than five minutes in all cases. In subsequent years, colonies were visited and attempts made to capture birds carrying data loggers were made.

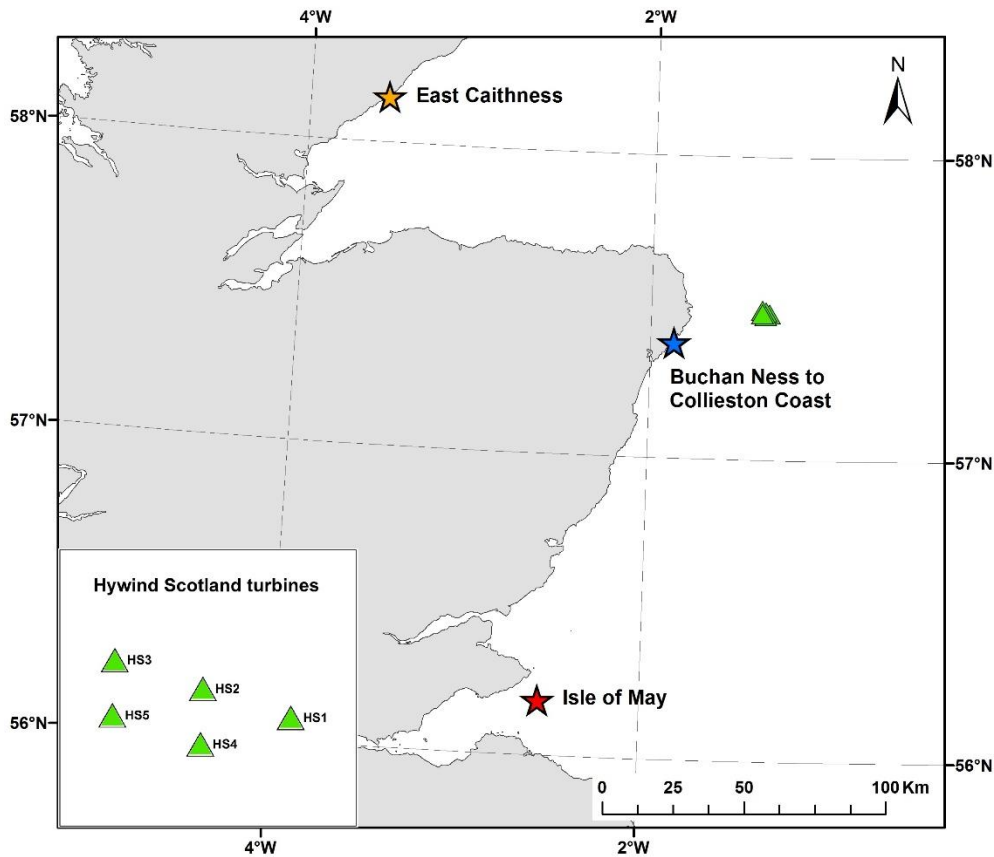


Fig. 1. Location of study colonies and Hywind Scotland Pilot Park Project (green triangles on main map and inset).

In total, 190 loggers were deployed in 2017, and 82 of these were retrieved in 2018 (representing 43% success rate). An additional 18 loggers were retrieved in 2019,

bringing the overall success rate over the project duration to 53%. Sample size of deployments and successful retrievals per colony and species is shown in Table 1.

| Colony and species | Deployed 2017 | Retrieved 2018 | Downloaded 2018 | Retrieved 2019 | Downloaded 2019 |
|------------------------------------|---------------|----------------|-----------------|----------------|-----------------|
| a) East Caithness | | | | | |
| Guillemot | 40 | 20 | 17 | 5 | 4 |
| Razorbill | 30 | 13 | 13 | 2 | 2 |
| b) Buchan Ness to Collieston Coast | | | | | |
| Guillemot | 40 | 24 | 24 | 4 | 4 |
| Razorbill | 20 | 2 | 2 | 3 | 3 |
| c) Isle of May | | | | | |
| Guillemot (as part of SEATRACK) | 30 | 12 | 12 | 4 | 4 |
| Razorbill | 30 | 11 | 11 | 0 | 0 |

Table 1: Summary of logger deployments in 2017 and number of birds tracked from each species and colony.

In 2018, a total of 193 loggers were deployed of which 84 were retrieved in 2019 (representing 44% success rate). Loggers were deployed on a mixture of individuals followed over 2017-18 and new individuals (see Appendix for details). Sample size of deployments and successful retrievals per colony and species is shown in Table 2.

| Colony and species | Deployed 2018 | Retrieved 2019 | Downloaded 2019 |
|------------------------------------|---------------|----------------|-----------------|
| a) East Caithness | | | |
| Guillemot | 40 | 22 | 22 |
| Razorbill | 30 | 5 | 5 |
| b) Buchan Ness to Collieston Coast | | | |
| Guillemot | 40 | 23 | 22 |
| Razorbill | 19 | 5 | 5 |
| c) Isle of May | | | |
| Guillemot (as part of SEATRACK) | 34 | 18 | 18 |
| Razorbill | 30 | 11 | 11 |

Table 2: Summary of logger deployments in 2018 and number of birds tracked from each species and colony.

2.2 Geolocation data processing

The geolocators measure light intensity every 60 seconds, and record the maximum value within each 10 minute interval. The loggers we deployed also have added sensors that record activity (wet – in the water or dry – on land/in flight) and temperature when the logger is submerged in water. The timing of twilight events (dawn and dusk) is estimated from thresholds in the light curves. To compute locations from the twilight events we used a new method involving an iterative forward step selection where each possible position is weighed using a set of parameters based on the species biology/behaviour (travel speeds) and environmental characteristics (sea surface temperature). The probabilistic algorithm used provides high accuracy of locations throughout the year, and substantially reduces location error particularly around the equinoxes due to the additional behavioural and environmental information incorporated (Merkel et al. 2016). This allowed us to retain in the processed dataset locations during the equinox periods when key migration movements often take place. All data

processing was carried out in R (version 3.6.1, R Core Team 2019) using package probGLS (Merkel et al. 2016; available from GitHub).

Although overall positions estimated using probGLS had considerably improved accuracy compared to those estimated using the standard threshold methods, plotting the processed data showed that some erroneous/implausible locations (where birds moved more than 500 km in a day) were included, particularly during the equinox periods. These represented 0.6% of the guillemot dataset and 1.4% of the razorbill dataset and were removed from the final dataset.

2.3 Geolocation data analysis

2.3.1 *Species utilisation distribution (UD)*

Utilisation distribution was determined for guillemots and razorbills from each study colony by calculating the kernel density of locations at sea. Locations were projected in Lambert azimuthal equal-area projection and kernel density was calculated in R (R development core team, 2019; package *adehabitatHR*, Calenge 2006), using a cell size of 1,000m² and a smoothing parameter h identified with the least-squares cross validation (LSCV) method (Worton 1989). For each species, the 50%, 75% and 90% UD contours were extracted in R (package *adehabitatHR*) and then mapped in ArcGIS. The 50% contour was used to define core areas and the 90% contour - to define the area of active use. Maps of population UDs were generated for the whole winter period (August to March) and for each month during the non-breeding season.

To assess the consistency of core areas among individuals, we determined individual UDs, extracted the 50% UD contours, converted these into grids with 2 km² cell size and calculated the total number of birds using each grid cell. The analysis was conducted in R using packages *adehabitatHR* and *raster* (Hijmans & van Etten 2018). We then generated maps of the cumulative individual core areas for the whole winter period in ArcGIS 10.4.1.

2.3.2 Overlap in utilisation distributions

To quantify variation in non-breeding distribution linked to breeding colony origin, we estimated the spatial similarity between the UD of birds from the three study colonies (based on the area of active use) using Bhattacharyya's affinity measure (Fieberg & Kochanny 2005). This measure ranges from zero (no similarity) to 1 (identical densities). Bhattacharyya's affinity measures are presented for each month and for the whole non-breeding period in 2017-18 and 2018-19.

2.3.3 Minimum adequate sample size of tracked birds

To establish whether the sample size of tracked individuals was adequate to estimate the core wintering areas used by the population of each species during the sampling period, we examined the relationship between size of core areas and number of individuals using a resampling procedure. This procedure was performed in R, and involved two steps: 1) calculating each individual's UD and 50% UD contour (using the `adehabitatHR` package within R); 2) calculating the total cumulative area for each sample size of birds ranging from 1 to n (where n denotes the total number of birds for which we had data) 1,000 times, by choosing birds randomly allowing replacement (Manly, 2009). The distribution of these areas across the 1,000 resampling rounds was used to quantify the typical at-sea area used for a given sample size of birds and the uncertainty associated with estimating this area. This analysis was carried out for the whole non-breeding period (August to March) in each year.

2.4 Diet studies

Diet of these two species has been studied on the Isle of May since the 1980s (Harris & Wanless 1985, 1986; Wilson et al 2004; Daunt et al. 2008; Thaxter et al 2013). To our knowledge, only one previous study of diet has been undertaken at Buchan Ness to Collieston Coast SPA (in 2006; Anderson et al. 2014), and no previous study of diet has been undertaken at East Caithness SPA.

The basic approach to assessing common guillemot and razorbill diet is observing prey carried in the bills of breeding adults returning to the colony to feed the young or, in the UKCEH report ... version 1.0

case of common guillemot only, to display to other adults. This is achieved by setting up an observation location with a clear view of a group of breeding birds with sufficient numbers of nests to obtain good sample sizes (up to 100 pairs), that is close enough that prey can be identified and a categorical size estimated using binoculars. All staff involved in observations had training and experience in these techniques. All data were collected in the field in notebooks and field data sheets. These were transcribed to spreadsheets after each check was made and files backed up on CEH servers. All notebooks and field sheets are archived at UKCEH.

In 2017, two hour watches covering the period of 04:00 – 22:00 twice were undertaken at East Caithness (13 June - 1 July) and Buchan Ness to Collieston Coast (19 June - 5 July). One plot was used for each species at East Caithness, and one plot for guillemots and two for razorbills at Buchan Ness to Collieston Coast. The rationale for using two plots for razorbills at the latter colony was that all chicks had fledged from the original plot, but staff were still *in situ*, so further samples were collected at a plot which still contained active nests in the remaining time available at the colony. On the Isle of May, two continuous dawn to dusk watches were carried out at separate plots used as part of the UKCEH long-term study on 21 June and 25 June. These two plots have been used for observation watches on the Isle of May since 1993. At all colonies, further observations were made opportunistically (range of dates the same as above for East Caithness and Buchan Ness to Collieston Coast; range of dates for Isle of May 2 April – 16 July).

In 2018, two-hour watches covering the period 04:00 - 22:00 twice were undertaken between 18 June and 4 July at East Caithness and between 25 June and 11 July at Buchan Ness to Collieston Coast. One plot was used for each species at both East Caithness and Buchan Ness to Collieston Coast. However, the plots used in 2018 at Buchan Ness to Collieston Coast were different to those used in 2017 for both species, because high failure rates at the original plots in 2018 resulted in few chicks and therefore limited opportunities for diet observations. On the Isle of May, dawn to dusk watches were carried out on 1 July and 3 July at the same two plots used in 2017. Further observations were made opportunistically (range of dates 28 April – 21 July).

In 2019, two-hour watches covering the period 04:00 - 22:00 twice were undertaken between 18 June and 4 July at East Caithness and between 10 June and 29 June at Buchan Ness to Collieston Coast. One plot was used for each species at both East Caithness and Buchan Ness to Collieston Coast. The plots used in 2019 at Buchan Ness to Collieston Coast were the same as those used in 2018 for both species. On the Isle of May, dawn to dusk watches were carried out on 24 June and 30 June at the same plots as in 2017 and 2018. Further observations were made opportunistically (range of dates 4 June to 17 July).

Guillemots return to the colony with a single prey item in their bill. During observation watches, time of arrival, prey species and prey size were recorded. For opportunistic observations, only the latter two were recorded. In all years of the long-term study on the Isle of May, the vast majority of prey are lesser sandeel or clupeids. Clupeids comprise sprat and herring, which are all but impossible to separate from observations in the field. In the current and previous years on the Isle of May, a small number of clupeids have been collected from breeding ledges and identified as sprat. However, all prey from this fish family are classed as clupeids in the dataset.

Prey were assigned to one of four size classes (very small, small, medium, large) using the bill length of the adult as a guide. When feeding a prey item to the chick the adult partly spreads its wings and arches its back, sometimes obscuring the view of the prey. All prey items presented in this way were presumed to have been eaten although classification by size and/or species was not always possible.

The protocol for collecting diet observations from razorbills was similar, except that razorbills may carry multiple prey in their bills. Thus, in addition to prey type and size, prey number was also estimated where possible, either as an exact estimate or as a broad category (“few”; “many”). Broader categories are necessary since estimating the number of prey in the bill is challenging. Fish found on guillemot breeding ledges allowed an estimate of the lengths of fish in each size category.

3 Results

3.1 Species winter distribution

3.1.1 Distribution maps

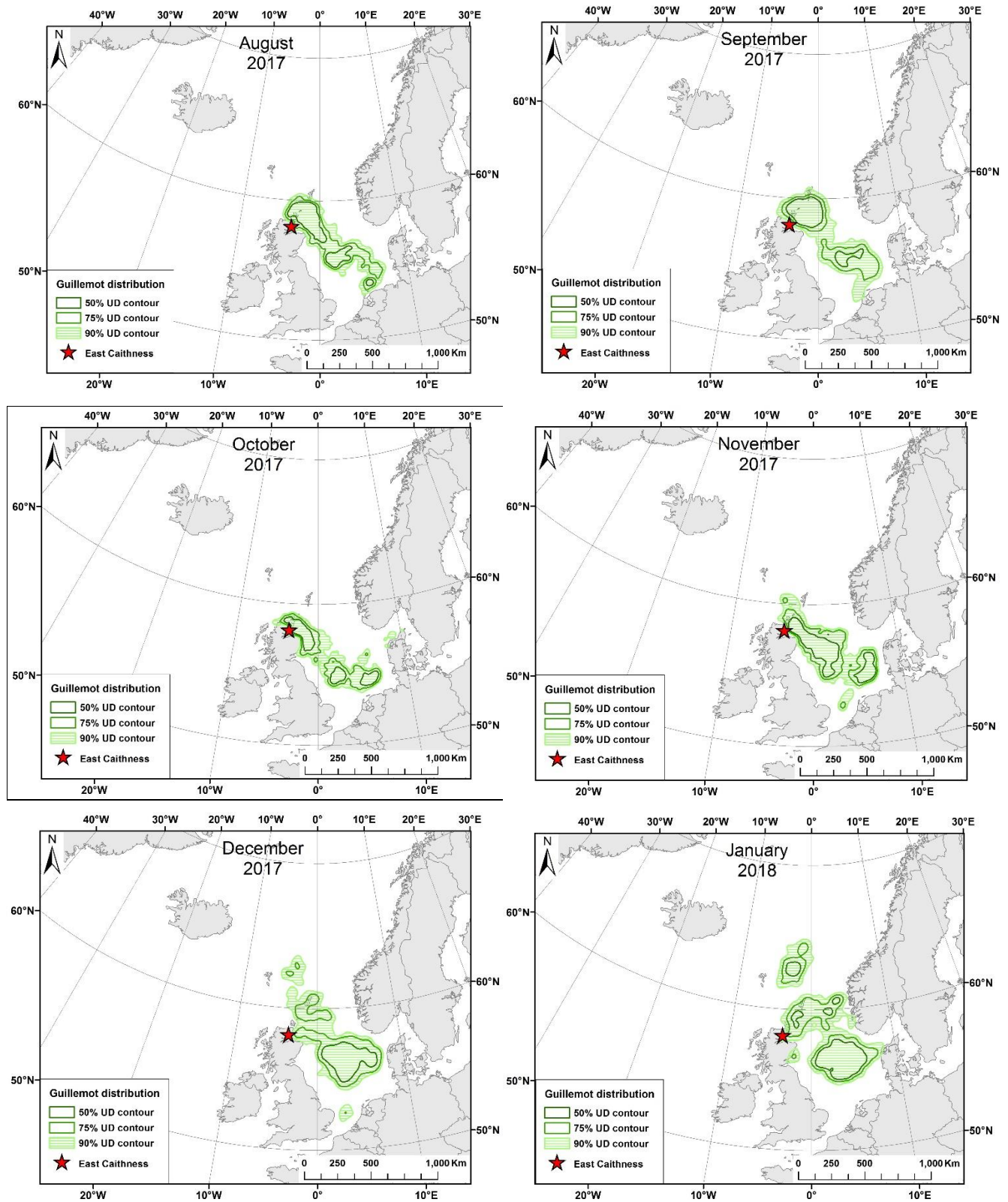
3.1.1.1 2017-18

Maps of utilisation distributions of guillemots in the first study year are provided in Figures 2-7. Corresponding maps for razorbills are presented in Figures 8-13.

After the breeding season in 2017, core areas of guillemots from East Caithness were located close to the colony and further south within the central North Sea (Fig. 2a). Between September and November areas near the colony and in central and south-western parts of the North Sea were used (Fig. 2b-d). In mid-winter most birds had moved further away from the colony, with core areas concentrated in the central and northern North Sea as well as the Norwegian Sea (Fig. 2e-f). In February, key areas extended from the central North Sea to the area of the colony, suggesting that part of the population returned close to the breeding grounds already in late winter (Fig. 2g). In March, core areas were located in the central North Sea, around the colony and in the Norwegian Sea indicating that some birds made substantial movements north in the pre-breeding period (Fig. 2h).

Overall, throughout the non-breeding period, the guillemots from East Caithness ranged across the North Sea and in parts of the Norwegian Sea but two key areas were apparent – one surrounding the colony and the other in the central parts of the North Sea (Fig. 2i). In line with this, individual core areas overlapped to greatest extent near the colony where areas were used by up to 12 out of 21 birds, and to lesser extent in the central parts of the North Sea - used by 5-9 out of 21 birds (Fig. 3).

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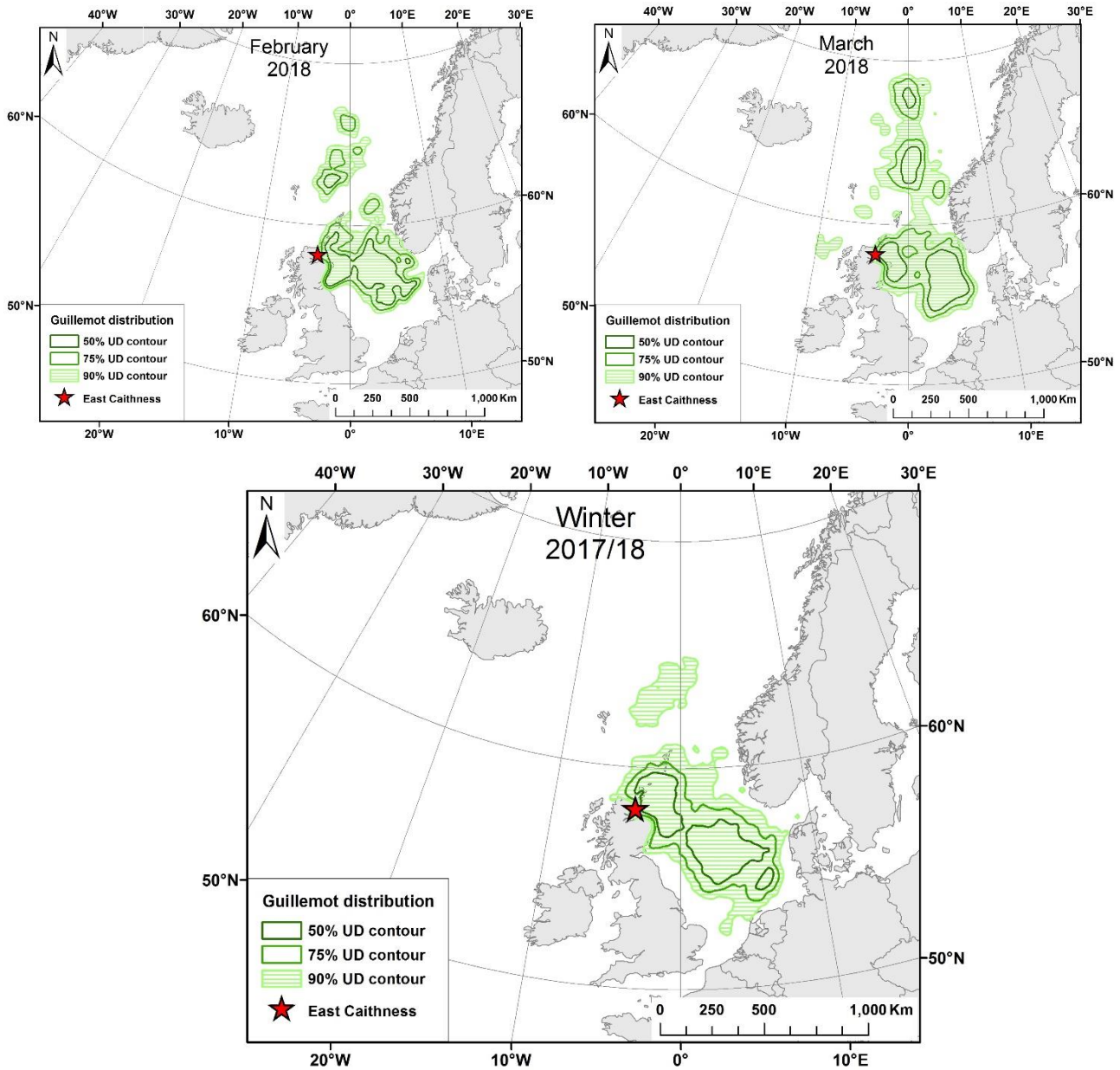


Fig. 2: Utilisation distributions (50%, 75%, 90% contours) for guillemots from East Caithness ($n=21$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

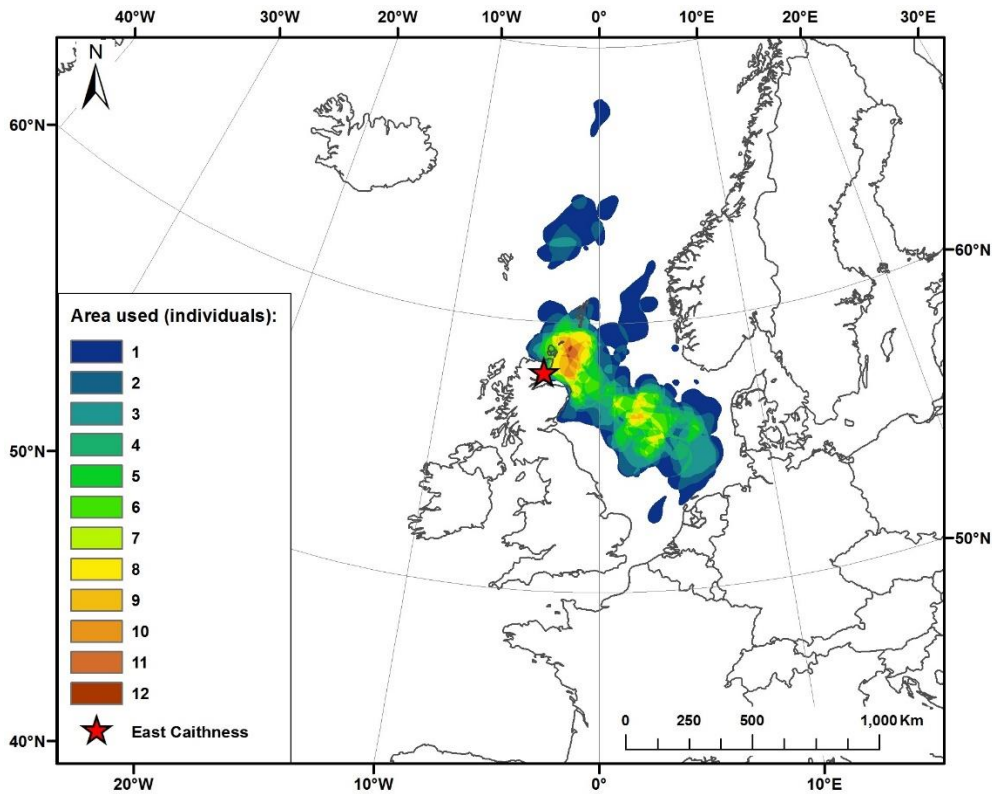
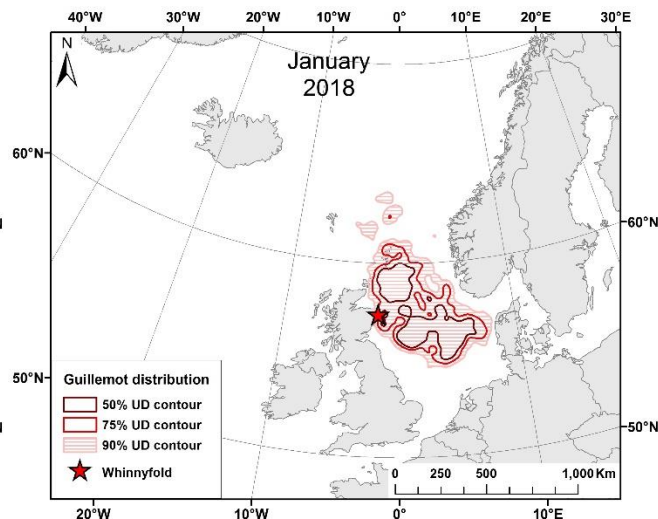
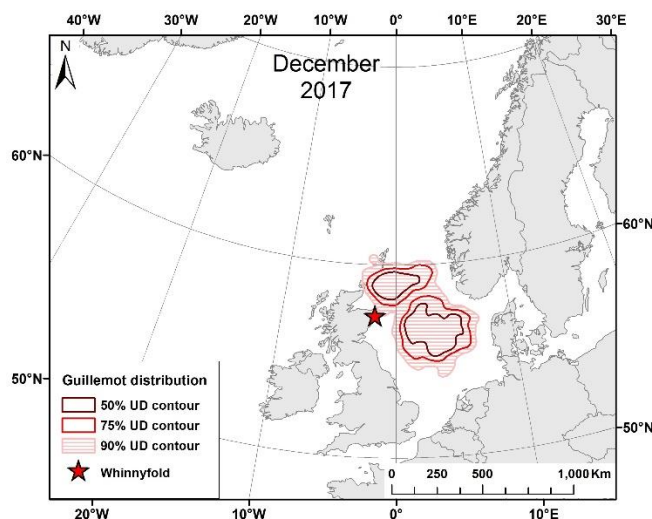
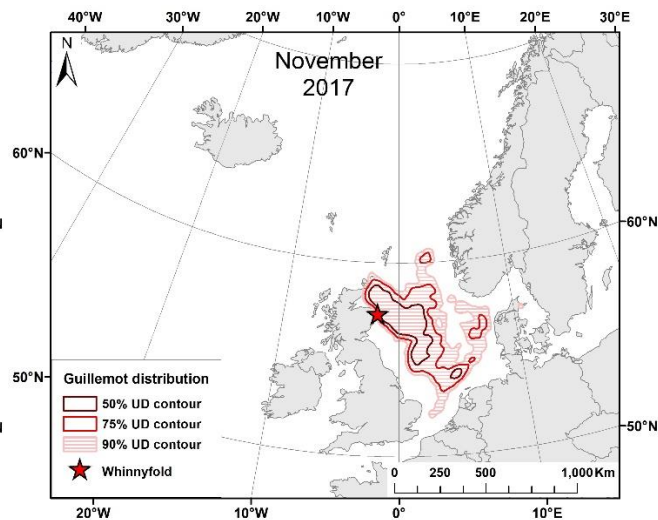
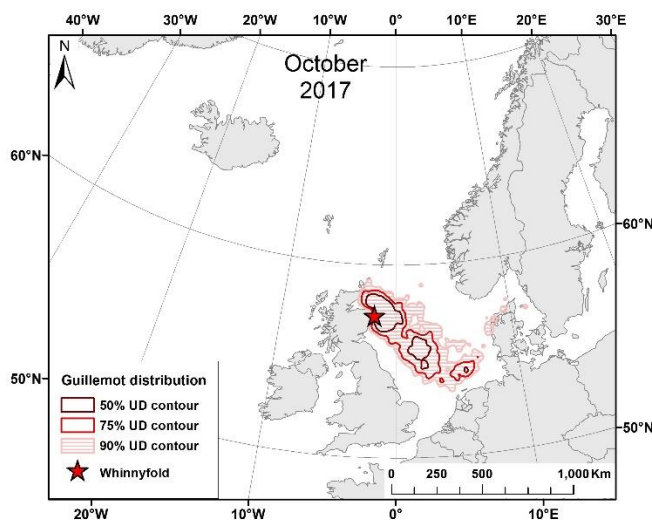
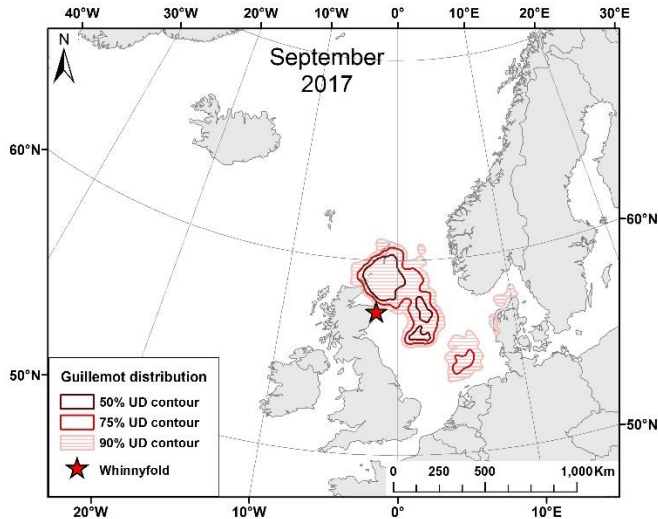
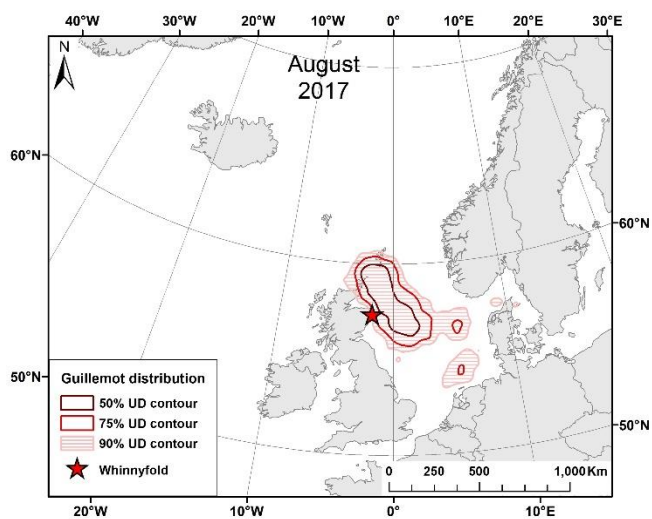


Fig. 3: Consistency in core areas (50% UD contours) of guillemots from East Caithness (n=21) in winter 2017-2018 (August – March).

In 2017, during the post-breeding period (August) guillemots from Buchan Ness to Collieston Coast remained within the northern North Sea, with core areas located close to the colony (Fig. 4a). Between September and November core areas were near the colony and in central and south-western parts of the North Sea (Fig. 4b-d). In December and January, two key areas were apparent - one in the central, the other in the northern North Sea (Fig. 4e,f). Similar to birds from East Caithness, in February and March core areas extended from the central North Sea, through the northern North Sea to the Norwegian Sea, including the area around the colony (Fig. 4g,h).

Overall, during the non-breeding period, guillemots from Buchan Ness to Collieston Coast ranged across the North Sea and parts of the Norwegian Sea but core areas were concentrated around the colony and in the central North Sea (Fig. 4i). Accordingly, highest overlap in individual core areas was observed within the area around and north of the colony, which was used by up to 18 out of 28 birds (Fig. 5).

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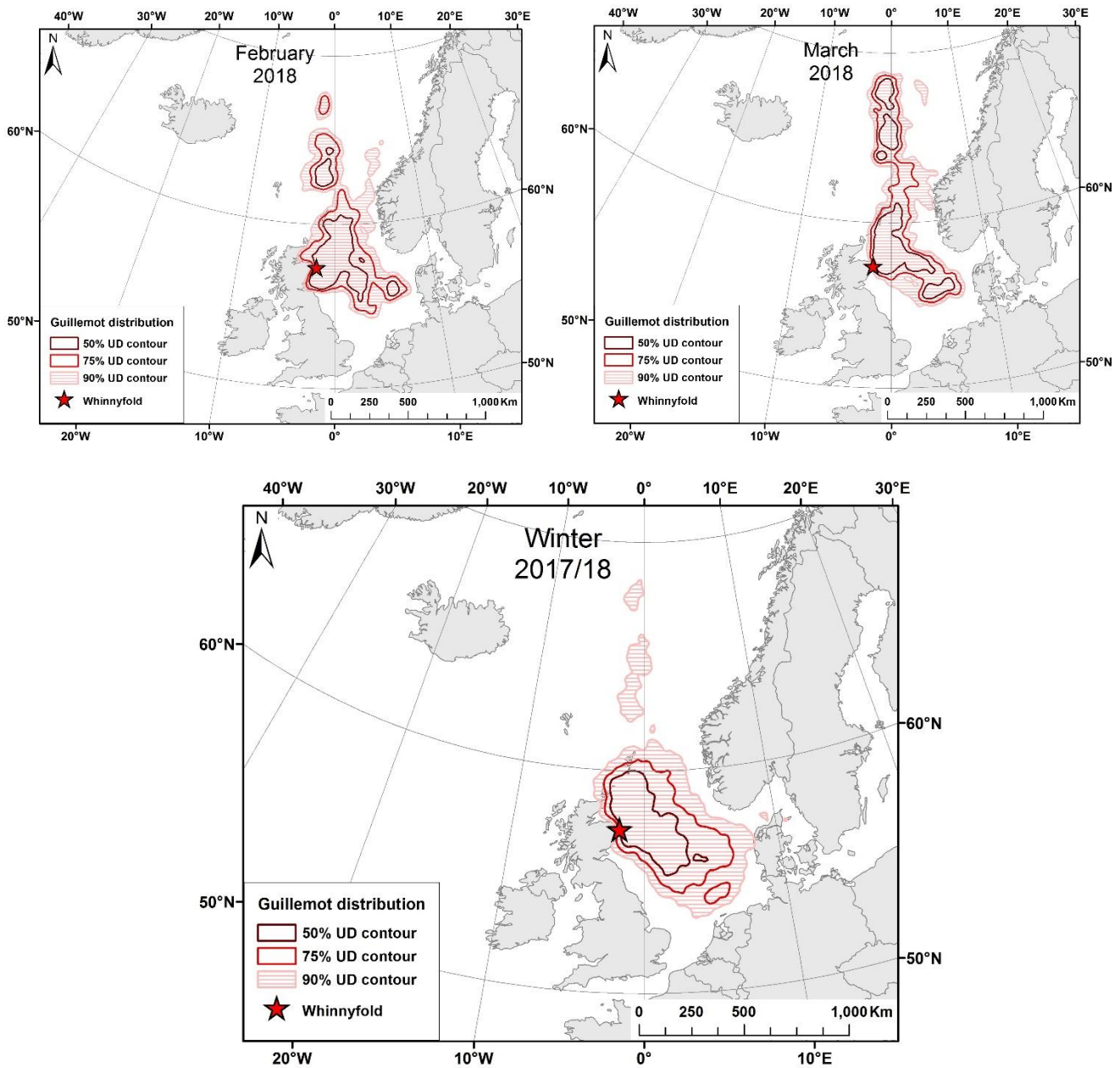


Fig. 4: Utilisation distributions (50%, 75%, 90% contours) for guillemots from Buchan Ness to Collieston Coast ($n=31$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

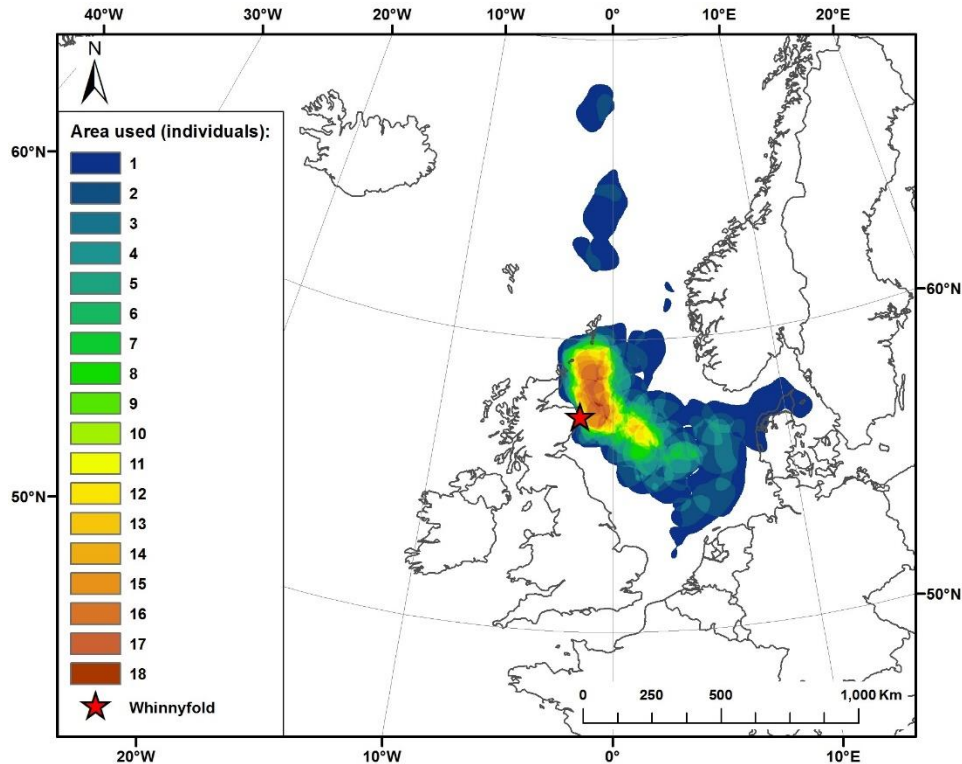
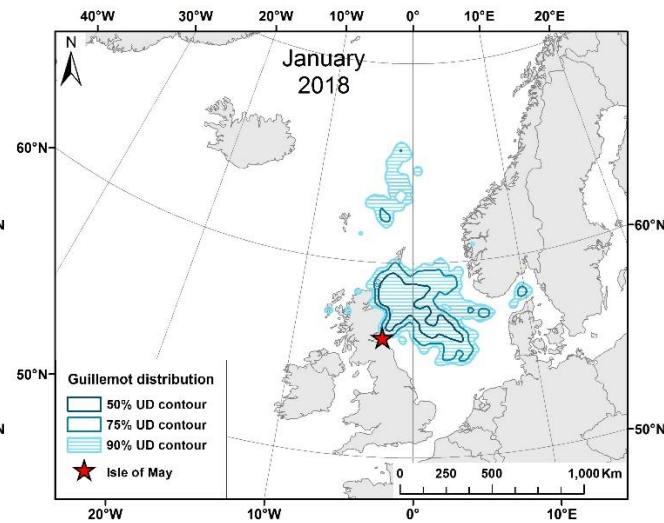
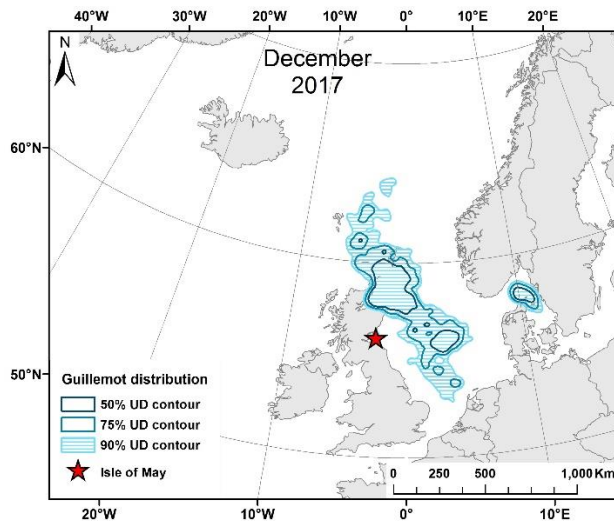
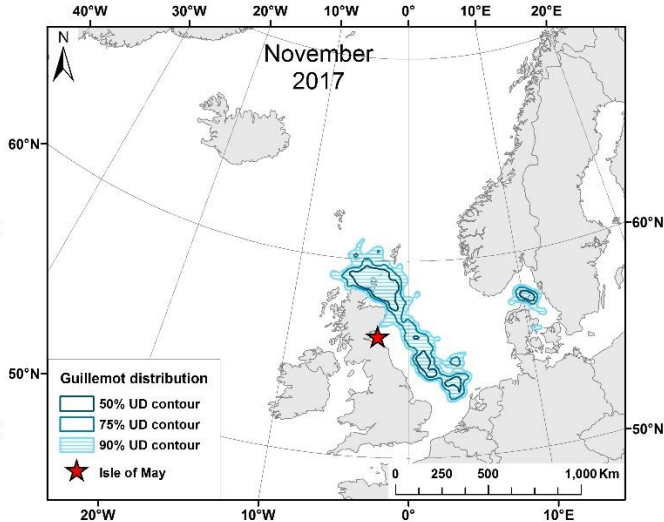
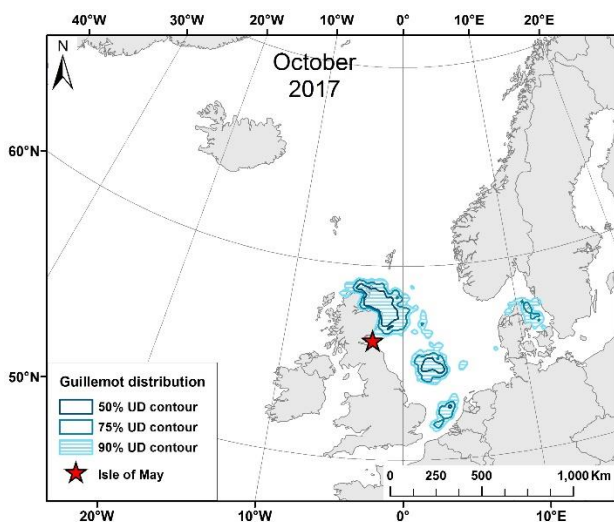
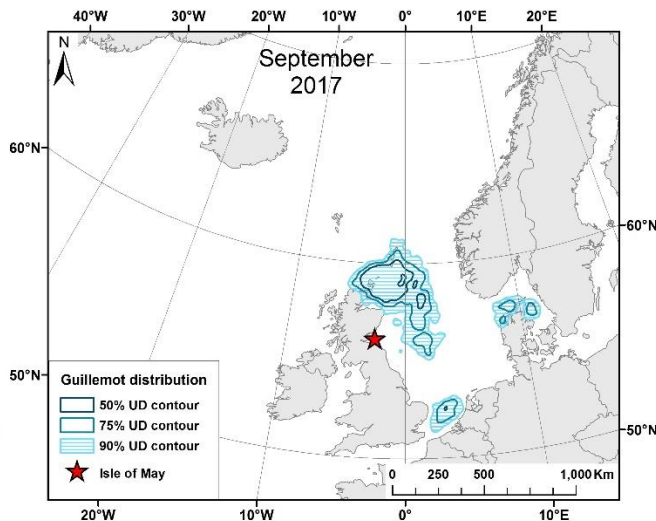
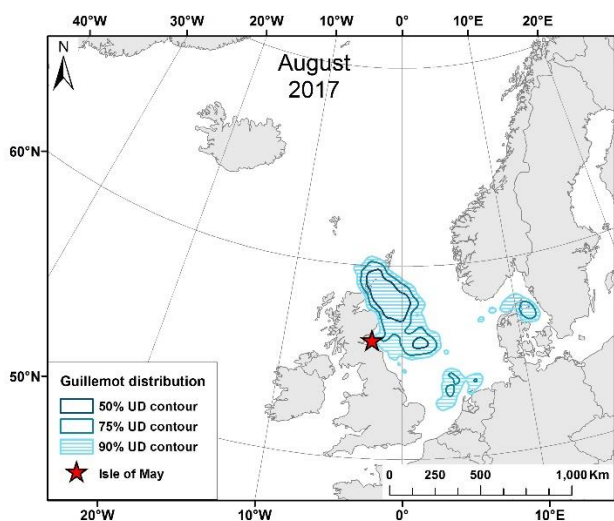


Fig. 5: Consistency in core areas (50% UD contours) of guillemots from Buchan Ness to Collieston Coast (n=31) in winter 2017-2018 (August – March).

During the post-breeding period in 2017, Isle of May guillemots were distributed mainly to the north and east of the colony, with some individuals moving to the southern North Sea and the Skagerrak/Kattegat area that links the North and Baltic Seas (Fig. 6a). Over the following months (September – December) core areas were located off north-east and northern Scotland, in the southern North Sea and around the Dogger Bank and in the Skagerrak/Kattegat area (Fig. 6b-e). In January the core areas were within the central and northern North Sea, including the area surrounding the colony (Fig. 6f). In February and March core areas extended from the northern North Sea to the Norwegian Sea, including the area around the Isle of May (Fig. 6g,h).

During the whole non-breeding period, guillemots from the Isle of May ranged widely across the North and Norwegian Seas but the key areas they used were mainly located in the waters off north-east Scotland (Fig. 6i). In line with this, highest overlap in individual core areas was observed within this area, which was used by up to 12 out of 16 birds (Fig. 7).



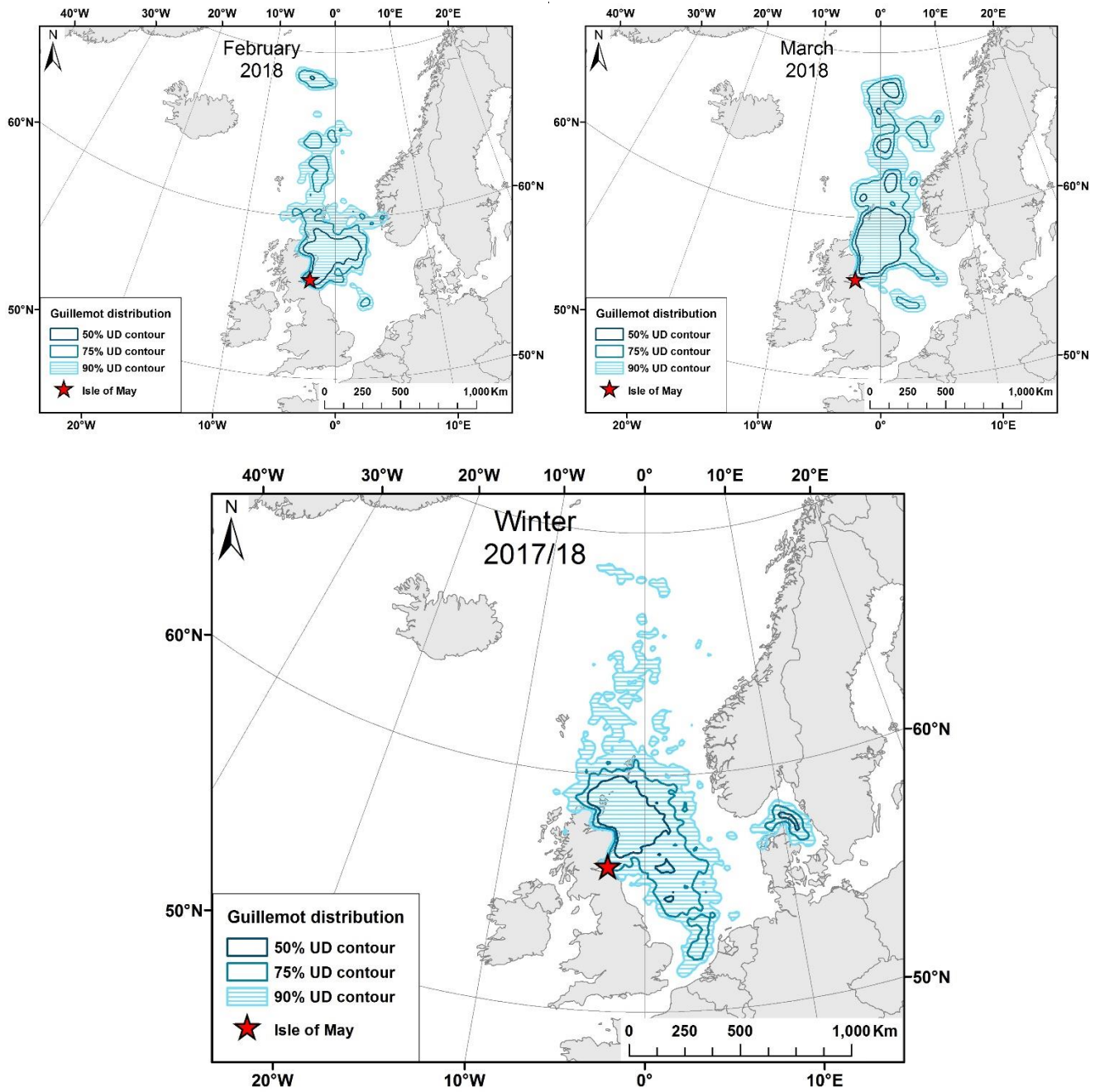


Fig. 6: Utilisation distributions (50%, 75%, 90% contours) for guillemots from Isle of May ($n=16$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

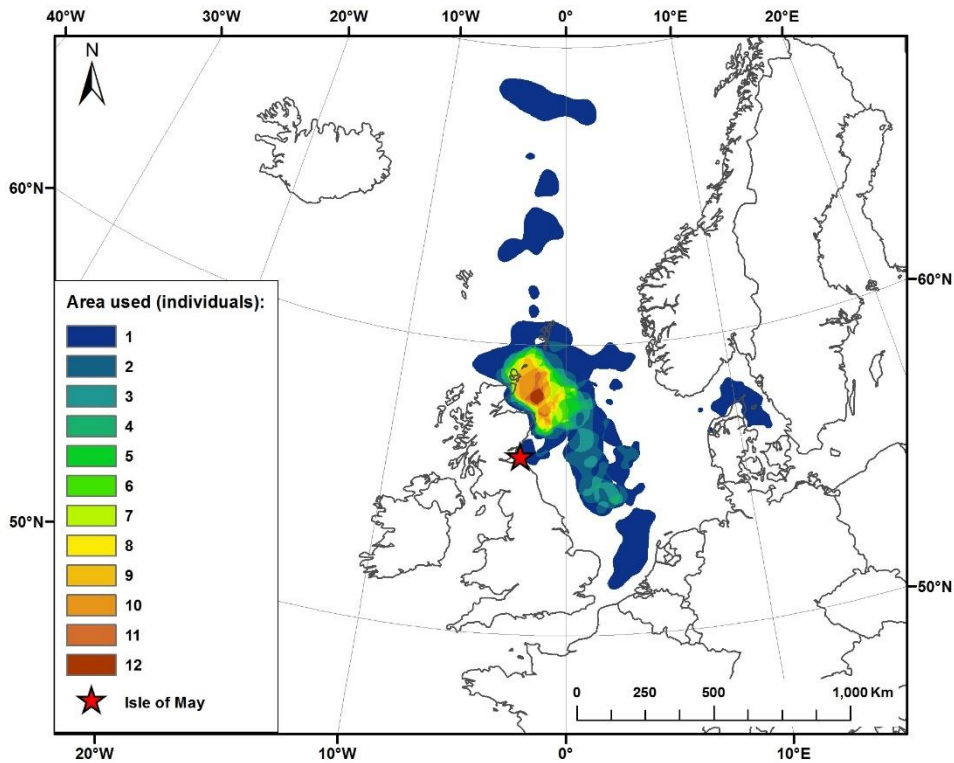
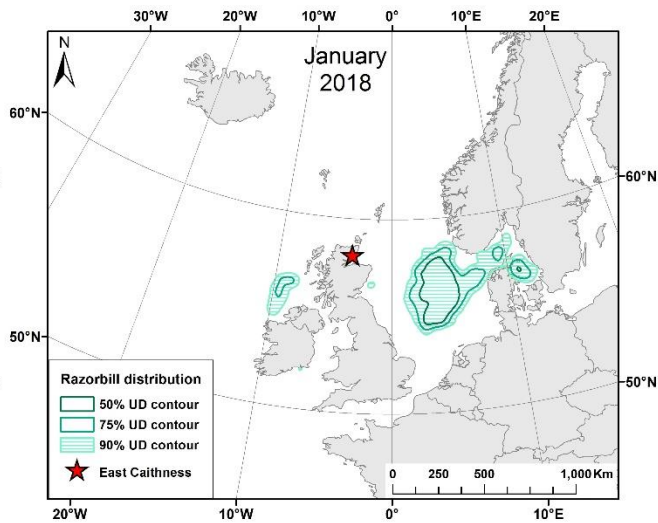
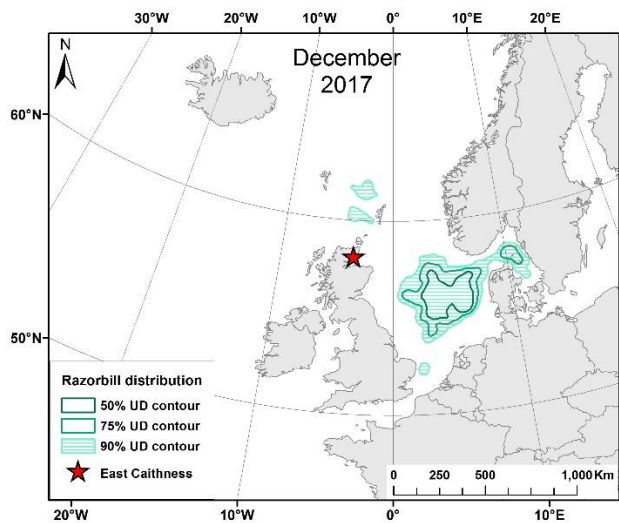
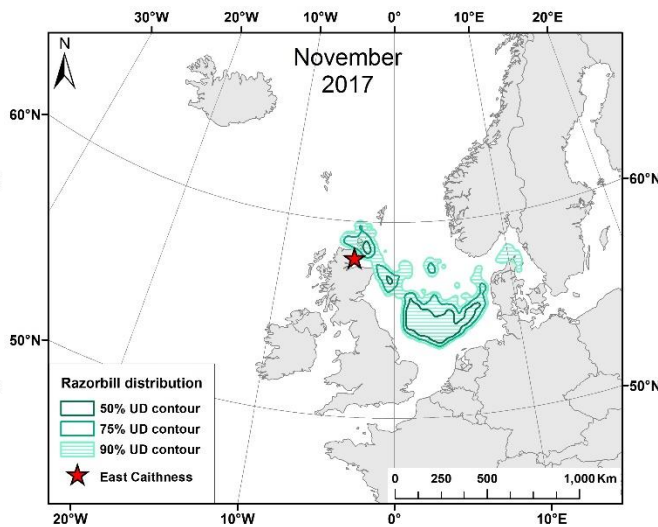
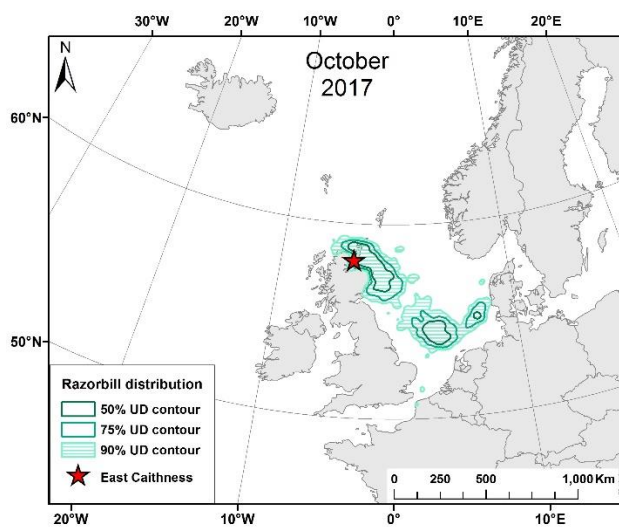
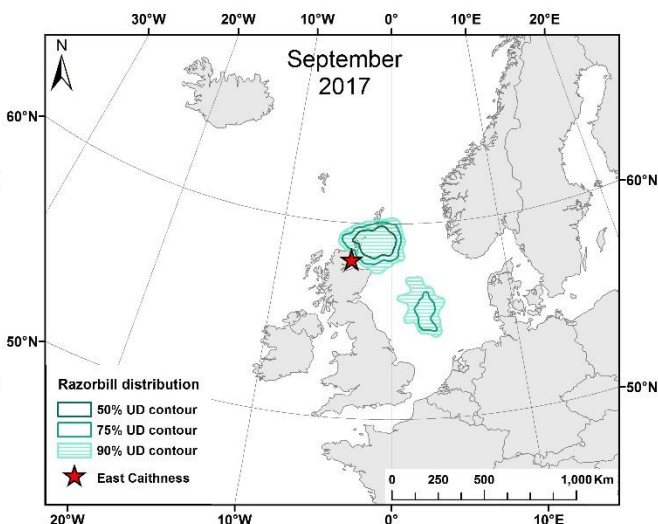
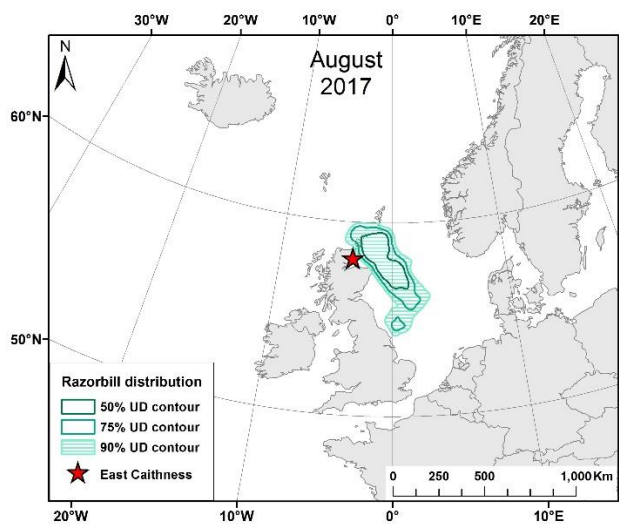


Fig. 7: Consistency in core areas (50% UD contours) of guillemots from Isle of May (n=16) in winter 2017-2018 (August – March).

After the breeding season in 2017, razorbills from East Caithness stayed mainly within the area surrounding the colony (Fig. 8a,b). During October and November, the birds gradually moved to the central North Sea where they remained throughout the winter months (Fig. 8c-g). In January and February, single birds moved to the Skagerrak/Kattegat region and to the north-east Atlantic, Irish and Celtic Seas (Fig. 8 g,h). By March, core areas were concentrated in the central North Sea and the east coast of Scotland including the area around the colony (Fig. 8h).

Overall, during the non-breeding period, the East Caithness razorbills ranged across the North Sea but two key areas were apparent – one in the central North Sea and the other off the east coast of Scotland (Fig. 8i). Accordingly, consistency in individual core areas was highest within these areas, with up to 8-11 out of 15 birds overlapping (Fig. 9).

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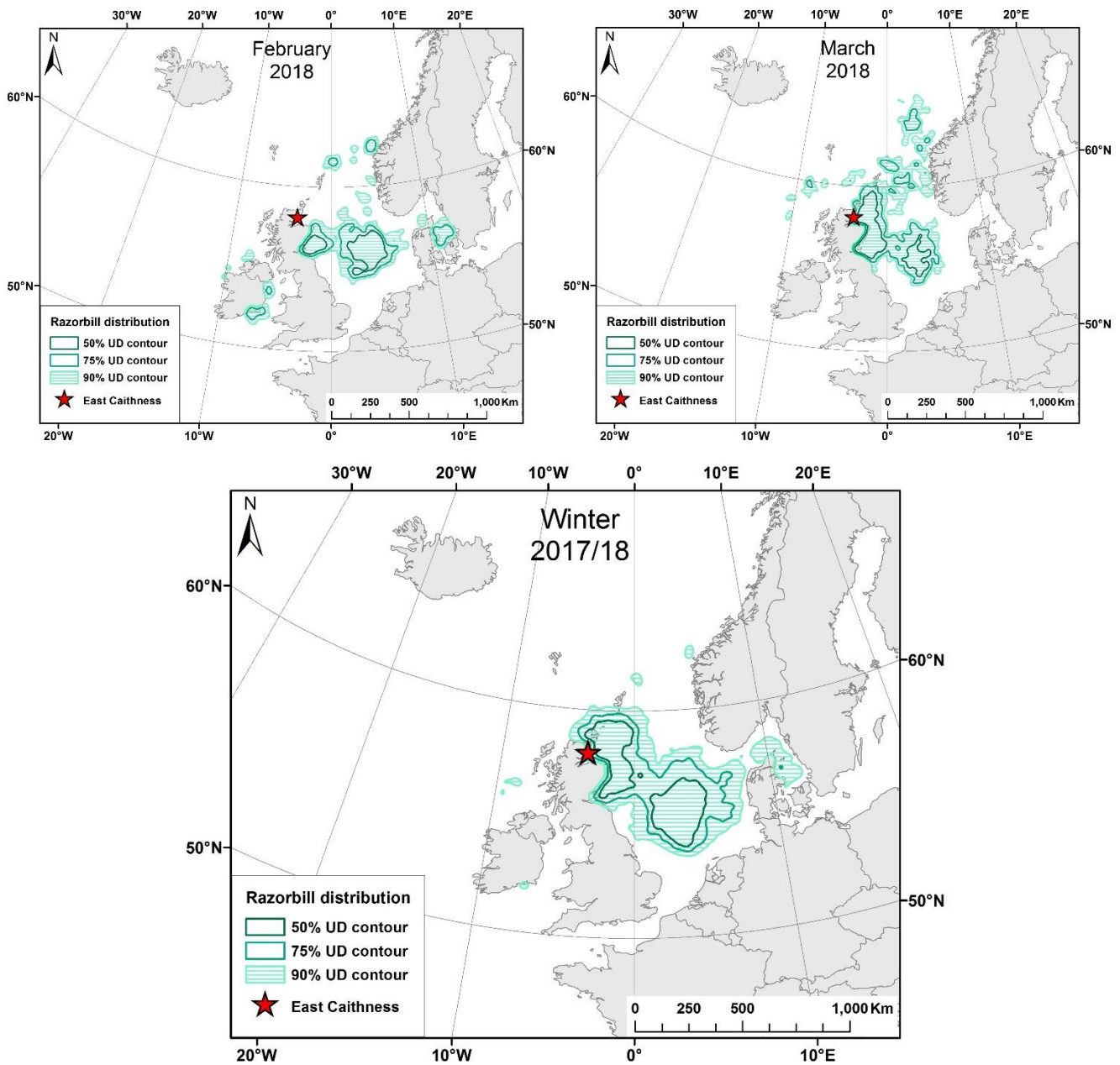


Fig. 8: Utilisation distributions (50%, 75%, 90% contours) for razorbills from East Caithness ($n=15$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

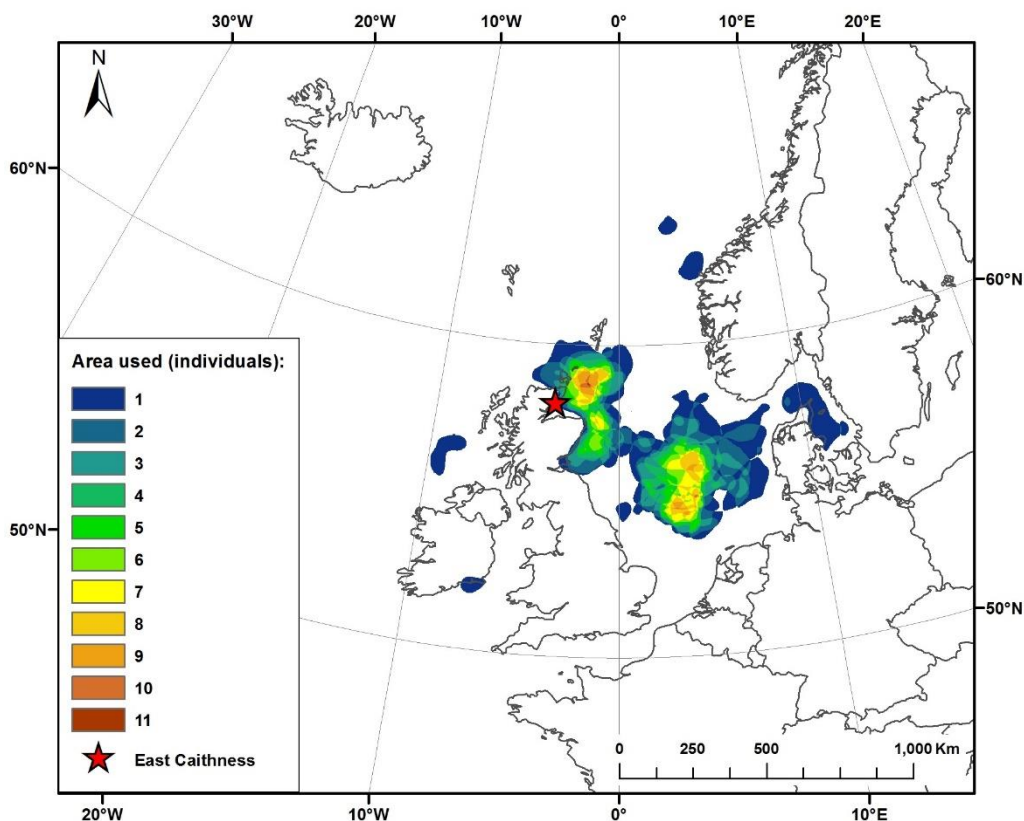
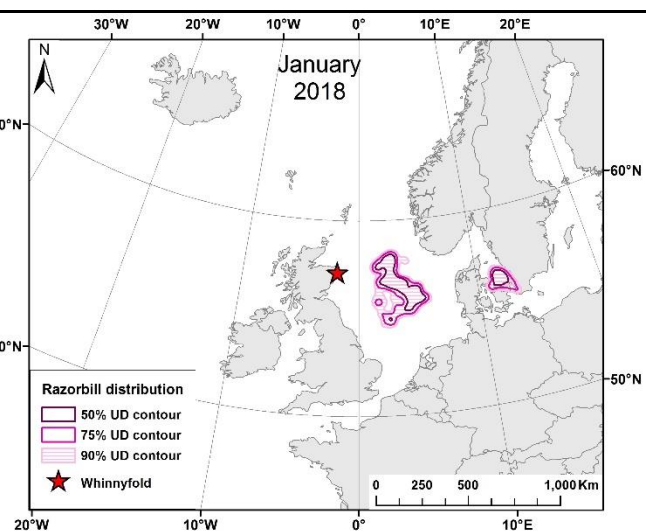
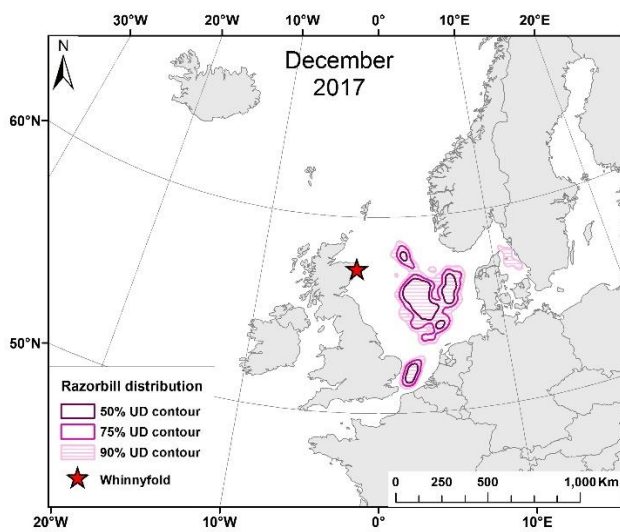
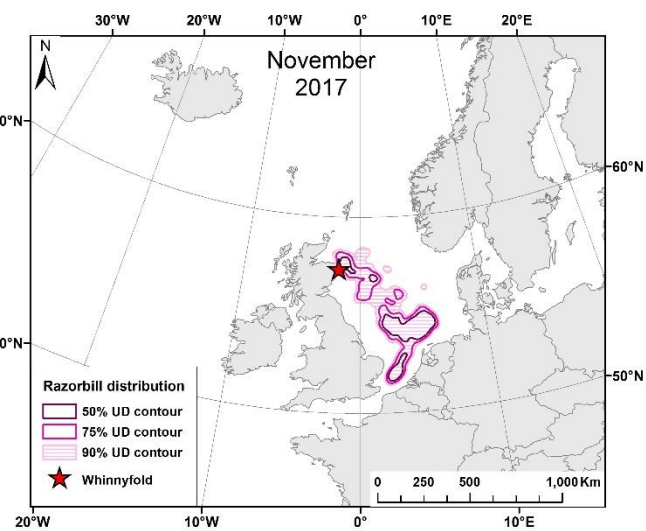
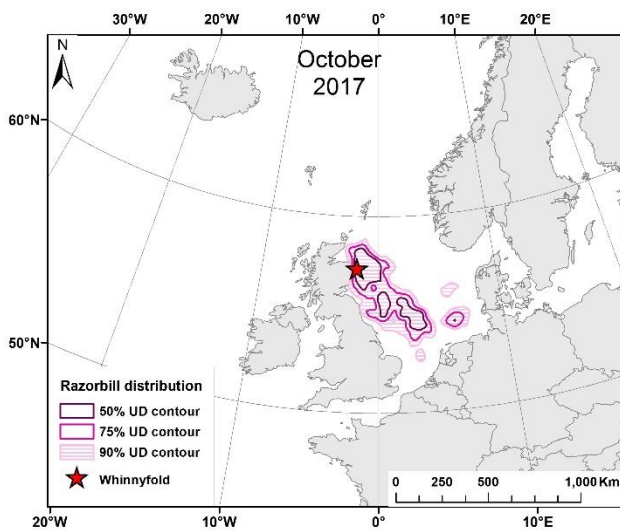
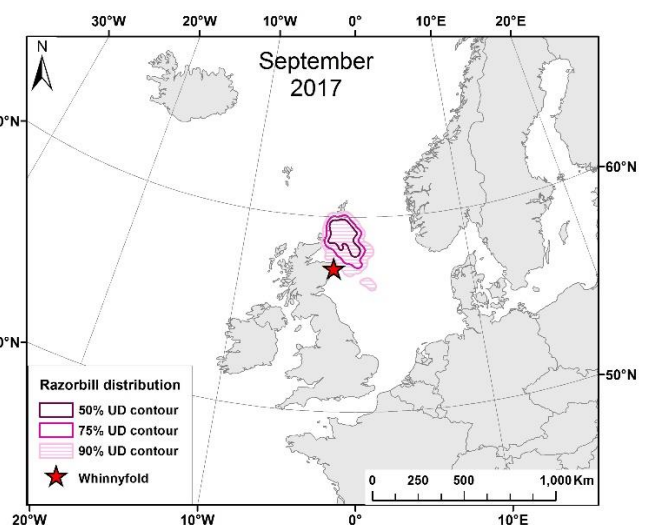
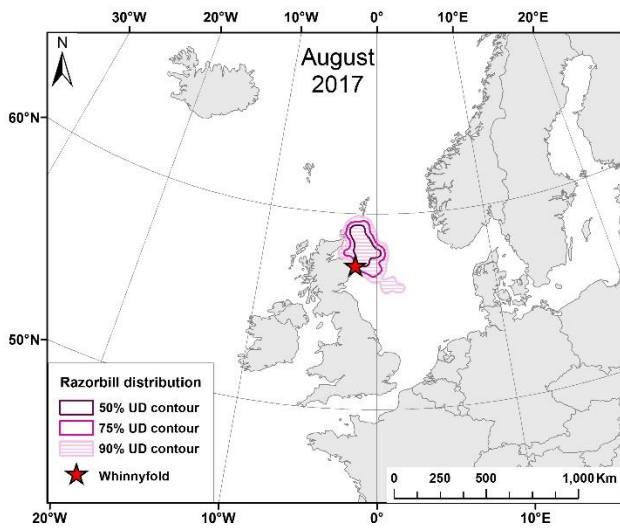


Fig. 9: Consistency in core areas (50% UD contours) of razorbills from East Caithness (n=15) in winter 2017-2018 (August – March).

Results for Buchan Ness to Collieston Coast in 2017 are based on data from few individuals and are likely to be less representative of the population's distribution compared to the other colonies. Similar to East Caithness razorbills, the birds from Buchan Ness to Collieston Coast stayed relatively close to the colony in the post-breeding period (Fig. 10a,b). In October, some of the birds were still in the area around the colony but others moved southward to the central North Sea and the Dogger Bank area (Fig. 10c). During the winter months core areas were located mainly in the central and southern North Sea, with a single bird migrating to the Skagerrak/Kattegat region (Fig. 10d-g). By March core areas were concentrated around the colony and in the central North Sea (Fig. 10h).

Over the whole non-breeding period, two key areas were apparent – one off north-east Scotland (including the area around the colony) and the other in the central North Sea (Fig. 10i). Overlap in individual core areas was highest near the colony (Fig. 11).

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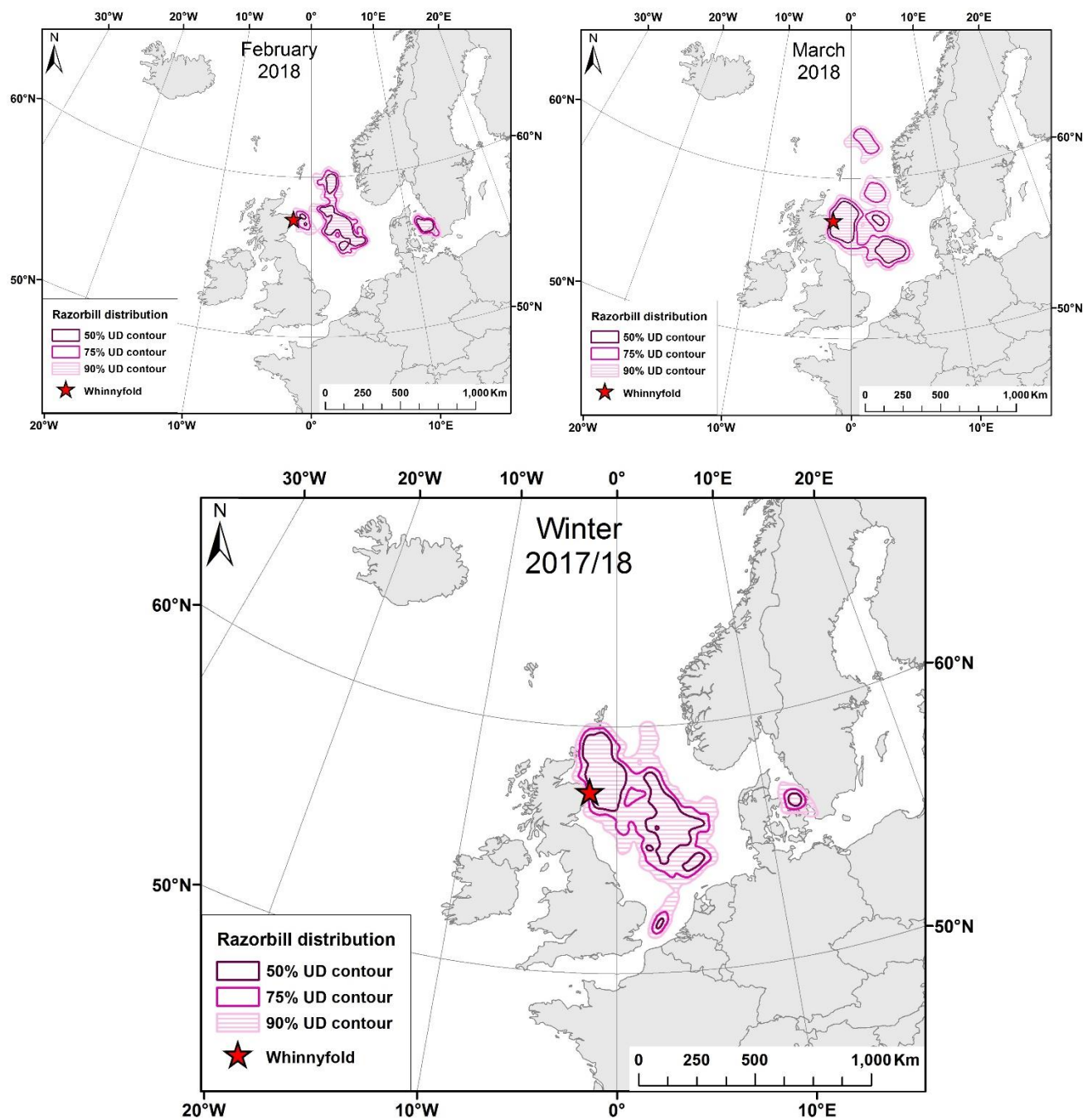


Fig. 10: Utilisation distributions (50%, 75%, 90% contours) for razorbills from Buchan Ness to Collieston Coast ($n=5$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

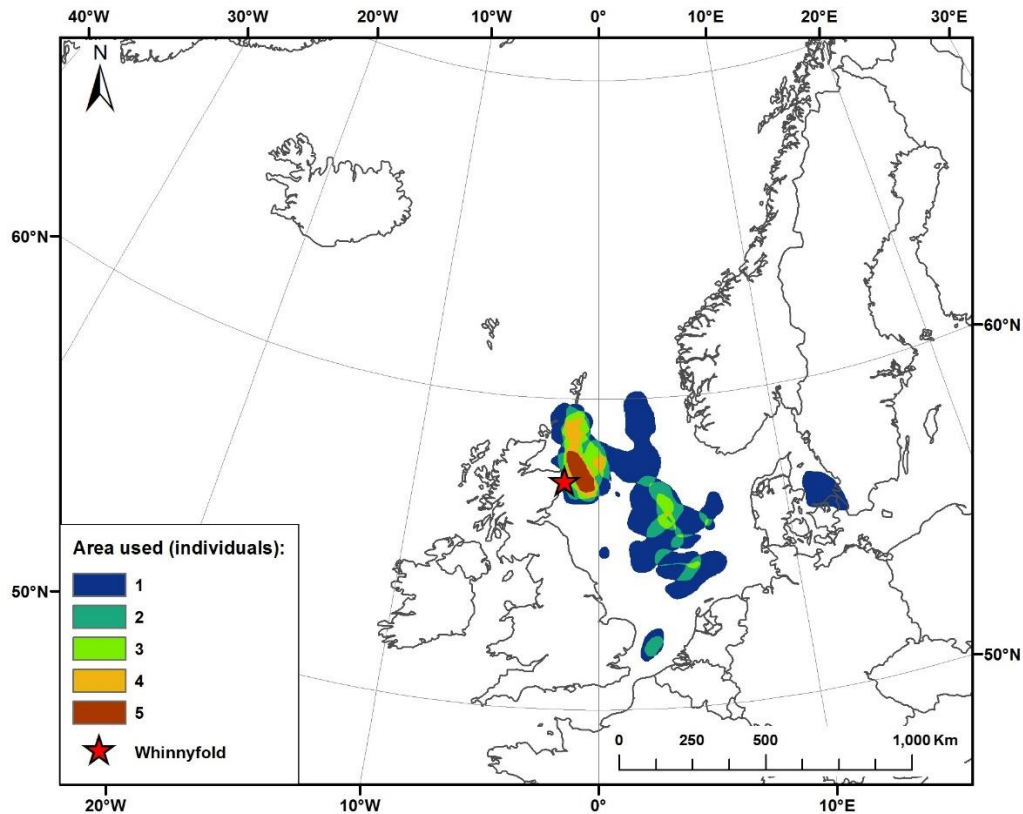
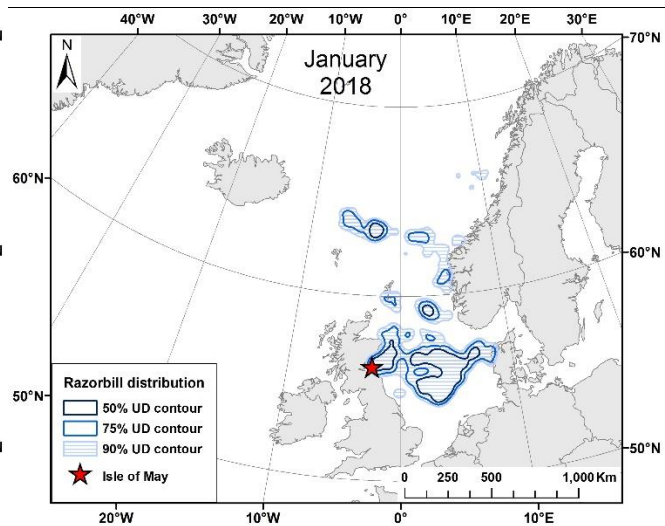
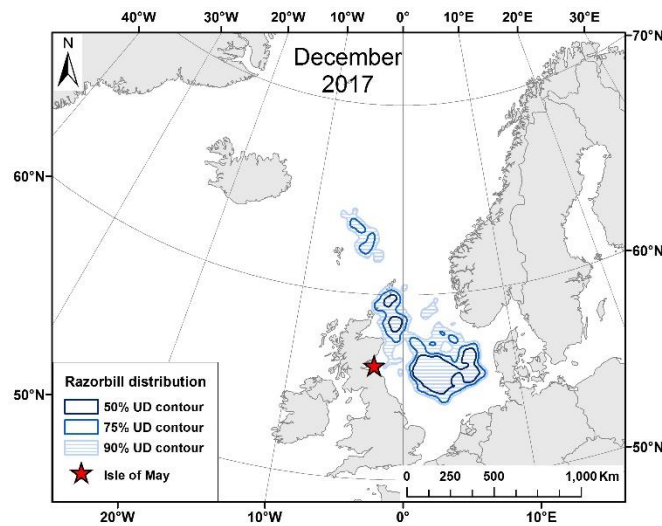
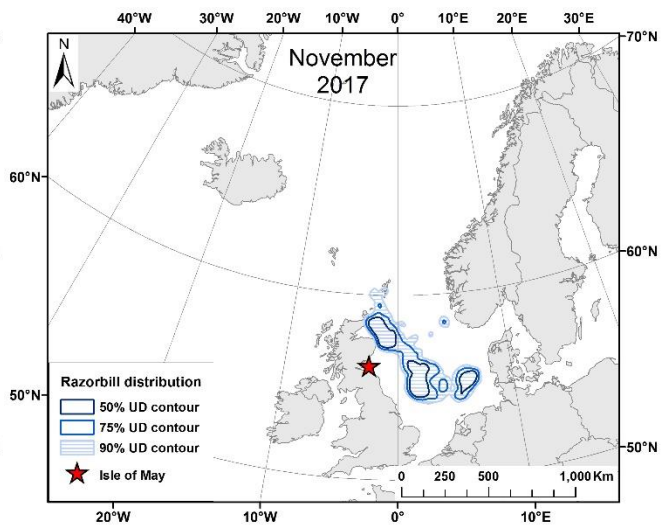
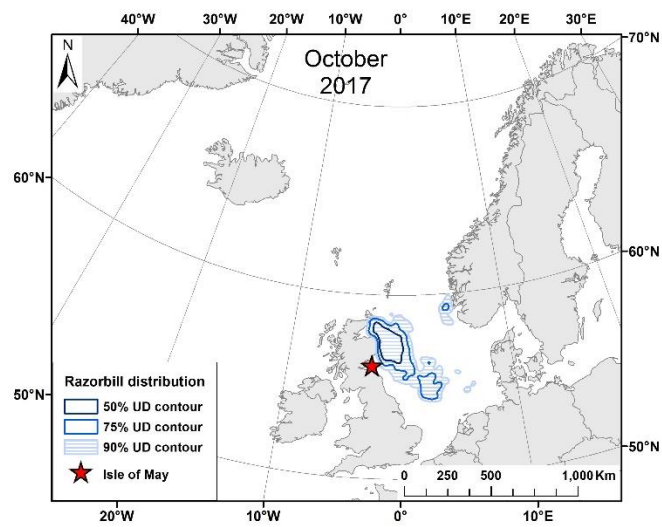
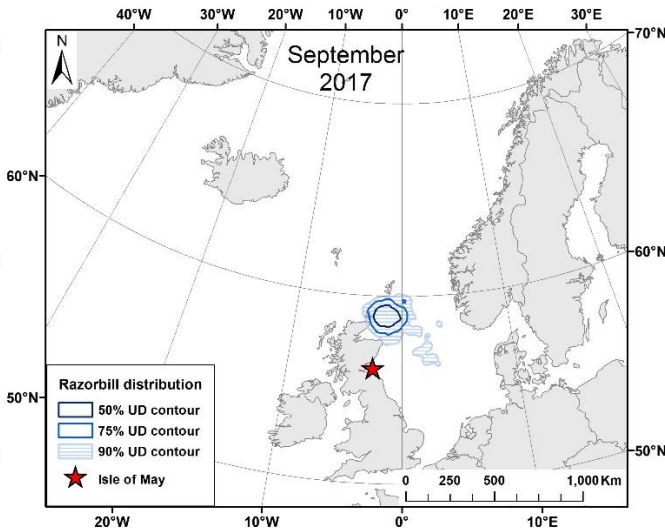
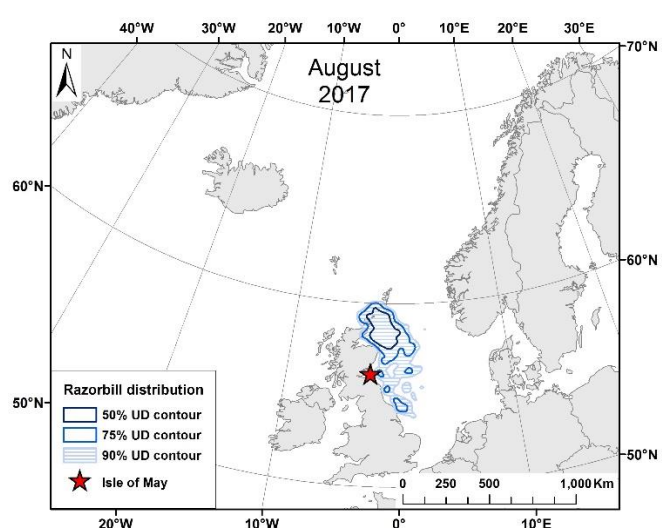


Fig. 11: Consistency in core areas (50% UD contours) of razorbills from Buchan Ness to Collieston Coast ($n=5$) in winter 2017-2018 (August – March).

After the breeding season, razorbills from the Isle of May stayed in the waters off north-east and east Scotland until November (Fig. 12a-c). In November and December core areas were located mainly in the central and northern North Sea (Fig. 12d,e). In January and February core areas were located around the colony, in the central North Sea and as far north as the Norwegian Sea (although the latter was driven by a single bird; Fig. 12f,g). In March, core areas were concentrated almost exclusively around the colony as the birds returned to breed (Fig. 12h).

Over the whole non-breeding period, razorbills from the Isle of May were distributed across the central and northern North Sea but key areas were located in the waters off east and north-east Scotland and in the Dogger Bank area (Fig. 12i). Consistency in individual core areas was highest around the Scottish coast, with 8-10 out of 11 birds overlapping (Fig. 13).

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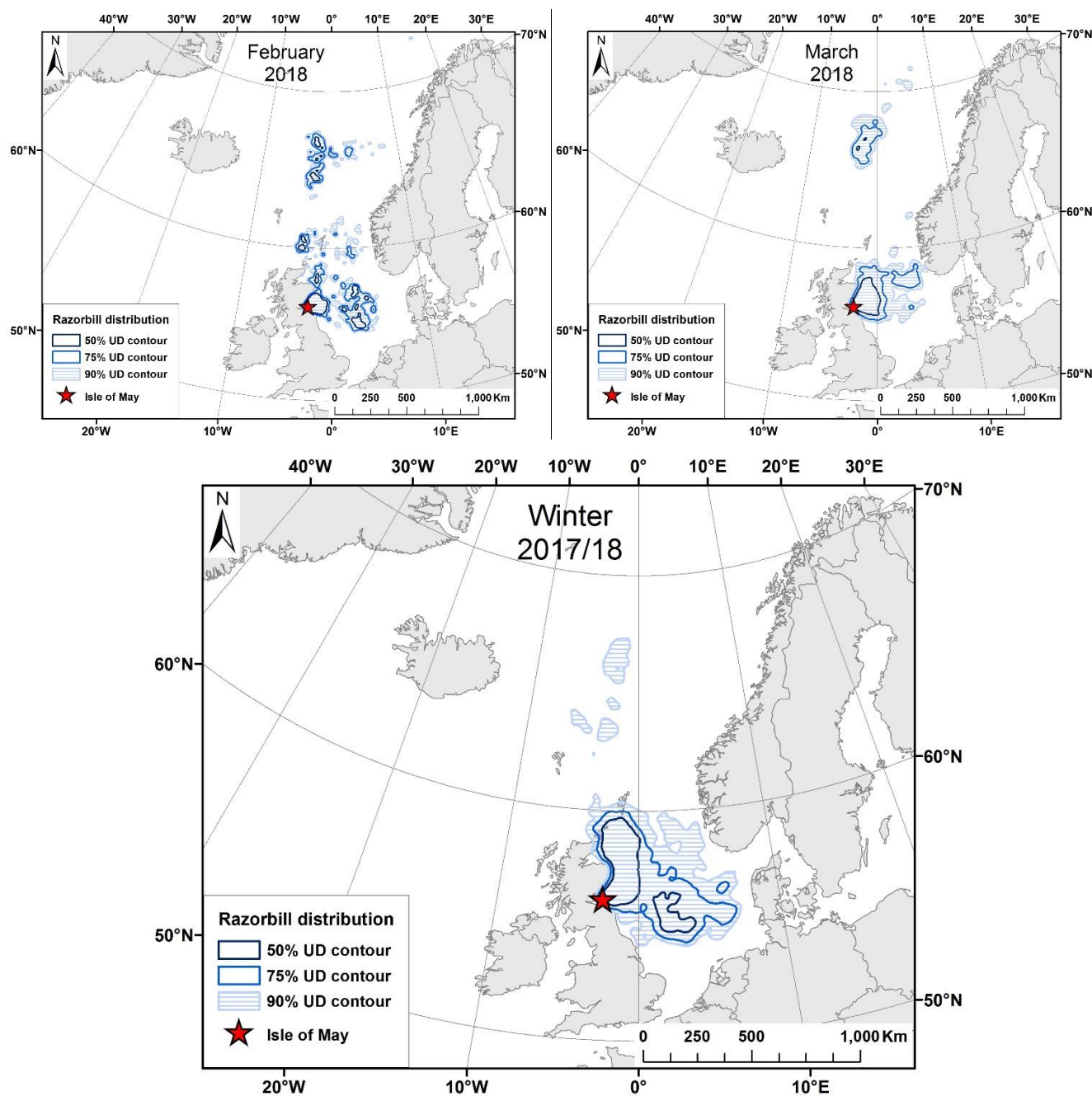


Fig. 12: Utilisation distributions (50%, 75%, 90% contours) for razorbills from Isle of May ($n=11$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2017-2018 (August – March).

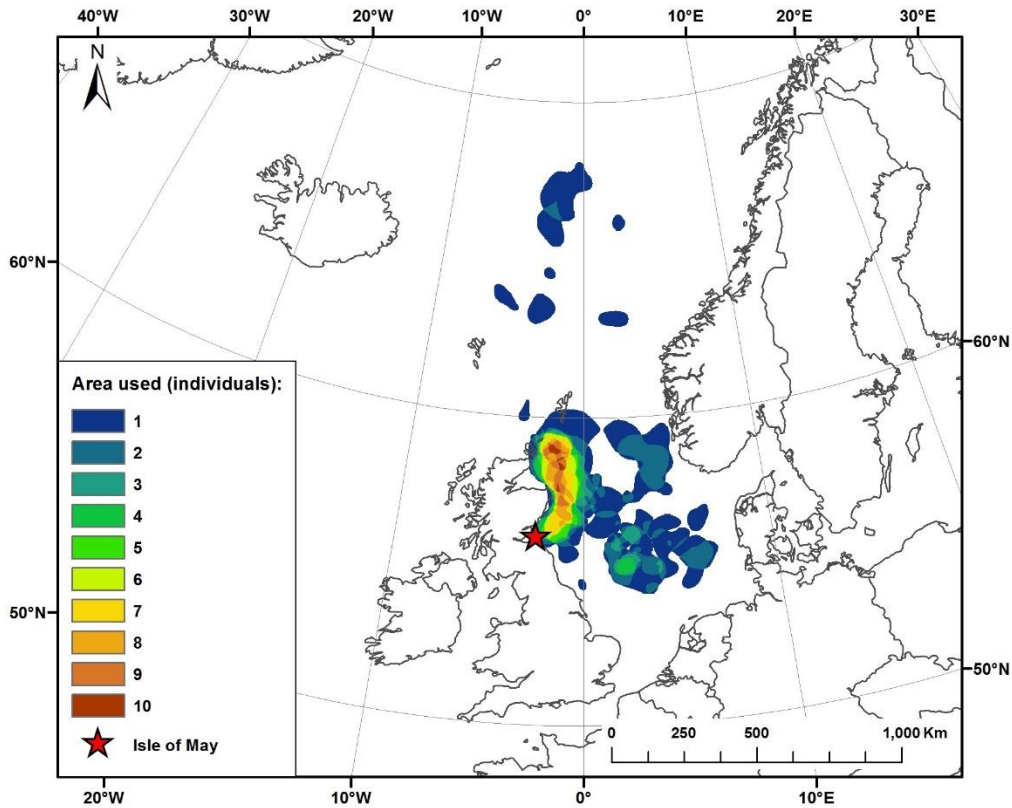


Fig. 13: Consistency in core areas (50% UD contours) of razorbills from Isle of May ($n=11$) in winter 2017-2018 (August – March).

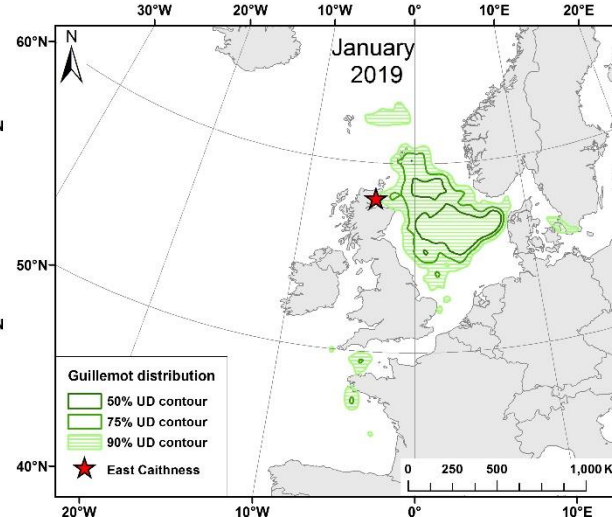
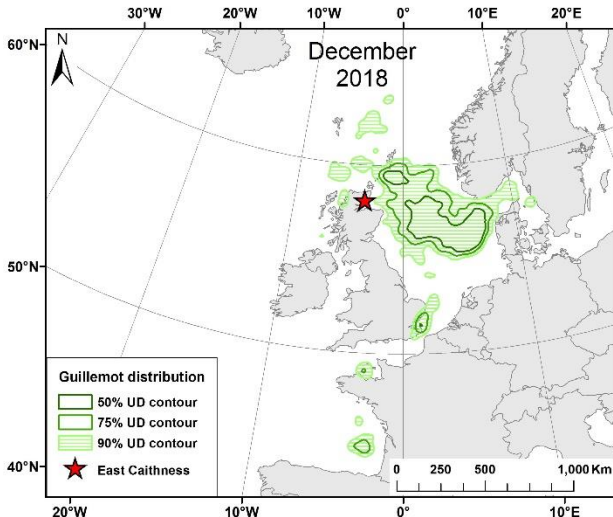
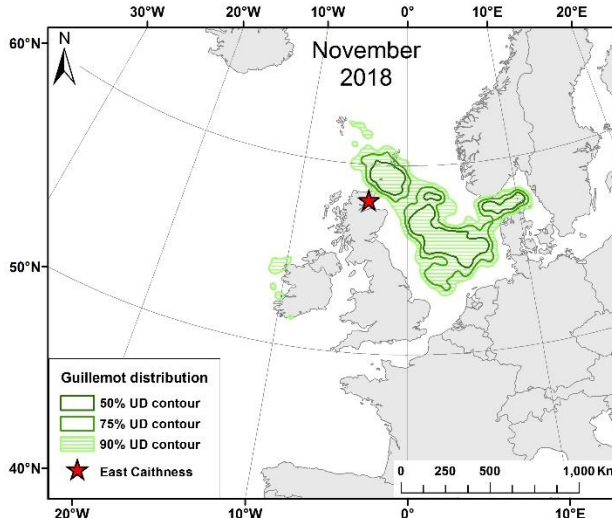
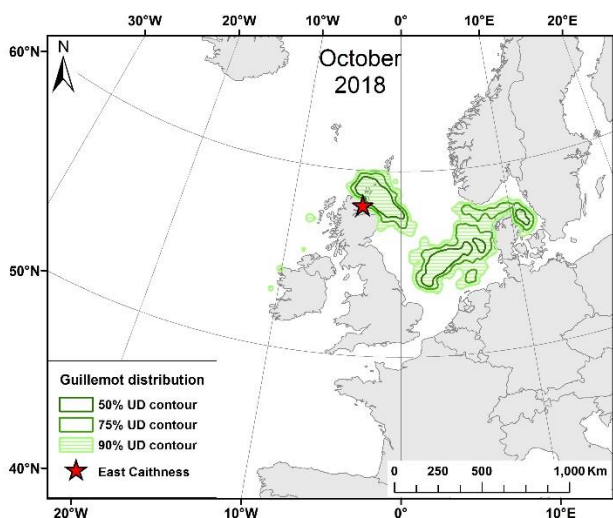
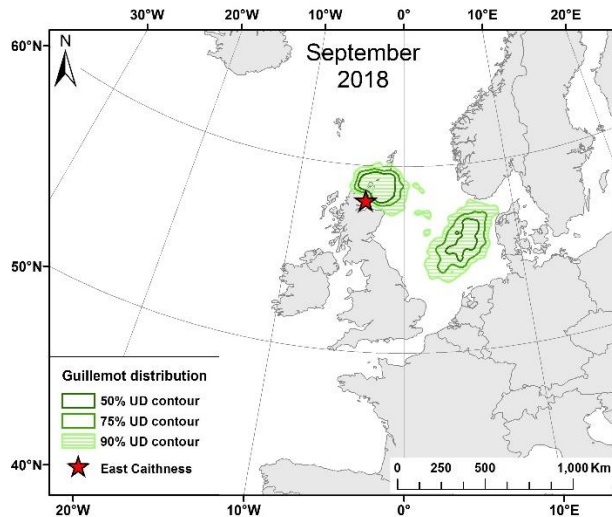
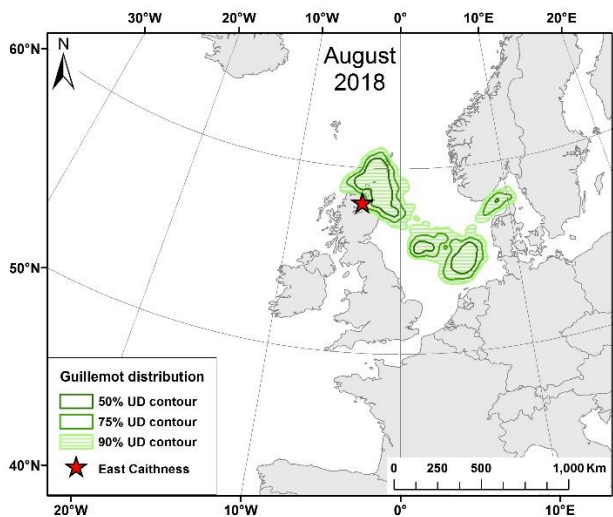
3.1.1.2 2018-19

Maps of utilisation distributions of guillemots in the second study year are provided in Figures 14-19. Corresponding maps for razorbills are presented in Figures 20-25.

Both the monthly and overall winter distributions of guillemots from East Caithness in 2018-19 were similar to those observed in 2017-18 (Fig. 2 and 14). The only notable differences were the more extensive use of the Skagerrak/Kattegat area in the second year and the intriguing migration of a single bird past the west coast of Ireland to the Celtic Sea, English Channel and Bay of Biscay (Fig. 14).

As in 2017-18, consistency in individual core areas was highest near the colony, with up to 13 out of 25 birds overlapping (Fig. 15). For comparison, key areas in the central North Sea were used by 7-10 out of 25 birds (Fig. 15).

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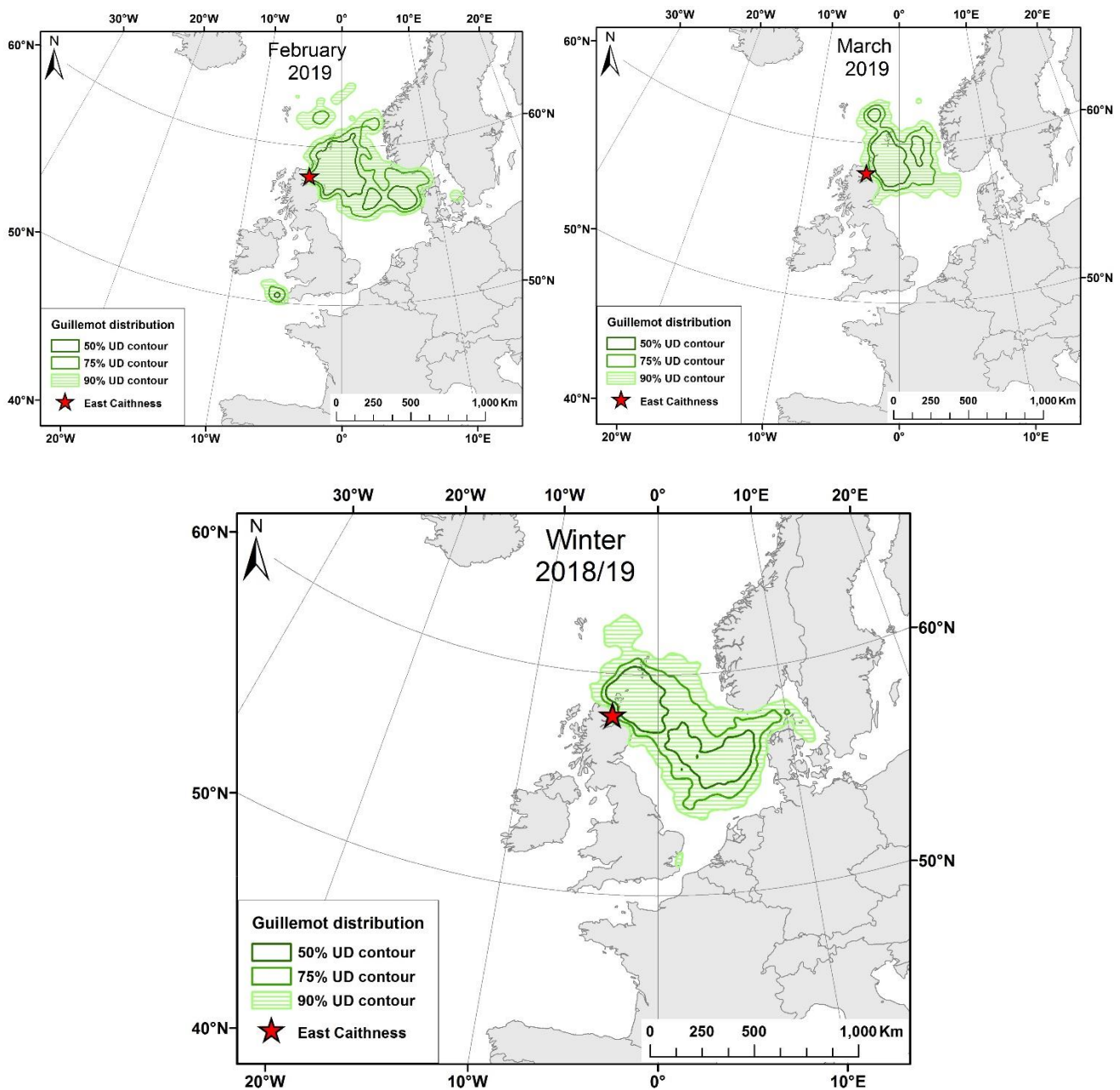


Fig. 14: Utilisation distributions (50%, 75%, 90% contours) for guillemots from East Caithness ($n=25$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

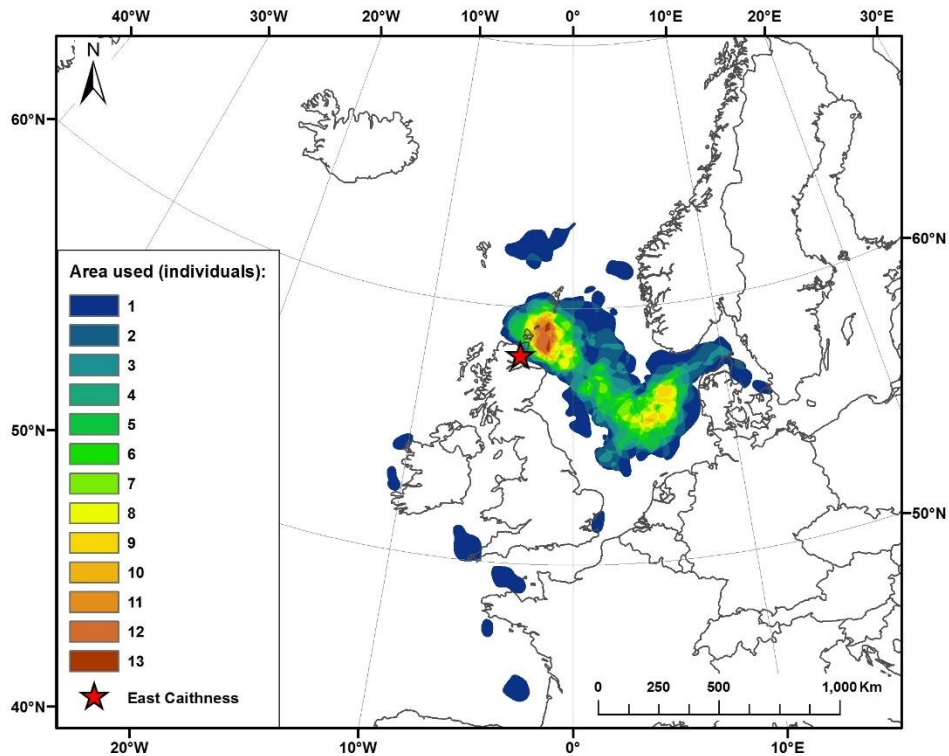
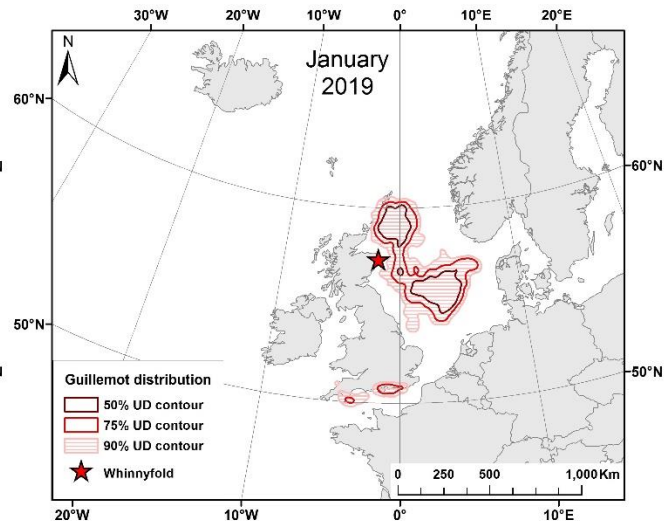
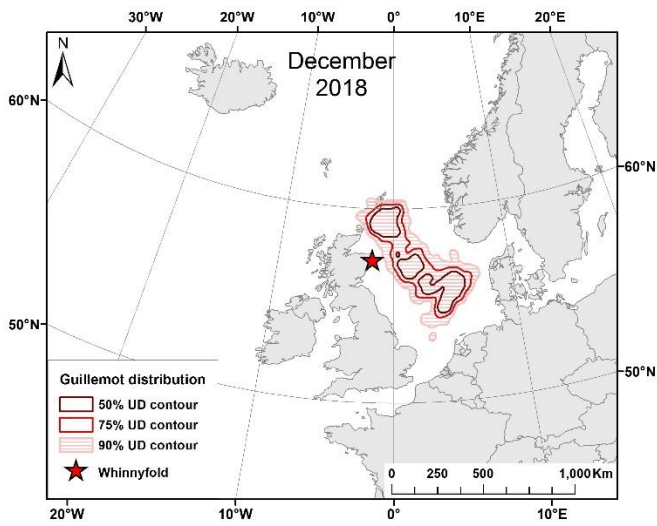
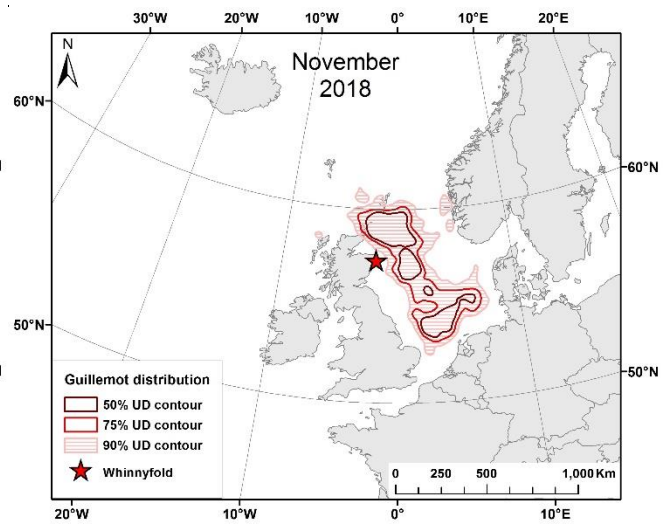
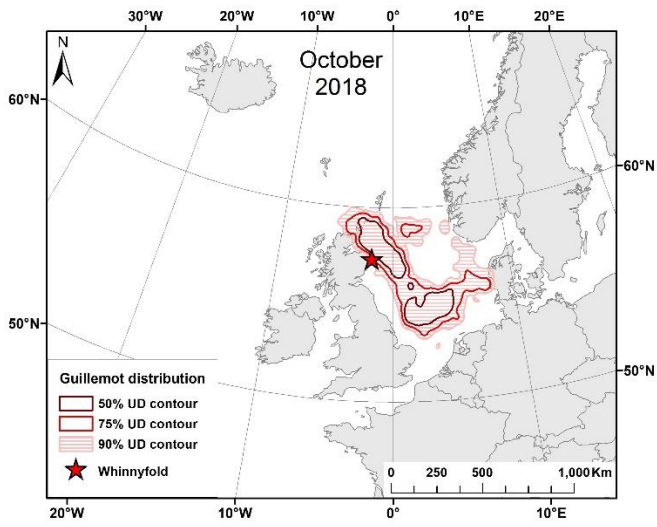
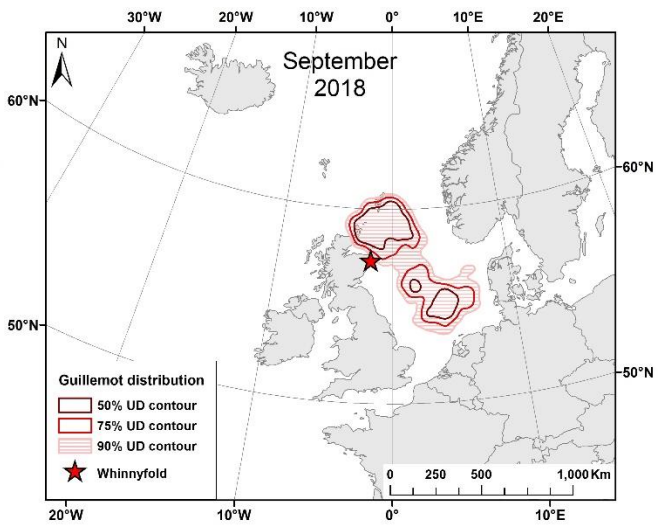
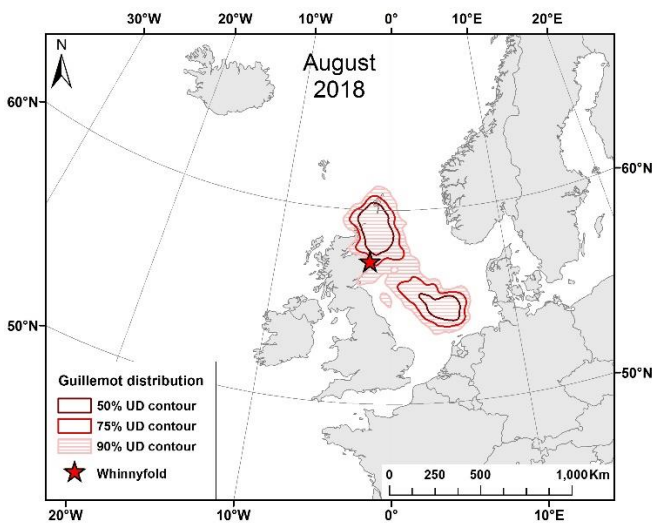


Fig. 15: Consistency in core areas (50% UD contours) of guillemots from East Caithness ($n=25$) in winter 2018-2019 (August – March).

As in East Caithness guillemots, both the monthly and overall winter distributions of guillemots from Buchan Ness to Collieston Coast in 2018-19 were similar to those in 2017-18 (Fig. 4 and 16). The only exception was the lack of movements towards the Norwegian Sea in February and March, resulting in a more restricted distribution in these months (Fig. 16).

As in 2017-18, consistency in individual core areas was highest near the colony, with up to 18 out of 26 birds overlapping (Fig. 17). However, the area of highest overlap was more restricted than in 2017-18. For comparison, areas in the central North Sea were used by 7-10 out of 26 birds (Fig. 17).

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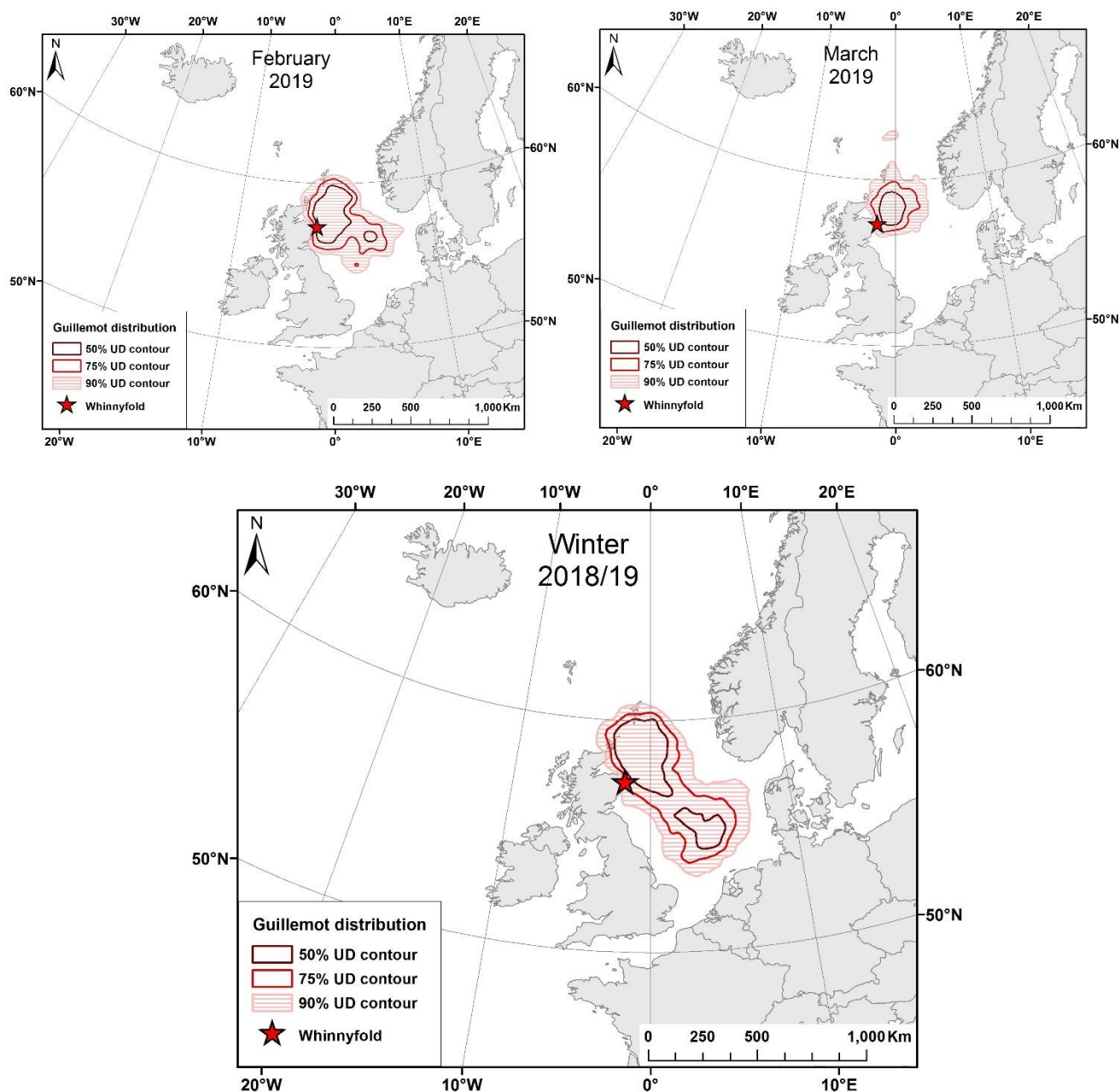


Fig. 16: Utilisation distributions (50%, 75%, 90% contours) for guillemots from Buchan Ness to Collieston Coast ($n=26$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

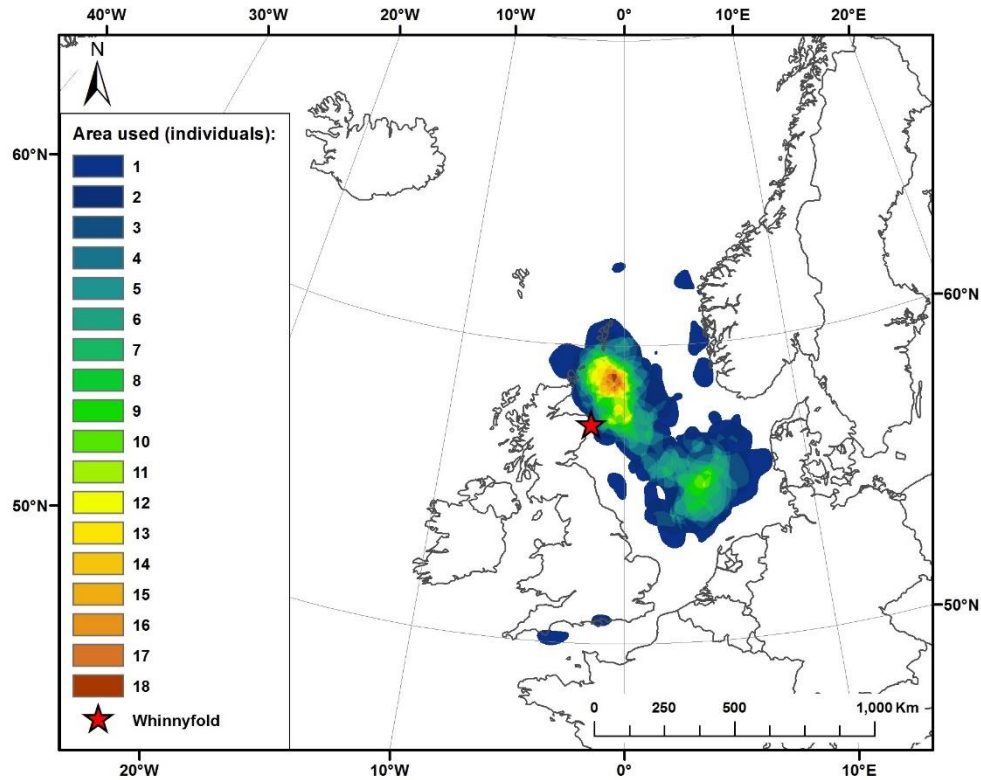
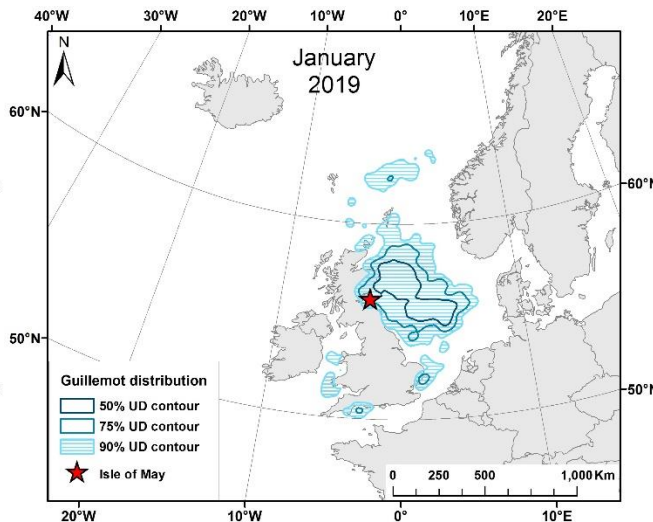
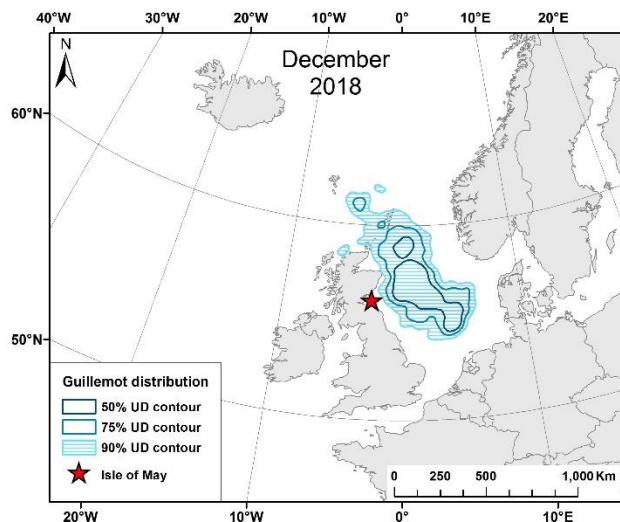
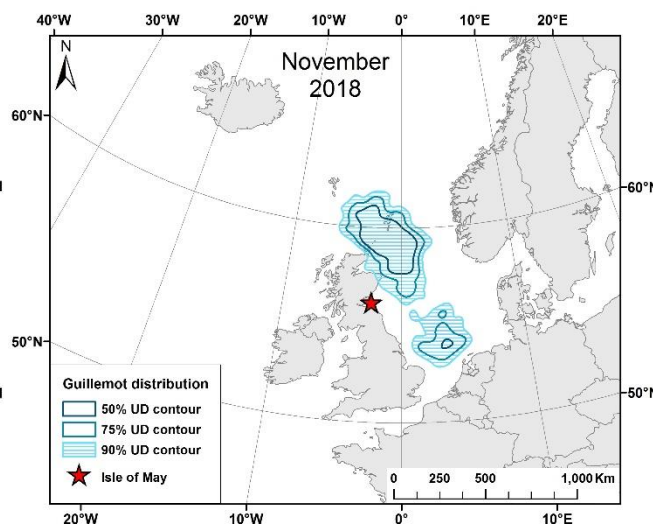
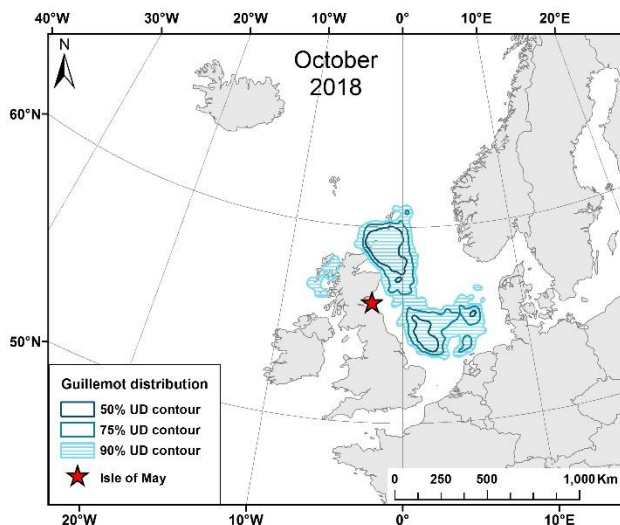
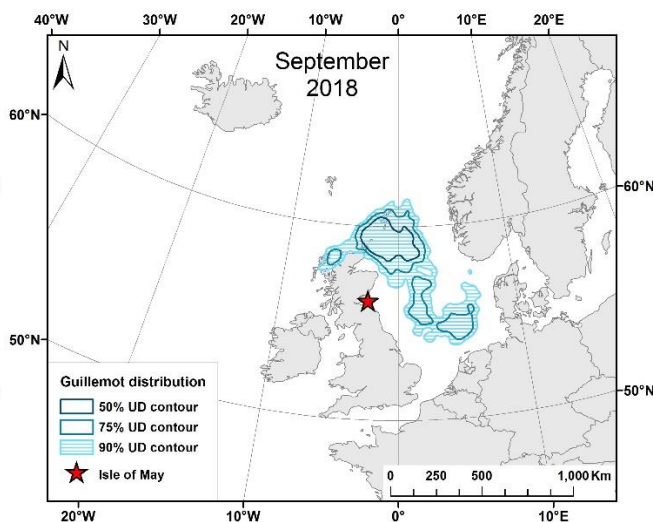
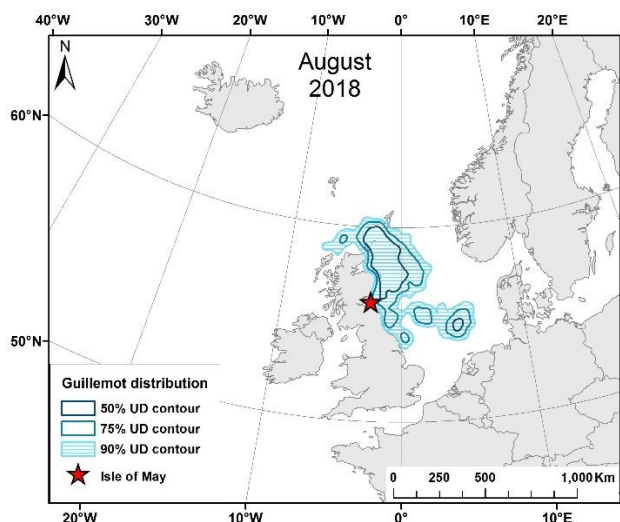


Fig. 17: Consistency in core areas (50% UD contours) of guillemots from Buchan Ness to Collieston Coast (n=26) in winter 2018-2019 (August – March).

As in the other colonies, the monthly and overall winter distributions of guillemots from Isle of May in 2018-19 were similar to those observed in 2017-18 (Fig. 6 and 18). The only notable differences were the absence of movements to the Skagerrak/Kattegat area in the second year and the less extensive use of the Norwegian Sea in February and March, resulting in a more restricted distribution in these months (Fig. 18).

As in 2017-18, consistency in individual core areas was highest in the waters off north-east Scotland, with up to 15 out of 22 birds overlapping (Fig. 19).



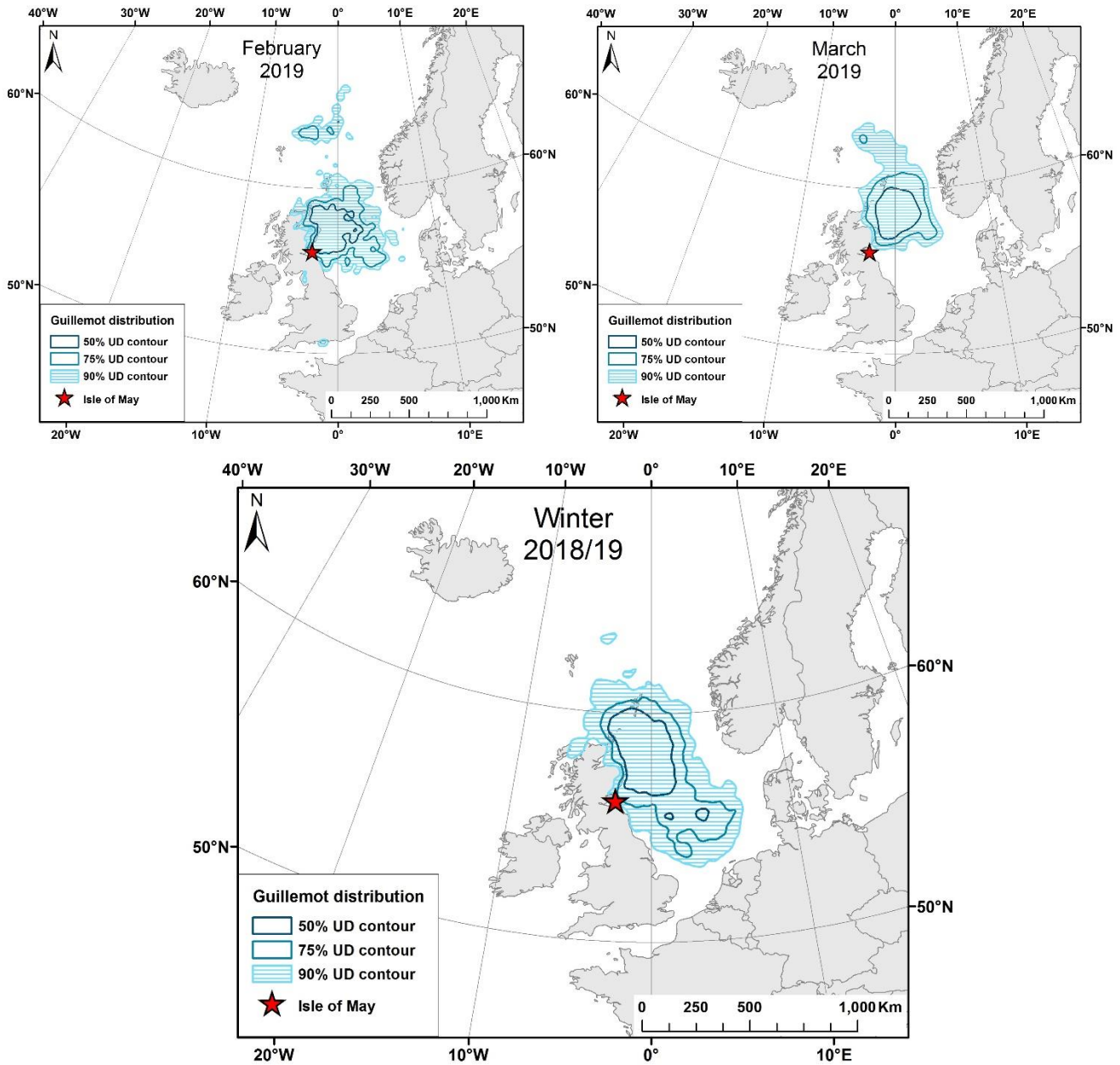


Fig. 18: Utilisation distributions (50%, 75%, 90% contours) for guillemots from Isle of May ($n=22$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

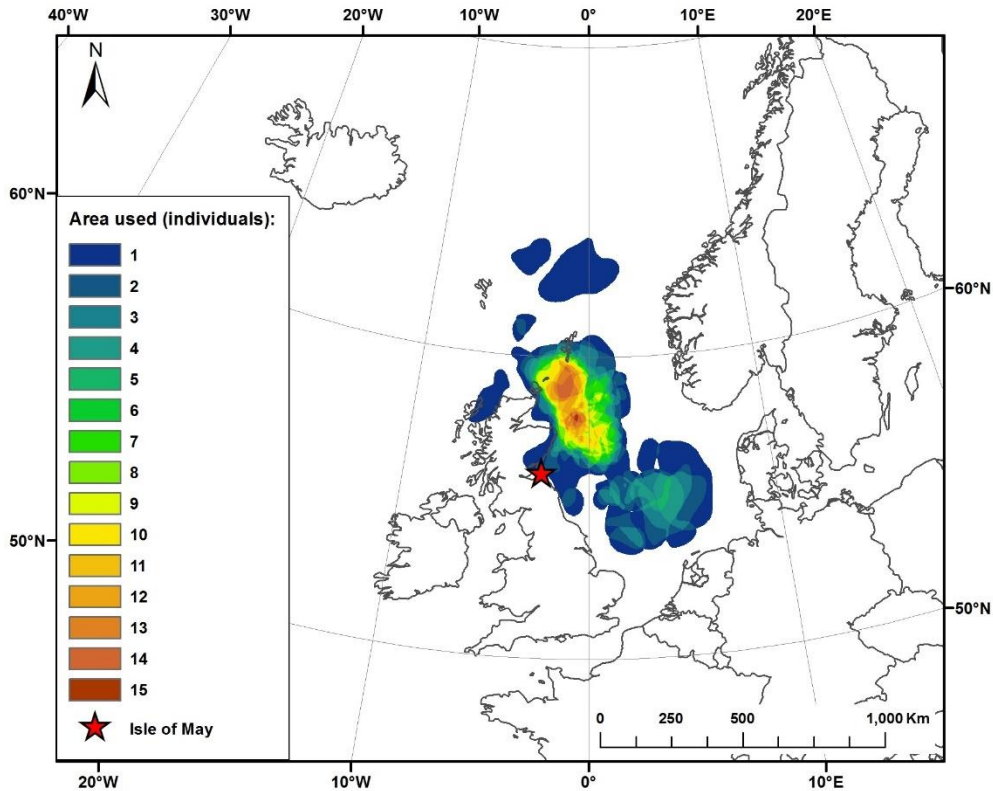
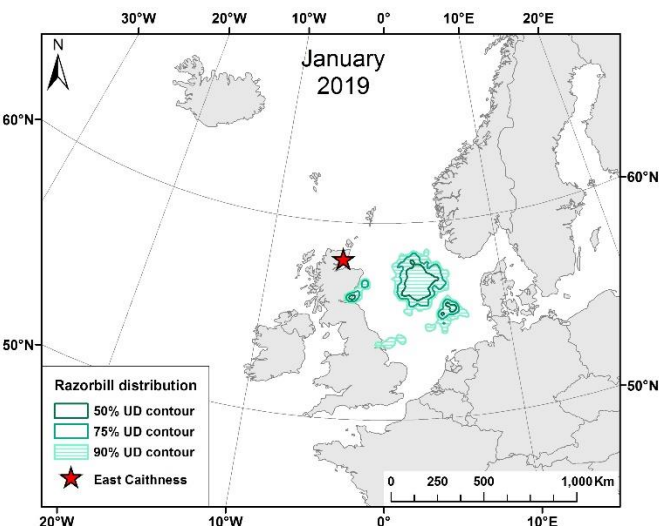
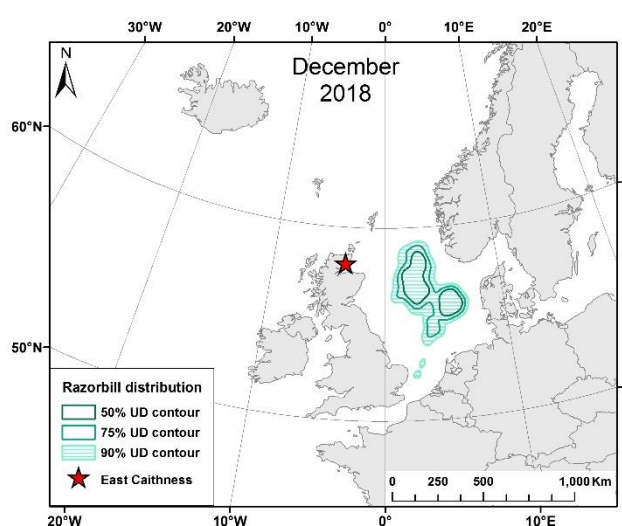
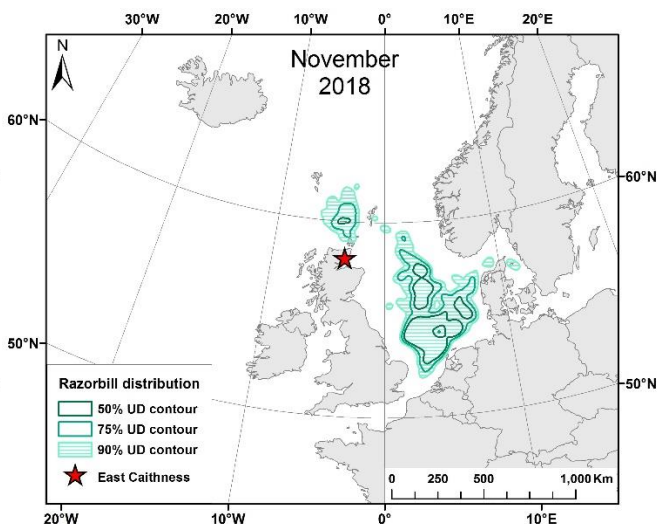
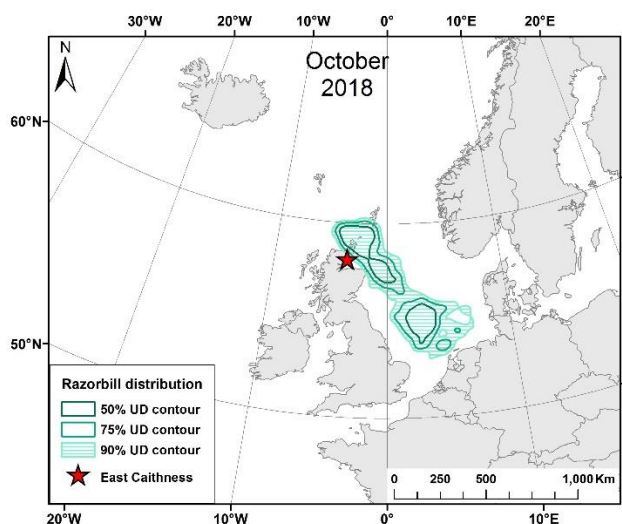
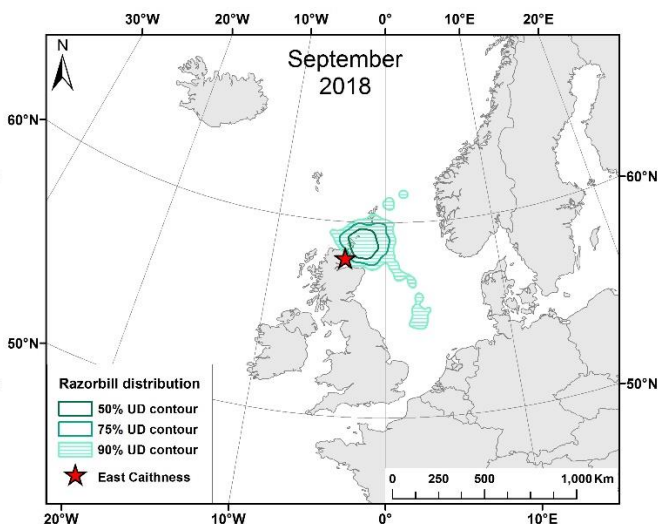
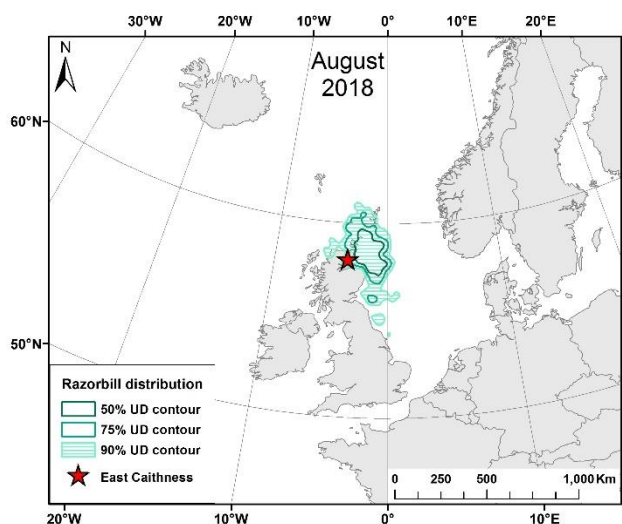


Fig. 19: Consistency in core areas (50% UD contours) of guillemots from Isle of May ($n=22$) in winter 2018-2019 (August – March).

The monthly and overall winter distributions of razorbills from East Caithness in 2018-19 were generally consistent with those observed in 2017-18 (Fig. 8 and 20). The distribution particularly in January to March was more restricted compared to the first year (Fig. 20), however this may be a consequence of the smaller sample size of birds in the second year.

Highest consistency in individual core areas was observed near the colony, with 5-7 out of 7 birds overlapping (Fig. 21). Consistency was less pronounced in the central parts of the North Sea (Fig. 21).

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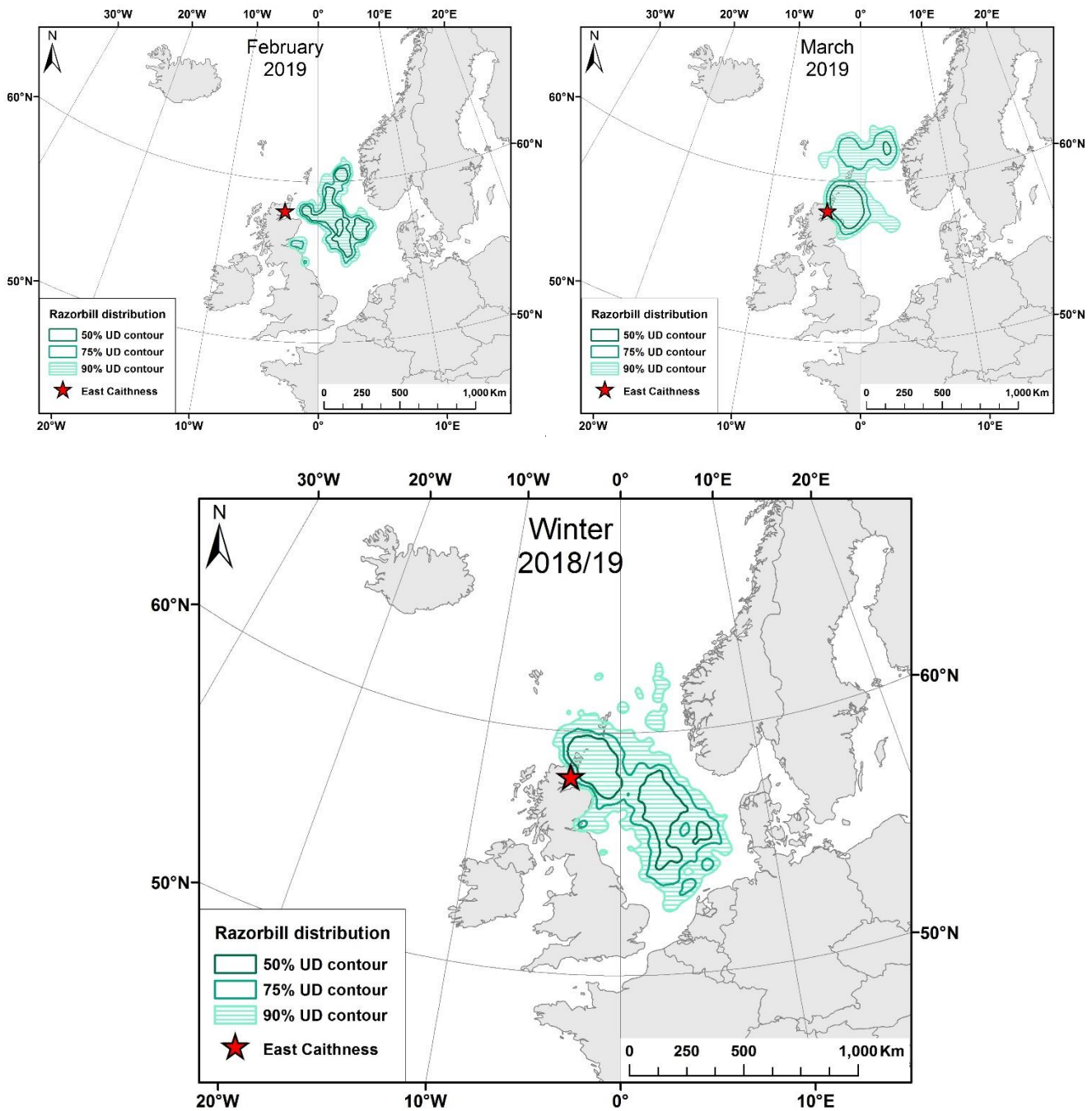


Fig. 20: Utilisation distributions (50%, 75%, 90% contours) for razorbills from East Caithness ($n=7$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

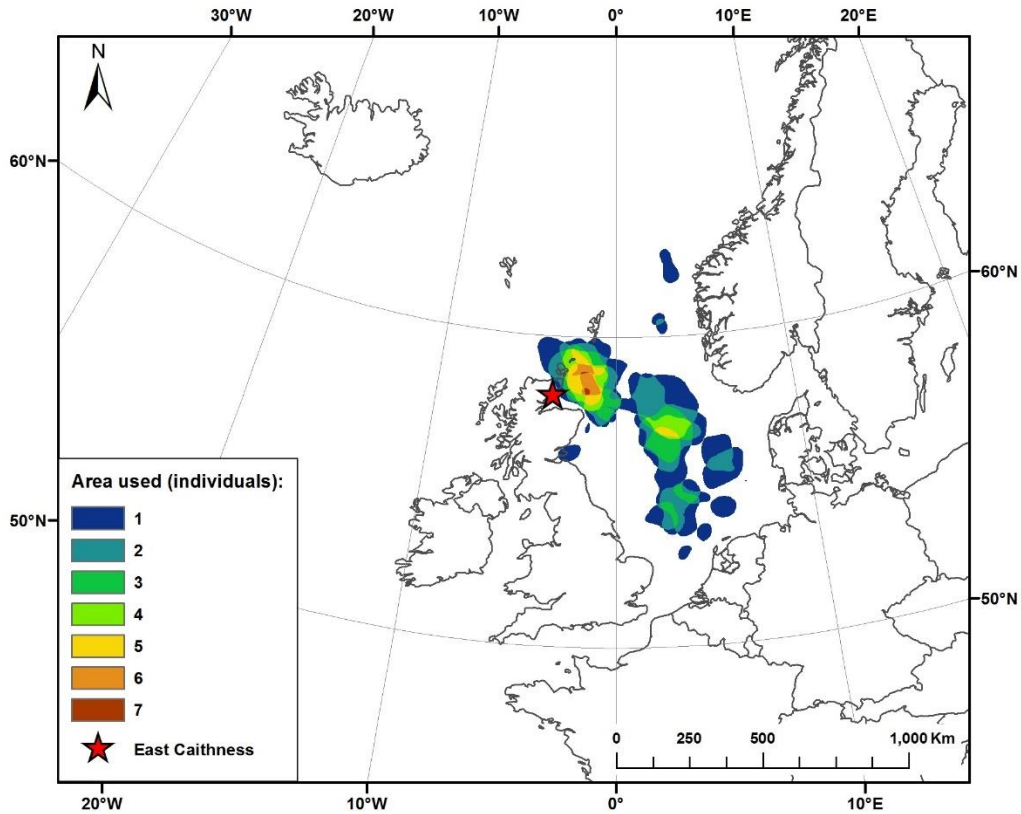
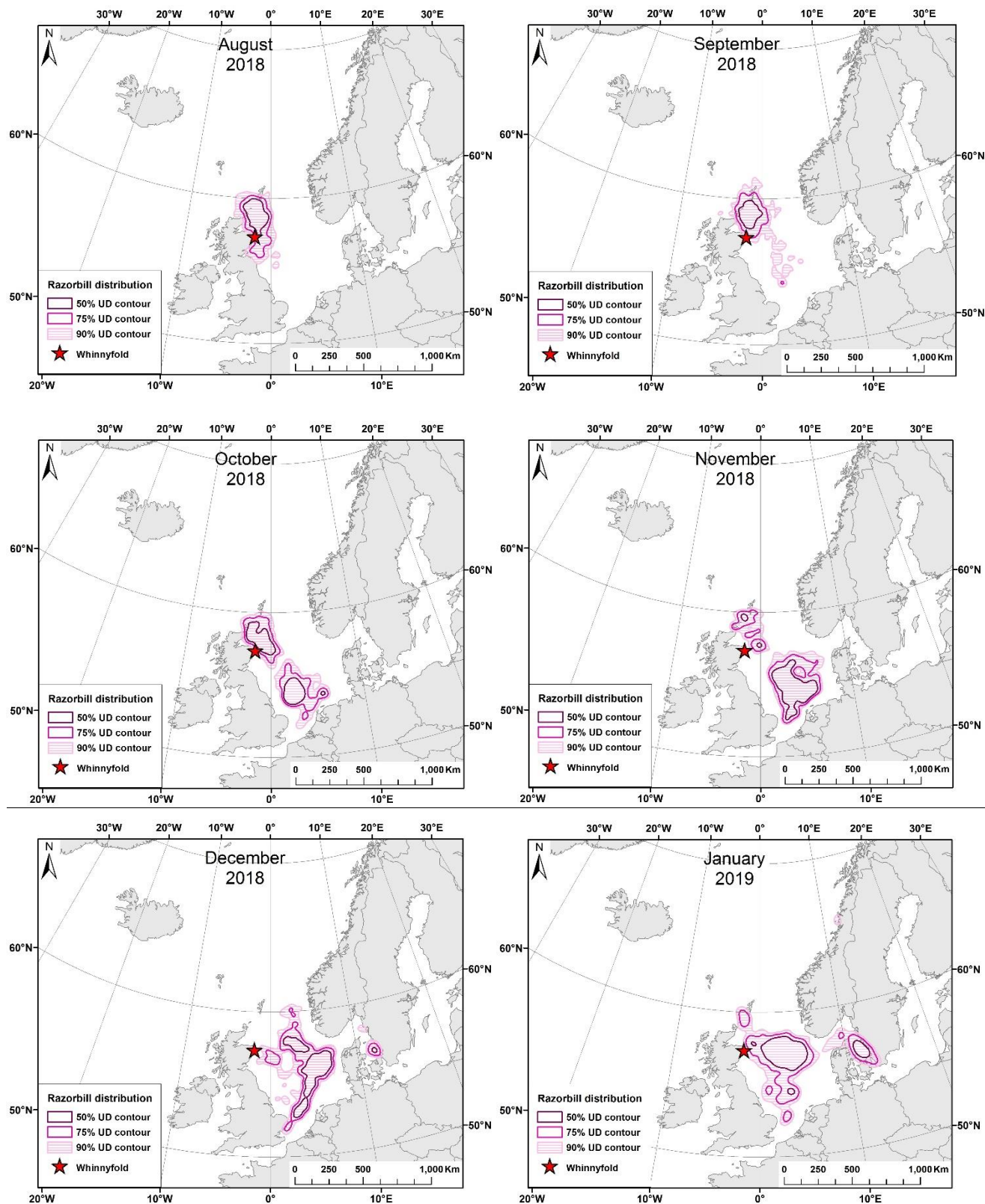


Fig. 21: Consistency in core areas (50% UD contours) of razorbills from East Caithness (n=7) in winter 2018-2019 (August – March).

The sample size of razorbills tracked at Buchan Ness to Collieston Coast in 2018-19 was slightly higher than in 2017-18 but remains relatively small and the results should therefore be treated with caution. Comparisons of the two years showed that the monthly and overall winter distributions were generally consistent (Fig. 10 and 22). Interestingly, a single individual undertook a movement along the coast of Norway, however there is no evidence it reached the Barents Sea as an Isle of May guillemot has previously done.

As in 2017-18, consistency in individual core areas was highest near the colony, with all 7 birds overlapping (Fig. 23).

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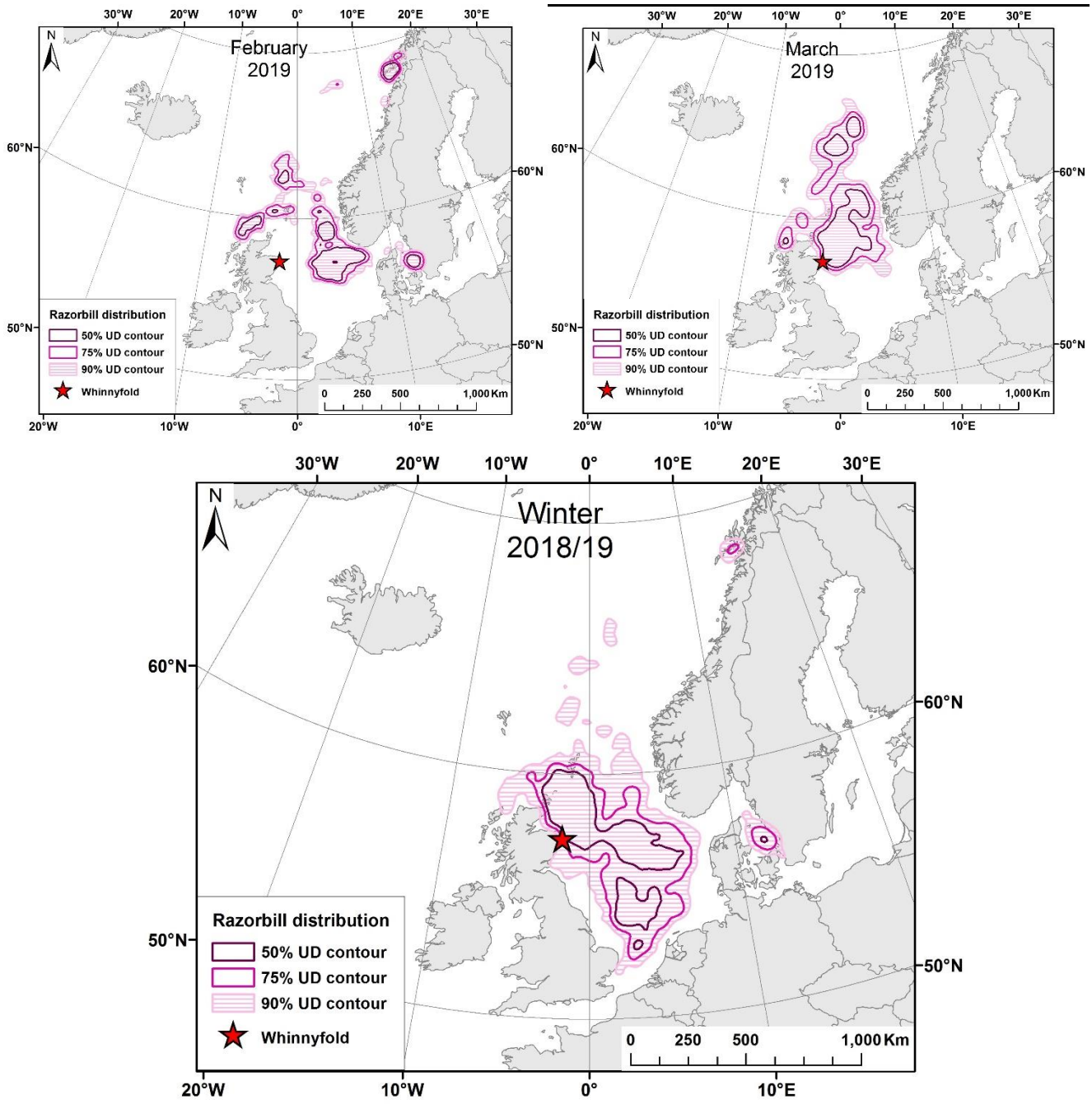


Fig. 22: Utilisation distributions (50%, 75%, 90% contours) for razorbills from Buchan Ness to Collieston Coast (n=8) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

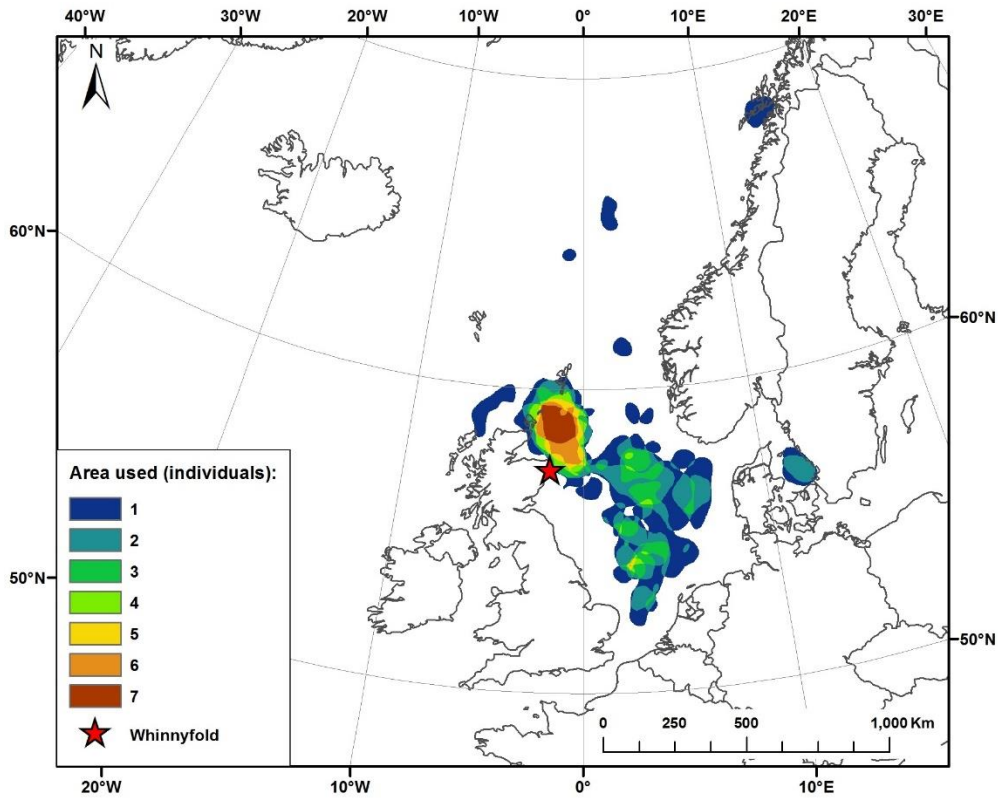
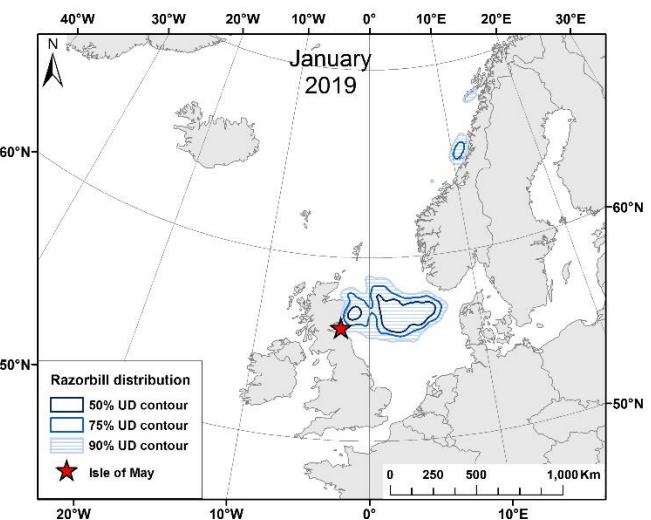
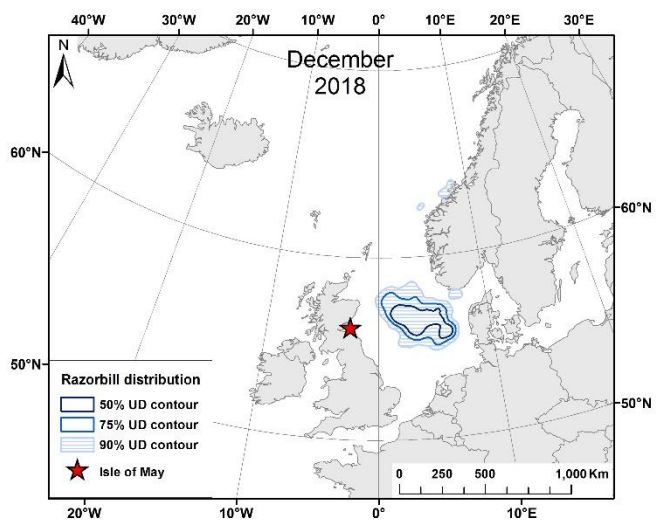
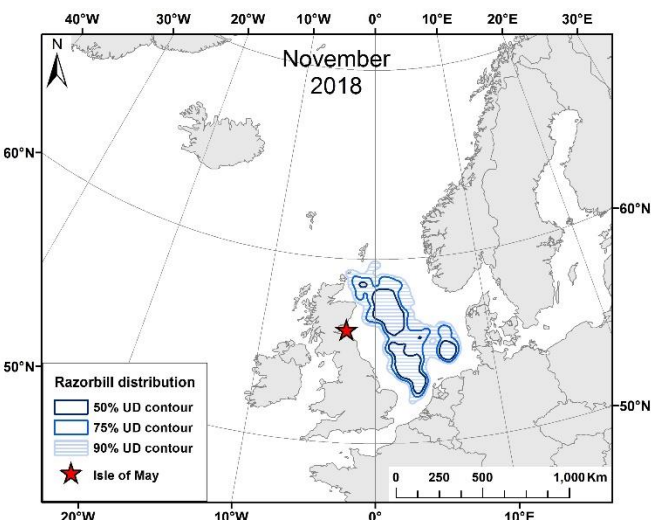
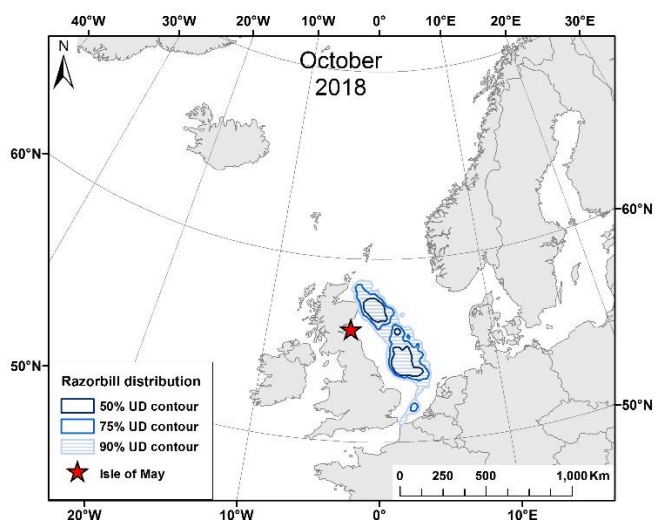
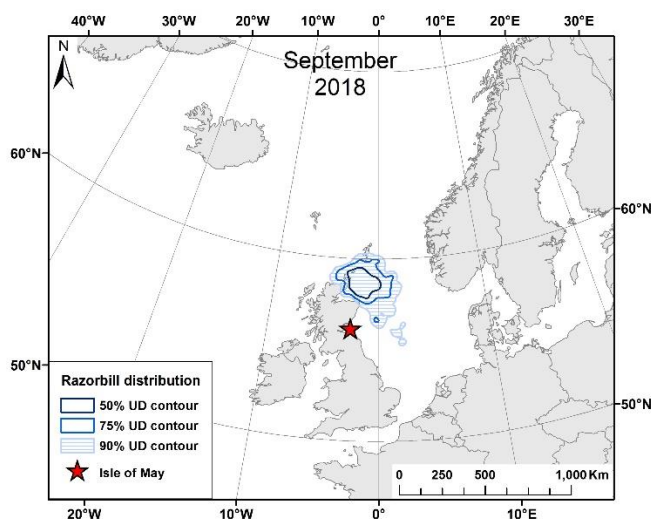
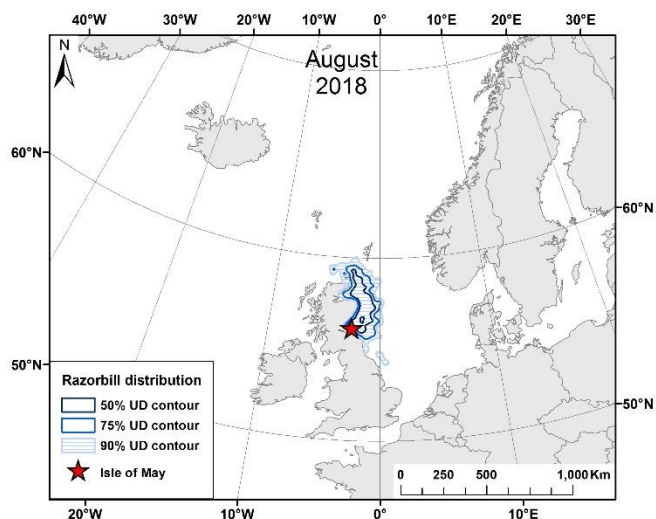


Fig. 23: Consistency in core areas (50% UD contours) of razorbills from Buchan Ness to Collieston Coast (n=8) in winter 2018-2019 (August – March).

The monthly and overall winter distributions of razorbills from Isle of May in 2018-19 were generally consistent with those observed in 2017-18 (Fig. 12 and 24). Similar to the case at Buchan Ness to Collieston Coast, a single bird undertook a movement along the coast of Norway.

Highest consistency in individual core areas was observed in the waters off the east/north-east coast of Scotland, with up to all 11 birds overlapping (Fig. 25). Consistency was less pronounced in the central parts of the North Sea (Fig. 25).

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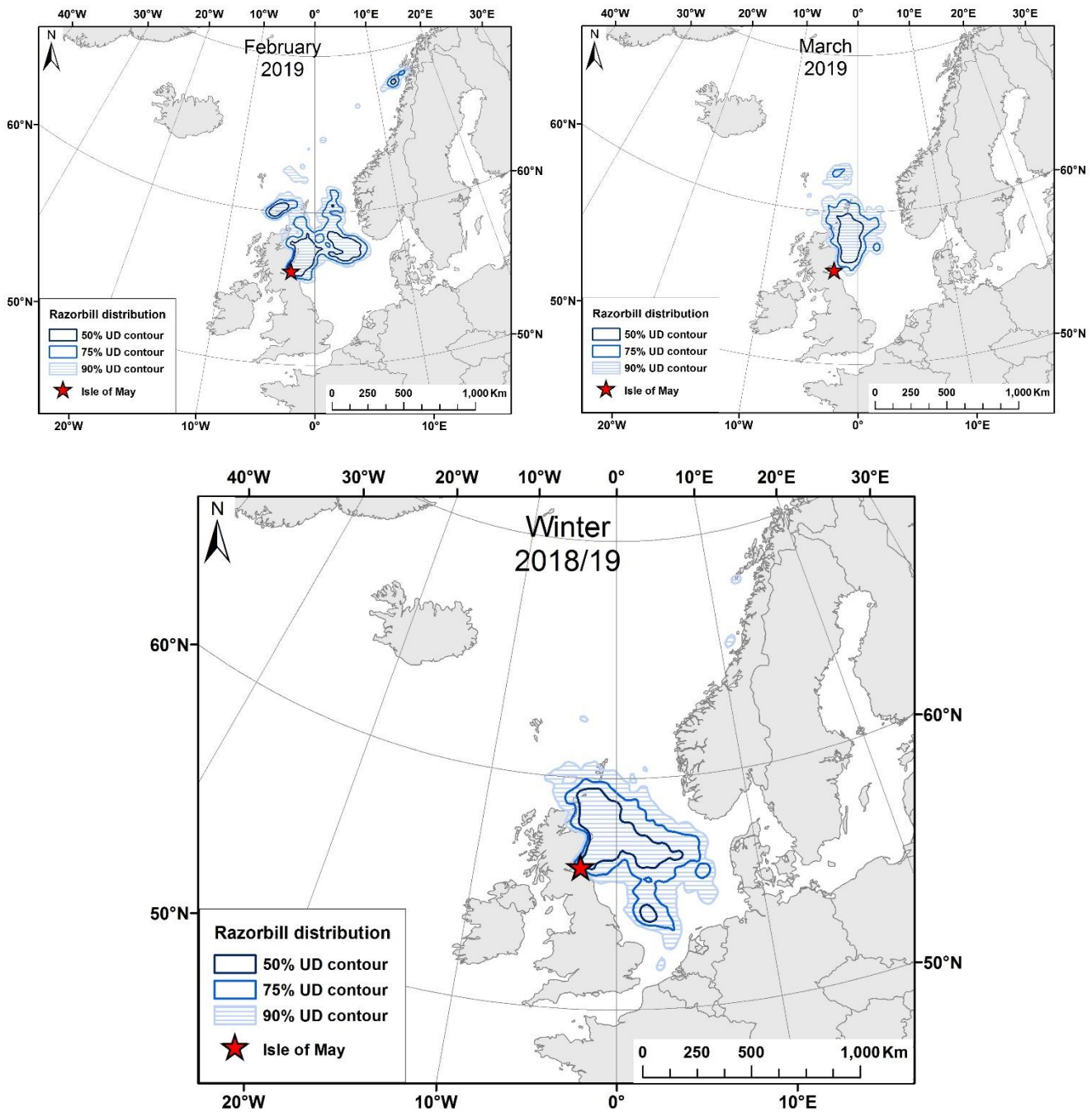


Fig. 24: Utilisation distributions (50%, 75%, 90% contours) for razorbills from Isle of May ($n=11$) in a) August, b) September, c) October, d) November, e) December, f) January, g) February, h) March and i) winter 2018-2019 (August – March).

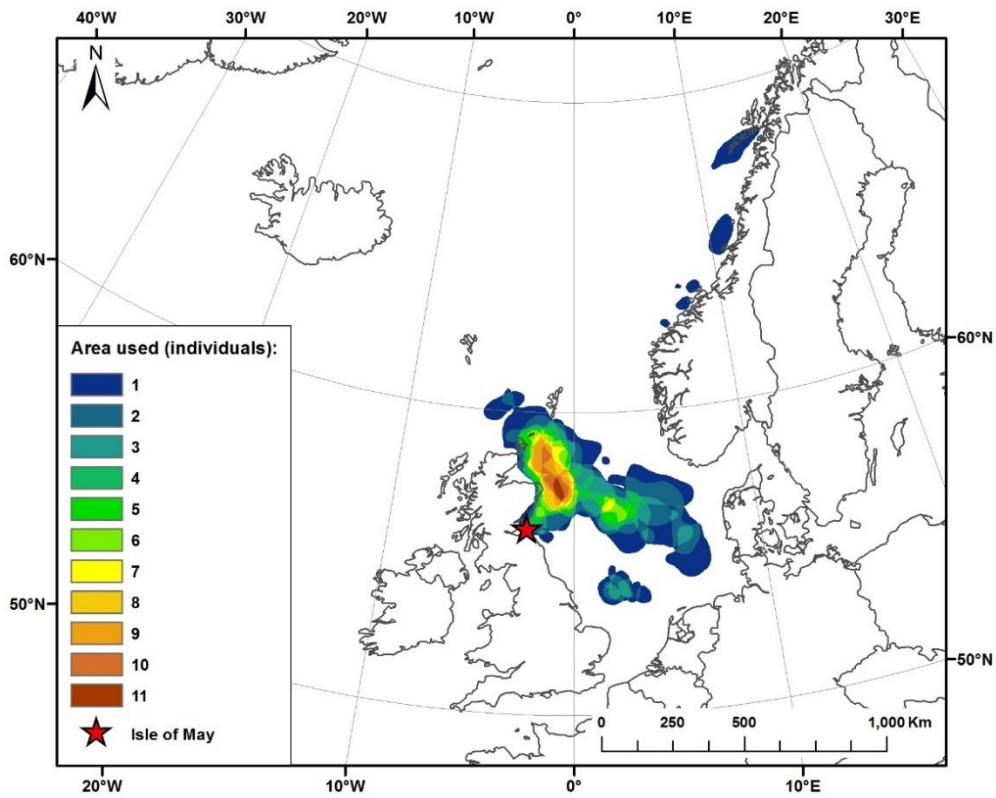


Fig. 25: Consistency in core areas (50% UD contours) of razorbills from Isle of May ($n=11$) in winter 2018-2019 (August – March).

3.1.2 Overlap in colony utilisation distributions

In both years, over the whole non-breeding period, spatial similarity in the UD of guillemots from the three study colonies was very high (Table 3).

| Colony | East Caithness | Buchan Ness to Collieston Coast | Isle of May |
|----------------|----------------|---------------------------------|-------------|
| a) 2017-18 | | | |
| East Caithness | | 0.95 | 0.86 |
| Buchan Ness | 0.95 | | 0.89 |
| Isle of May | 0.86 | 0.89 | |
| b) 2018-19 | | | |
| East Caithness | | 0.91 | 0.88 |
| Buchan Ness | 0.91 | | 0.95 |
| Isle of May | 0.88 | 0.95 | |

Table 3. Whole winter pairwise spatial similarity of UD of guillemots from the three study colonies in: a) 2017-18; b) 2018-19. Similarity was estimated using Bhattacharyya's affinity measure, ranging from zero (no similarity) to 1 (identical densities).

In 2017-18, pairwise comparisons indicated a moderate to high monthly spatial similarity in UD of guillemots from the three colonies (Fig. 26a). Throughout the non-breeding period, similarity between East Caithness and Buchan Ness to Collieston Coast was consistently high whereas similarity between these two colonies and the Isle of May was generally lower, particularly in November (Fig. 26a).

In 2018-19, monthly spatial similarity in UD was high among birds from all three colonies, although slightly lower between East Caithness and Isle of May compared to the other two pairs of colonies (Fig. 26b).

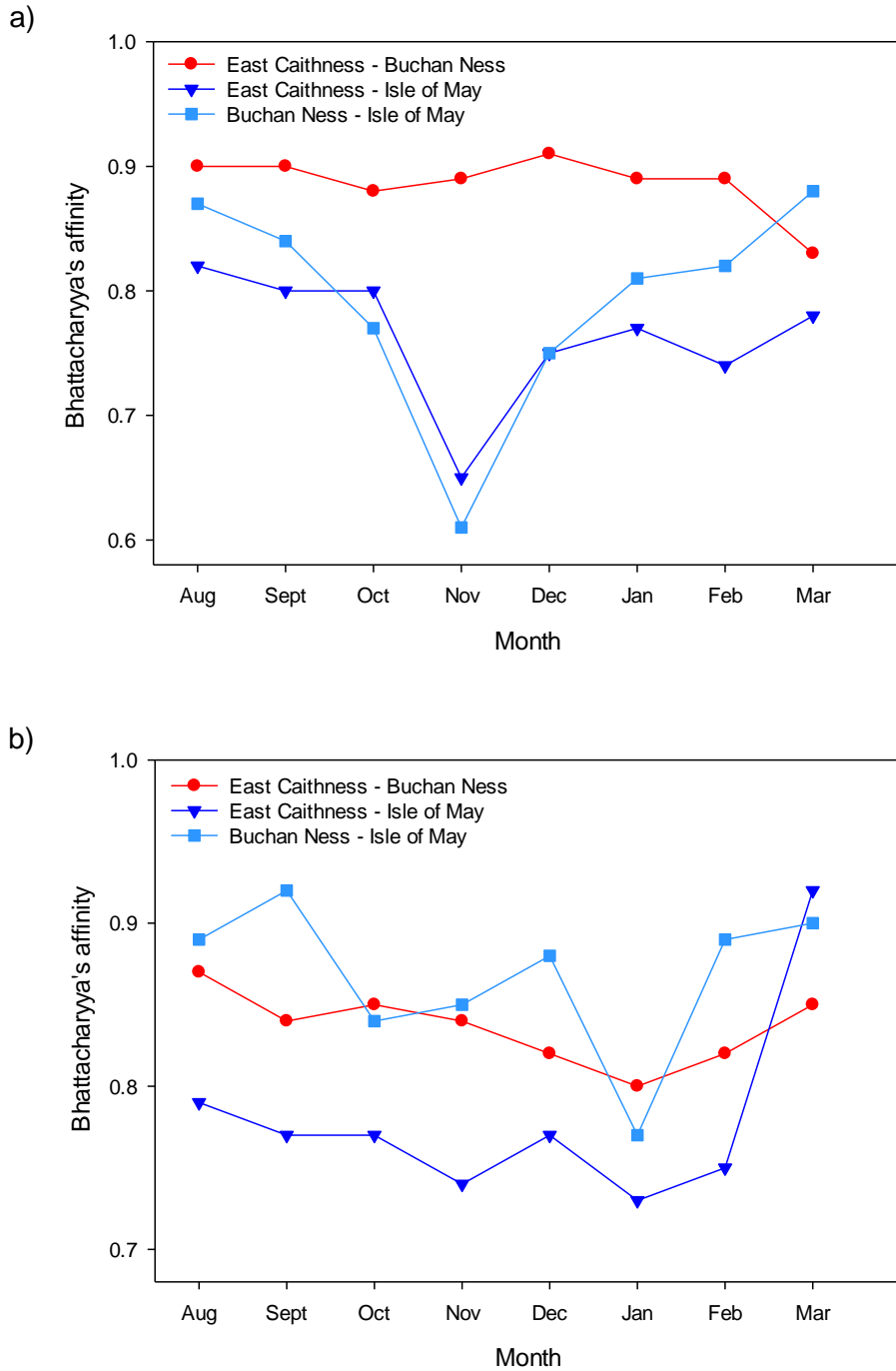


Fig. 26: Monthly pairwise spatial similarity of UD densities of guillemots from the three study colonies in: a) 2017-18; b) 2018-19. Similarity was estimated using Bhattacharyya's affinity measure, ranging from zero (no similarity) to 1 (identical densities).

Similar to guillemots, whole winter spatial similarity in the UD densities of razorbills between the three study colonies was very high in both years (Table 4).

| Colony | East Caithness | Buchan Ness to Collieston Coast | Isle of May |
|----------------|----------------|---------------------------------|-------------|
| a) 2017-18 | | | |
| East Caithness | | 0.89 | 0.87 |
| Buchan Ness | 0.89 | | 0.85 |
| Isle of May | 0.87 | 0.85 | |
| b) 2018-19 | | | |
| East Caithness | | 0.92 | 0.92 |
| Buchan Ness | 0.92 | | 0.89 |
| Isle of May | 0.92 | 0.89 | |

Table 4. Whole winter pairwise spatial similarity of UD_s of razorbills from the three study colonies in: a) 2017-18; b) 2018-19. Similarity was estimated using Bhattacharyya's affinity measure, ranging from zero (no similarity) to 1 (identical densities).

In 2017-18, monthly spatial similarity in UD_s of razorbills from the three colonies was very high in the post-breeding period, then gradually decreased and was lowest in late winter (Fig. 27a). Similarity between East Caithness and Buchan Ness to Collieston Coast was generally highest (Fig. 27a).

In 2018-19, a similar pattern was apparent, with highest similarity in the post-breeding period and lowest in late winter (Fig. 27b).

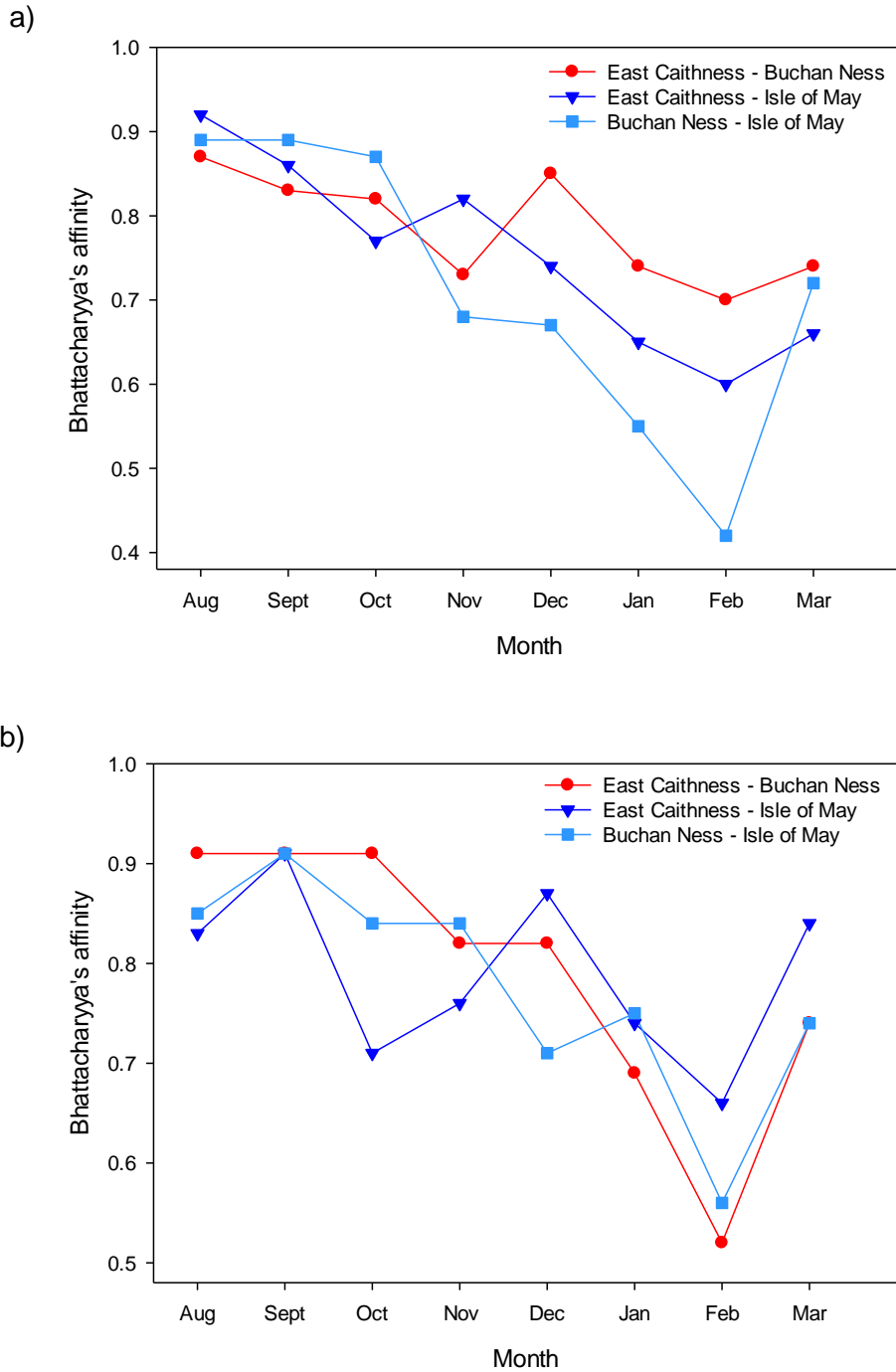


Fig. 27: Monthly pairwise spatial similarity of UD densities of razorbills from the three study colonies in: a) 2017-18; b) 2018-19. Similarity was estimated using Bhattacharyya's affinity measure, ranging from zero (no similarity) to 1 (identical densities).

3.1.3 Minimum adequate sample size of tracked birds

3.1.3.1 2017-2018

In guillemots from East Caithness, the resampling procedure using 50% UD contours indicated a gradual increase in the size of core areas used with increasing sample size of birds (Fig. 28a). The increment in cumulative area size was largest with sample size of up to around 6 birds, after which it was less than 5% with each additional bird (Fig. 28b). Randomized samples of 5 and 10 birds covered 57% and 79% of the area identified using all study birds, respectively (Fig. 28b).

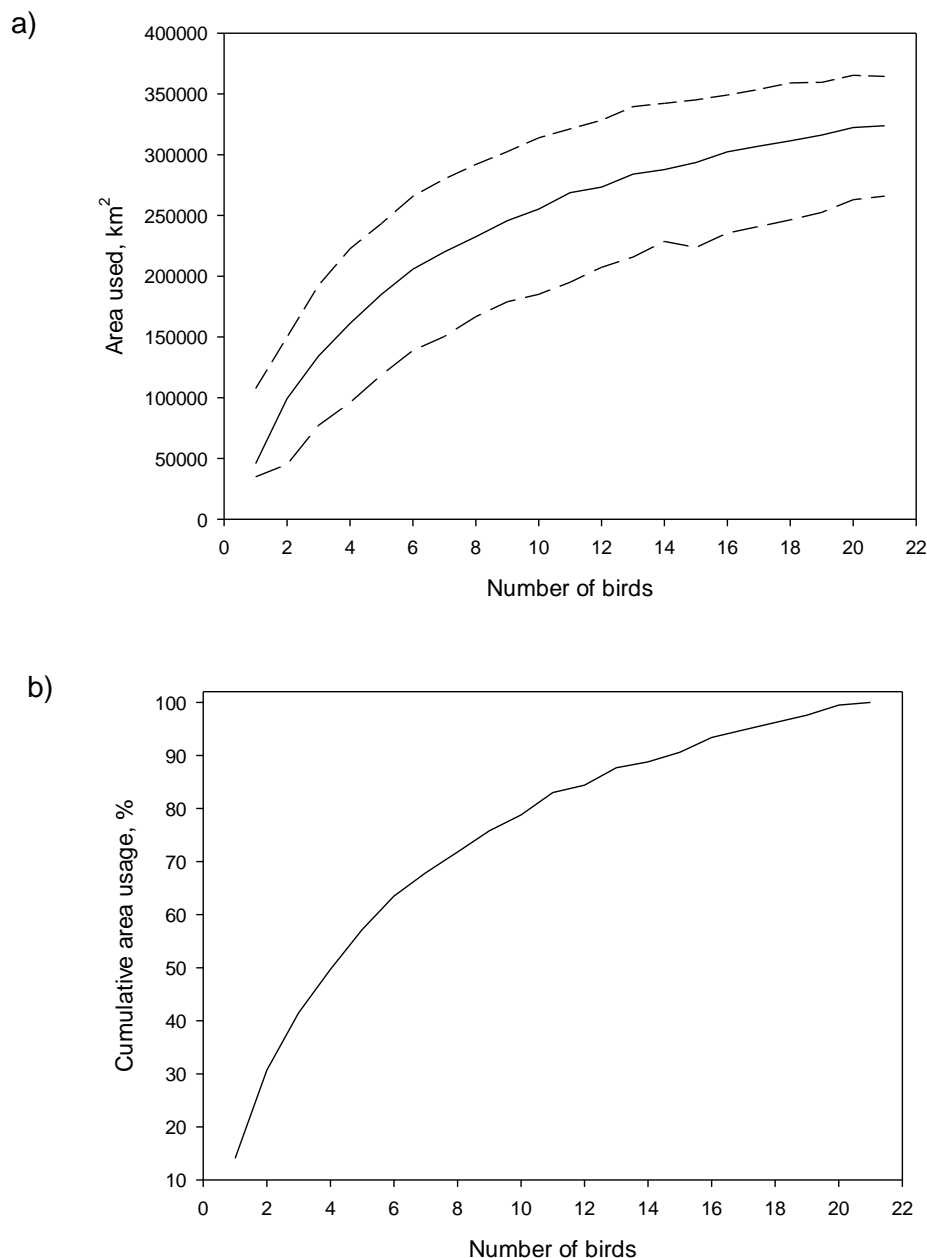


Fig. 28: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from East Caithness in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

In guillemots from Buchan Ness to Collieston Coast, a gradual non-linear increase in the size of core areas with increasing sample size of birds was apparent (Fig. 29a). The increment in cumulative area size was largest with sample size of up to 6 birds, after which it was less than 5% with each additional bird; for sample sizes over 14 birds the increment was less than 3% (Fig. 29b). Randomized samples of 5 and 10 birds covered 49% and 69% of the area identified using all study birds, respectively (Fig. 29b).

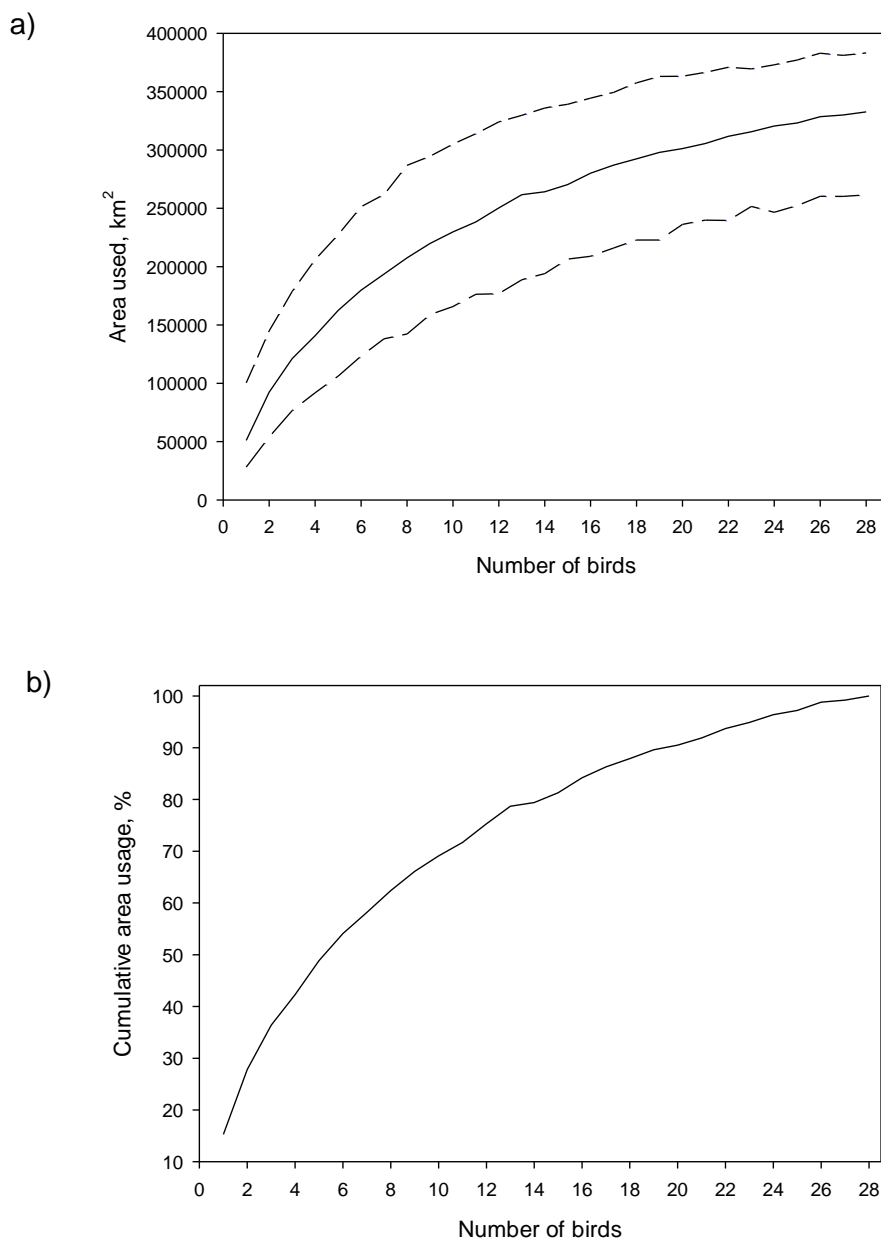


Fig. 29: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from Buchan Ness to Collieston Coast in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

In guillemots from the Isle of May, the size of core areas gradually increased with increasing sample size of birds (Fig. 30a). The cumulative curve did not appear to level off, suggesting the minimum adequate sample size was not reached at this colony (Fig. 30b). Randomized samples of 5 and 10 birds covered 51% and 79% of the area identified using all study birds, respectively (Fig. 30b).

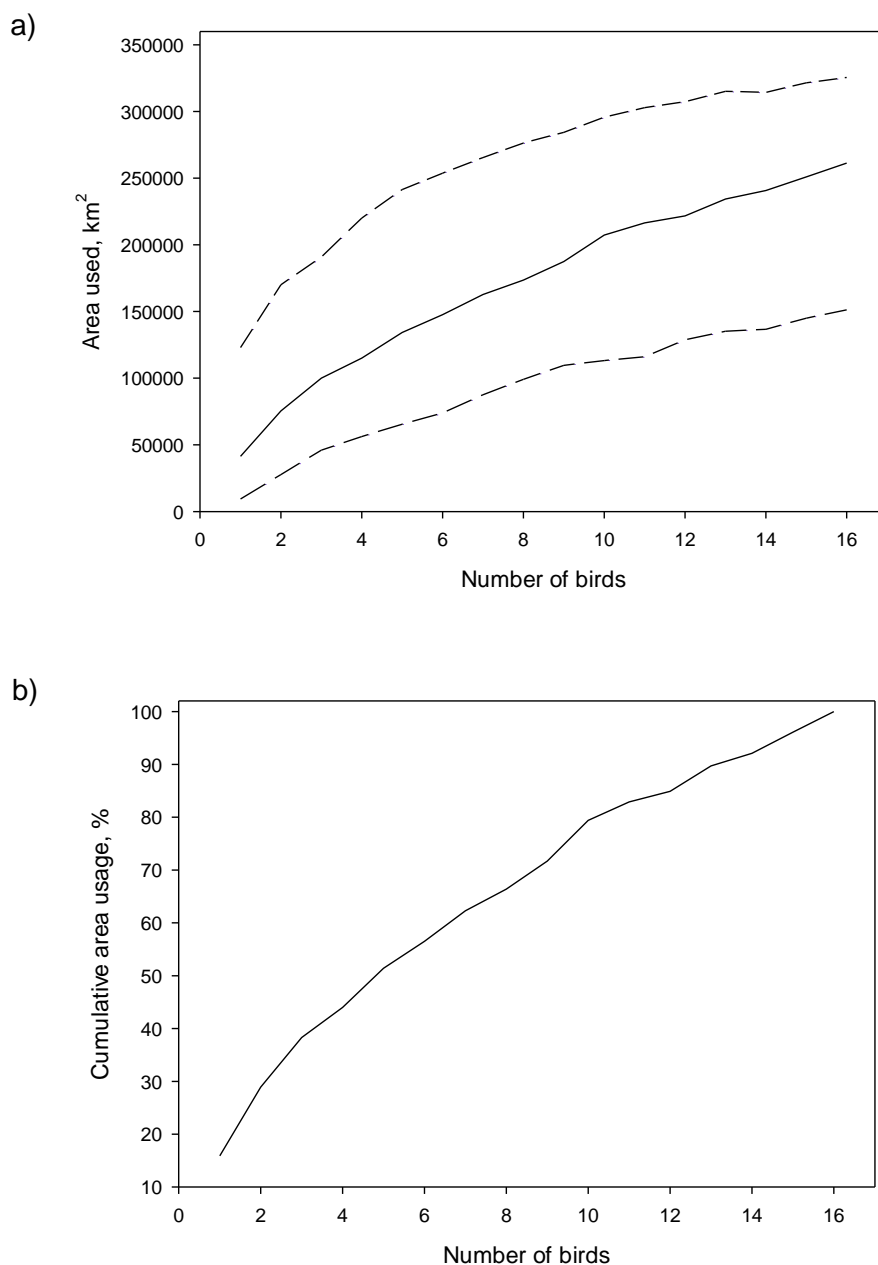


Fig. 30: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from Isle of May in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

In razorbills from East Caithness, as with guillemots, the size of core areas increased gradually with increasing sample size of birds (Fig. 31a). The increment in cumulative area size was larger with sample size up to 7 birds, after which it was less than 5% with each additional bird (Fig. 31b). Randomized samples of 5 and 10 birds covered 66% and 88% of the area identified using all study birds, respectively (Fig. 31b).

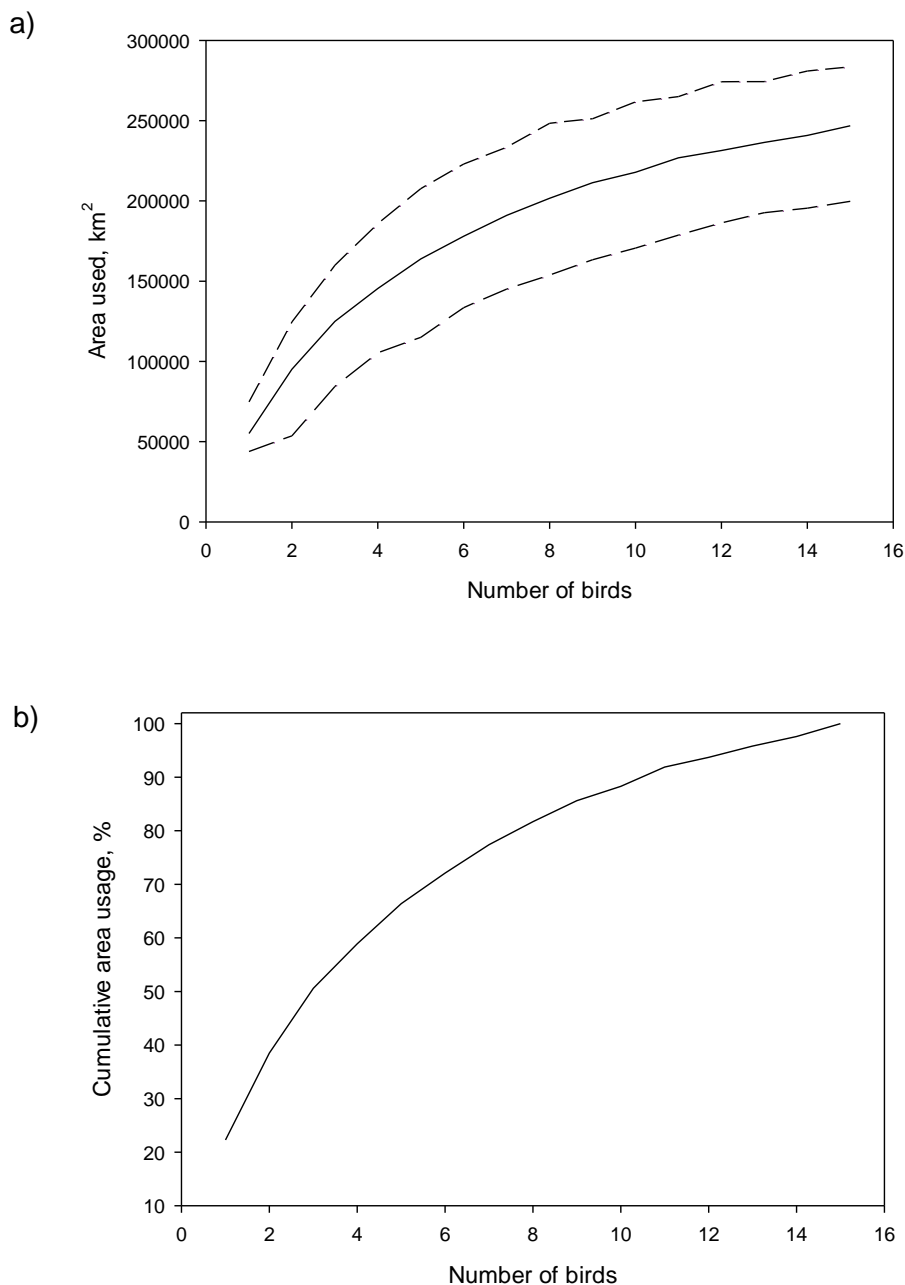


Fig. 31: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from East Caithness in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

At Buchan Ness to Collieston Coast only a total of 5 loggers were retrieved from razorbills. A gradual increase in the size of core areas with increasing number of birds was observed as in the other colonies and species (Fig. 32a) but unsurprisingly the cumulative curve did not level off due to the small sample size (Fig. 32b).

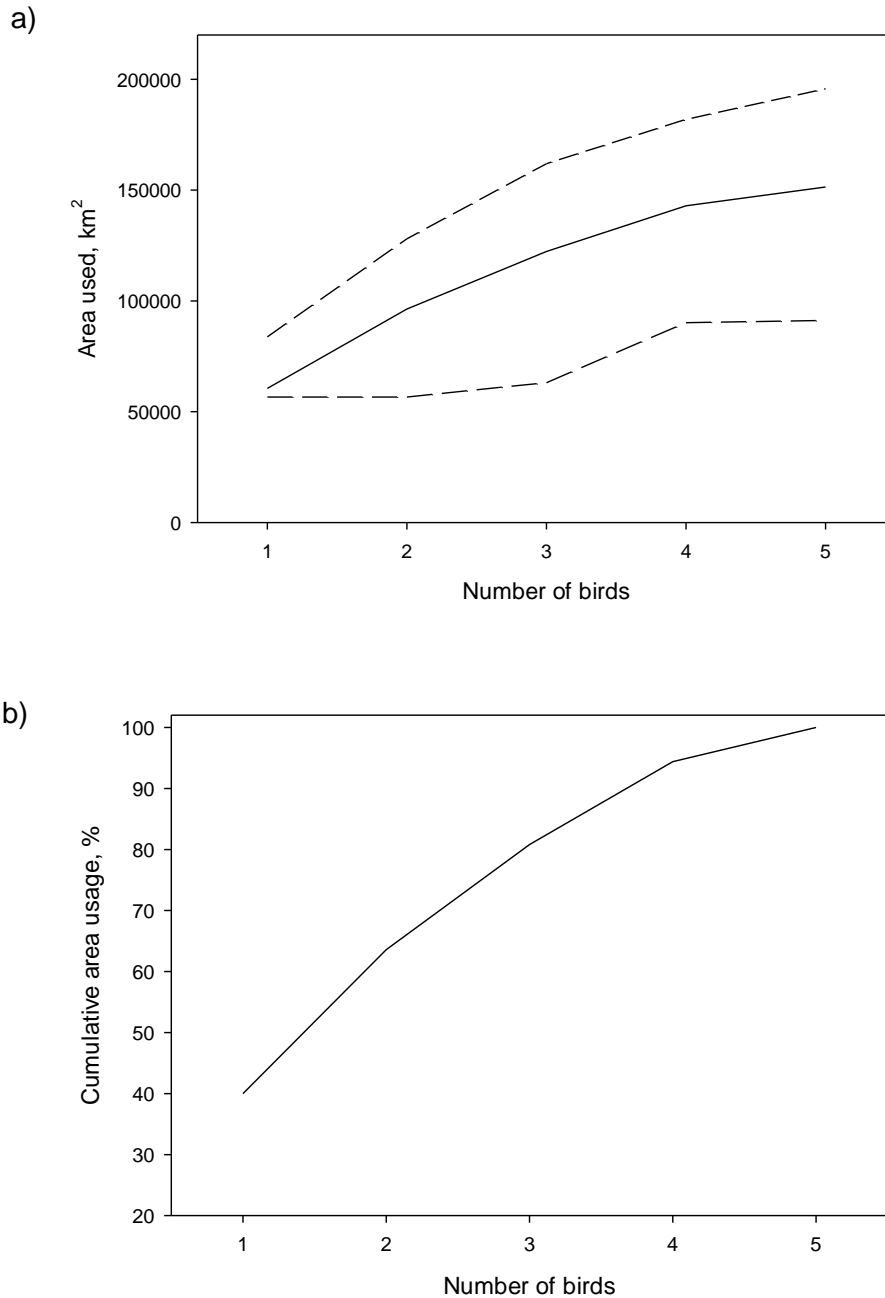


Fig. 32: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from Buchan Ness to Collieston Coast in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

The increase in size of core areas with increasing sample size of birds in Isle of May razorbills is shown on Fig. 33a. The increment in cumulative area size was larger with sample size up to around 7 birds, after which it declined to less than 5% with each additional bird (Fig. 33b). Randomized sample of 5 birds covered 71% of the area identified using all study birds (Fig. 33b).

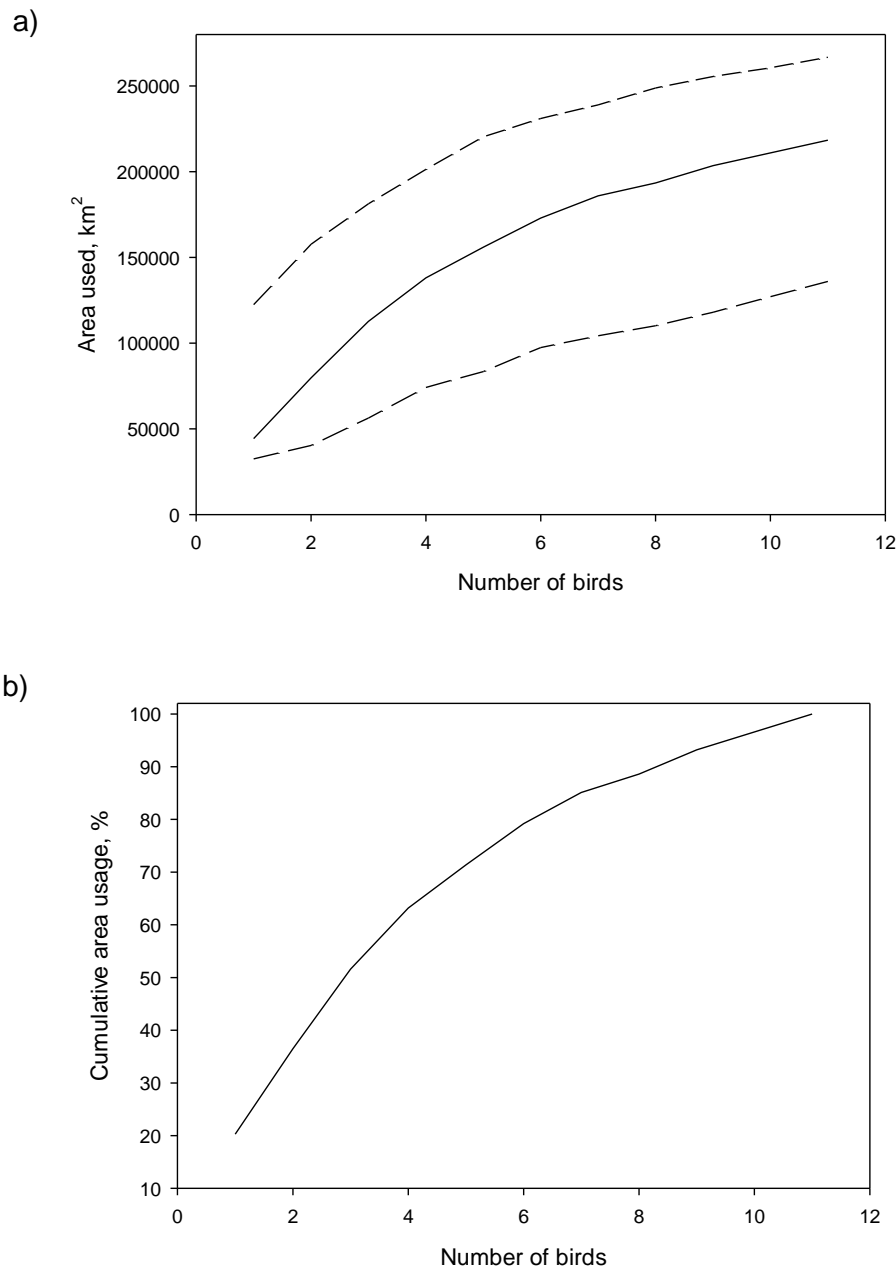


Fig. 33: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from Isle of May in 2017-18. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

3.1.3.2 2018-2019

As in 2017-18, East Caithness guillemots showed the typical non-linear increase in size of core areas used with increasing sample size of birds (Fig. 34a). The increment in cumulative area size was larger with sample size up to around 6 birds, after which it was less than 5% with each additional bird. For sample sizes above 12 birds the increment was 1-2% (Fig. 34b). Randomized samples of 5 and 10 birds covered 51% and 72% of the area identified using all study birds, respectively (Fig. 34b).

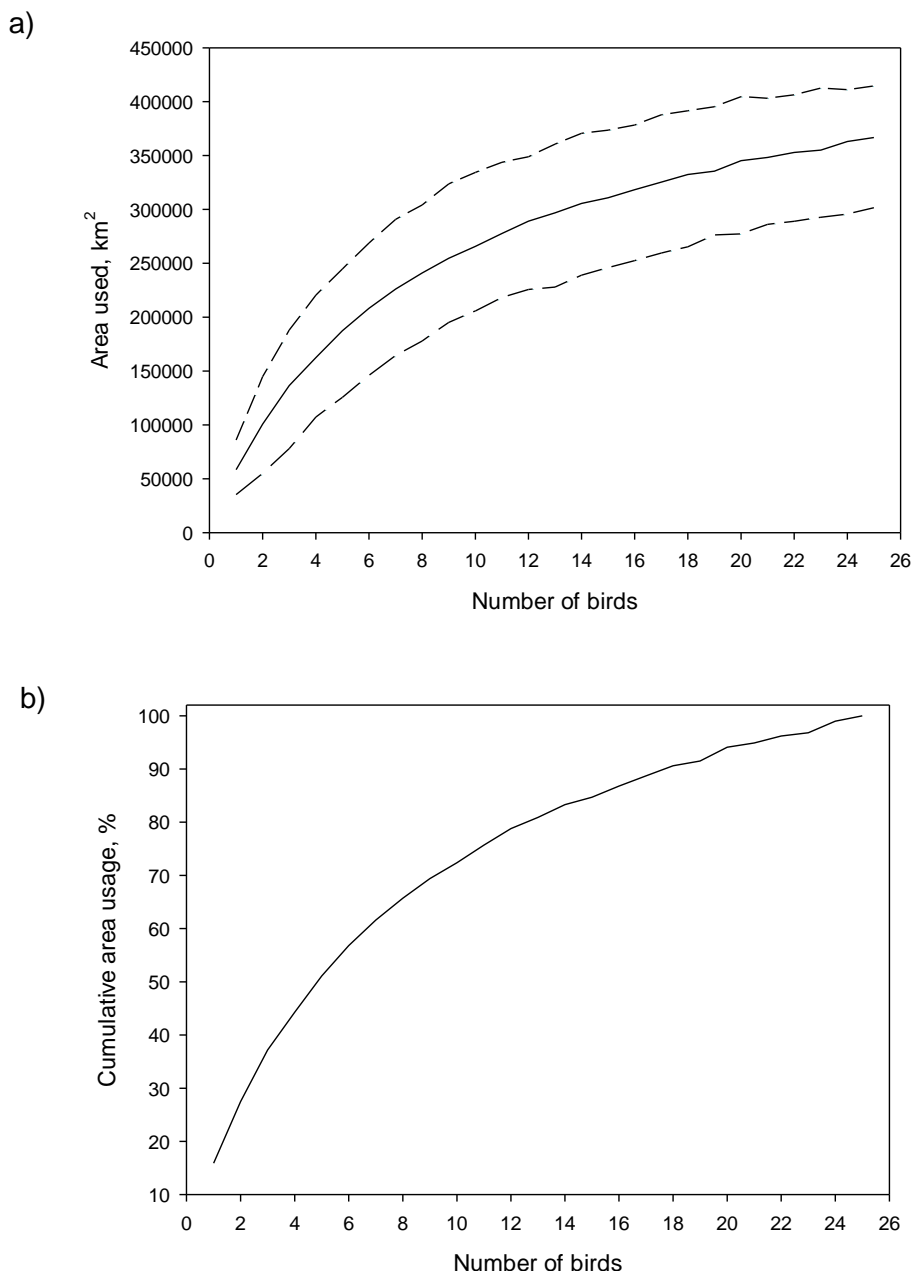


Fig. 34: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from East Caithness in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

In guillemots from Buchan Ness to Collieston Coast, a non-linear increase in size of core areas with increasing sample size of birds was evident (Fig. 35a). The increment in cumulative area size was larger with sample size up to around 7 birds, after which it was less than 5% with each additional bird. For sample sizes above 14 birds the increment was 1-2% (Fig. 35b). Randomized samples of 5 and 10 birds covered 55% and 75% of the area identified using all study birds, respectively (Fig. 35b).

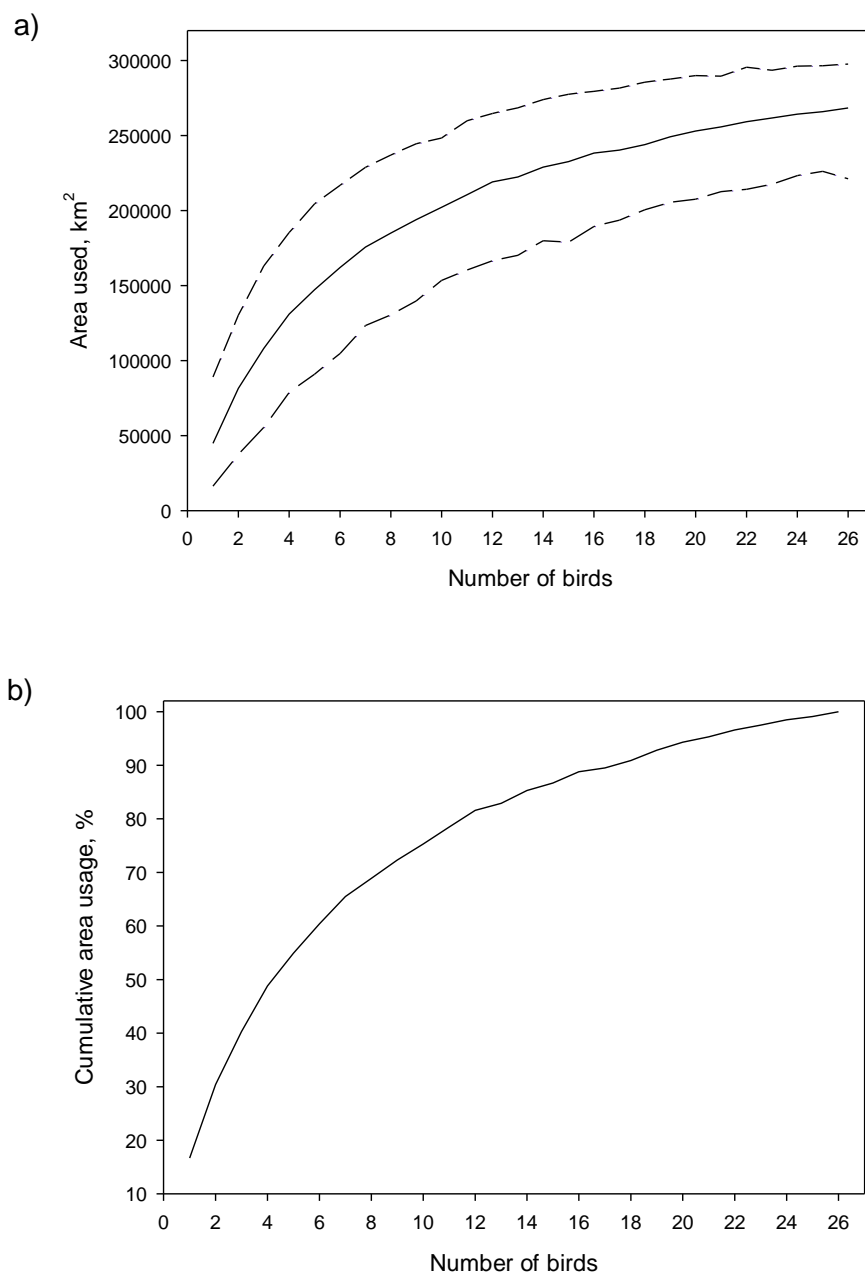


Fig. 35: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from Buchan Ness to Collieston Coast in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

As guillemots at the other two colonies, Isle of May guillemots showed the typical non-linear increase in size of core areas with increasing sample size of birds (Fig. 36a). The increment in cumulative area size was larger with sample size up to around 6 birds, after which it was less than 5% with each additional bird (Fig. 36b). Randomized samples of 5 and 10 birds covered 54% and 75% of the area identified using all study birds, respectively (Fig. 36b).

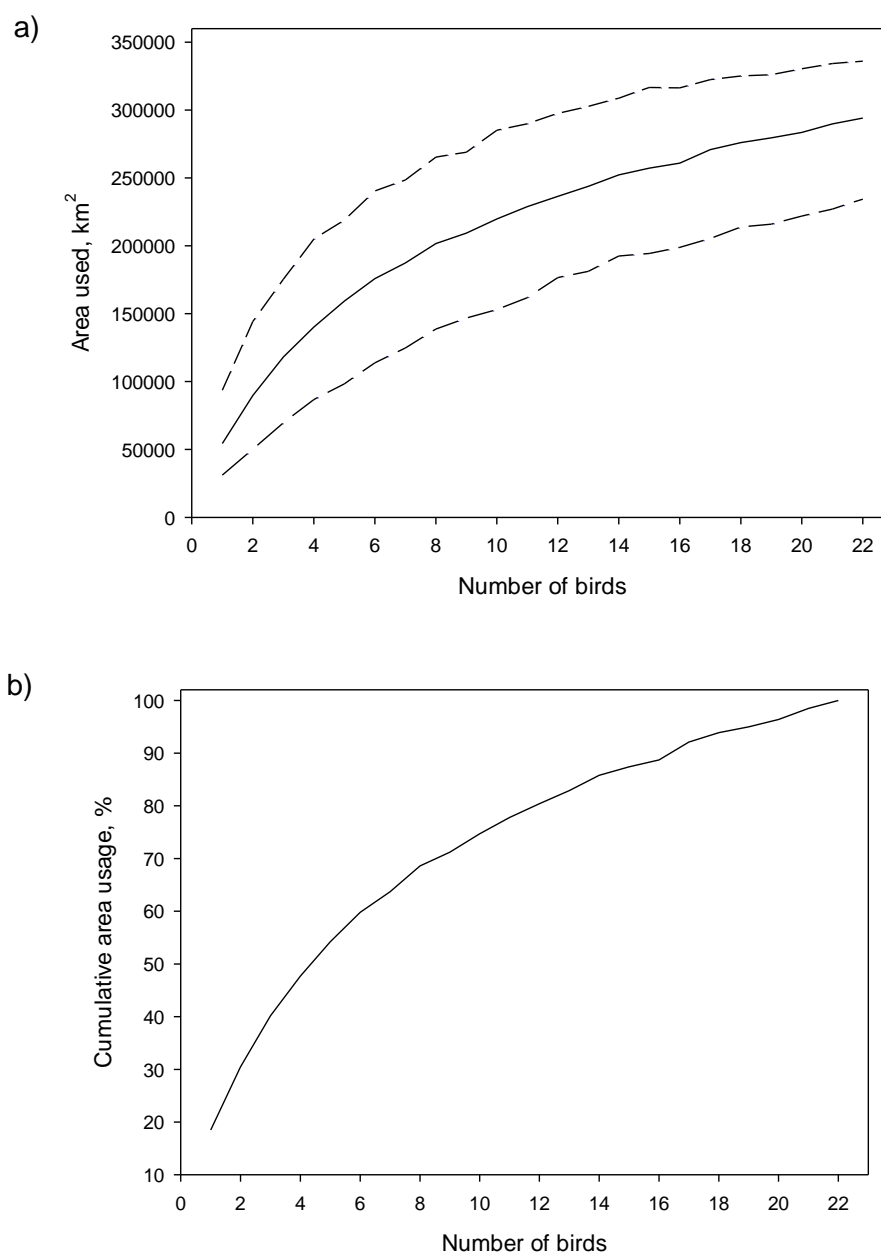


Fig. 36: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in guillemots from Isle of May in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

In razorbills from East Caithness, a non-linear increase in the size of core areas with increasing number of birds was observed as in the other colonies and species (Fig. 37a). However, the cumulative curve did not level off due to the relatively small sample size of birds (Fig. 37b).

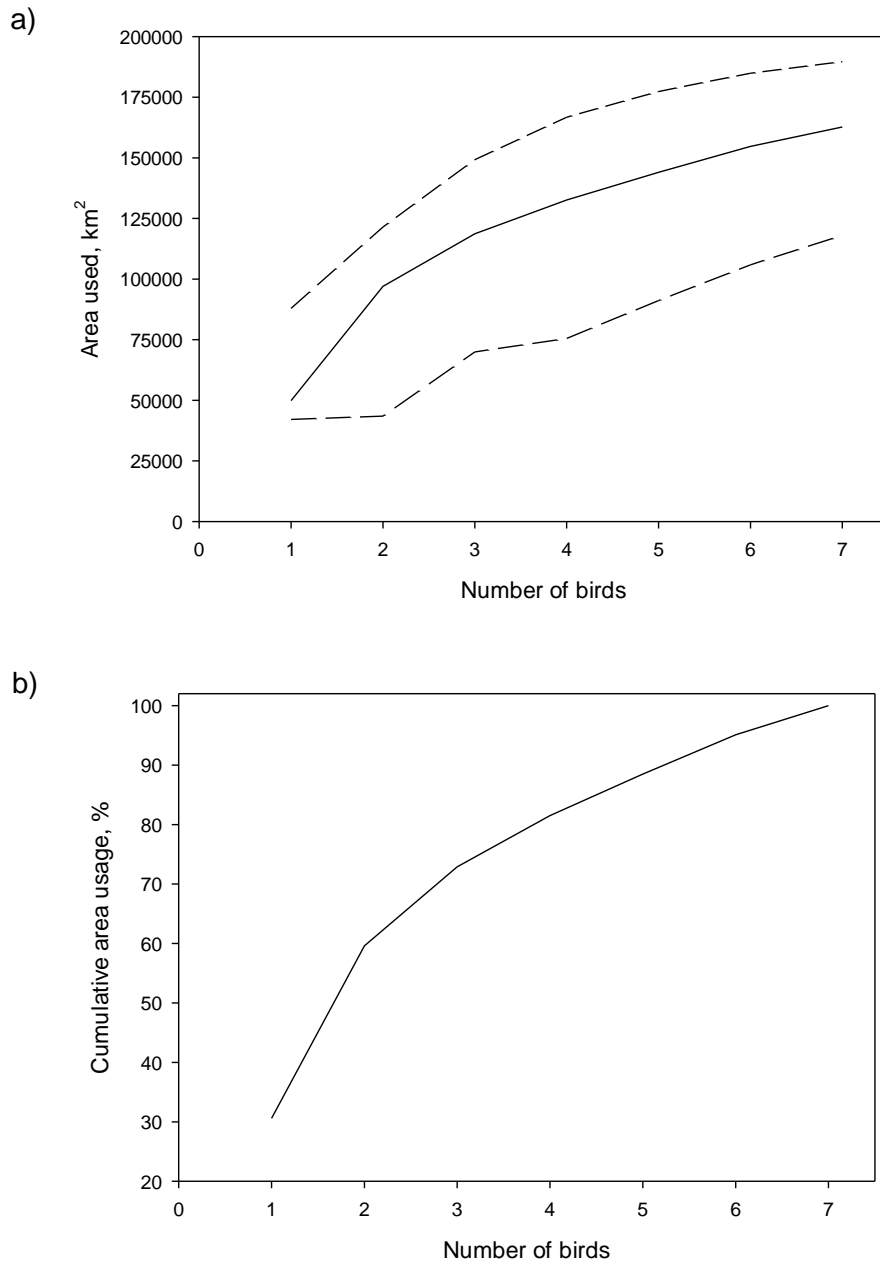


Fig. 37: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from East Caithness in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

Razorbills from Buchan Ness to Collieston Coast showed the expected non-linear increase in the size of core areas with increasing number of birds (Fig. 38a) but, similar to East Caithness, the cumulative curve did not level off, reflecting the relatively small sample size of birds (Fig. 38b).

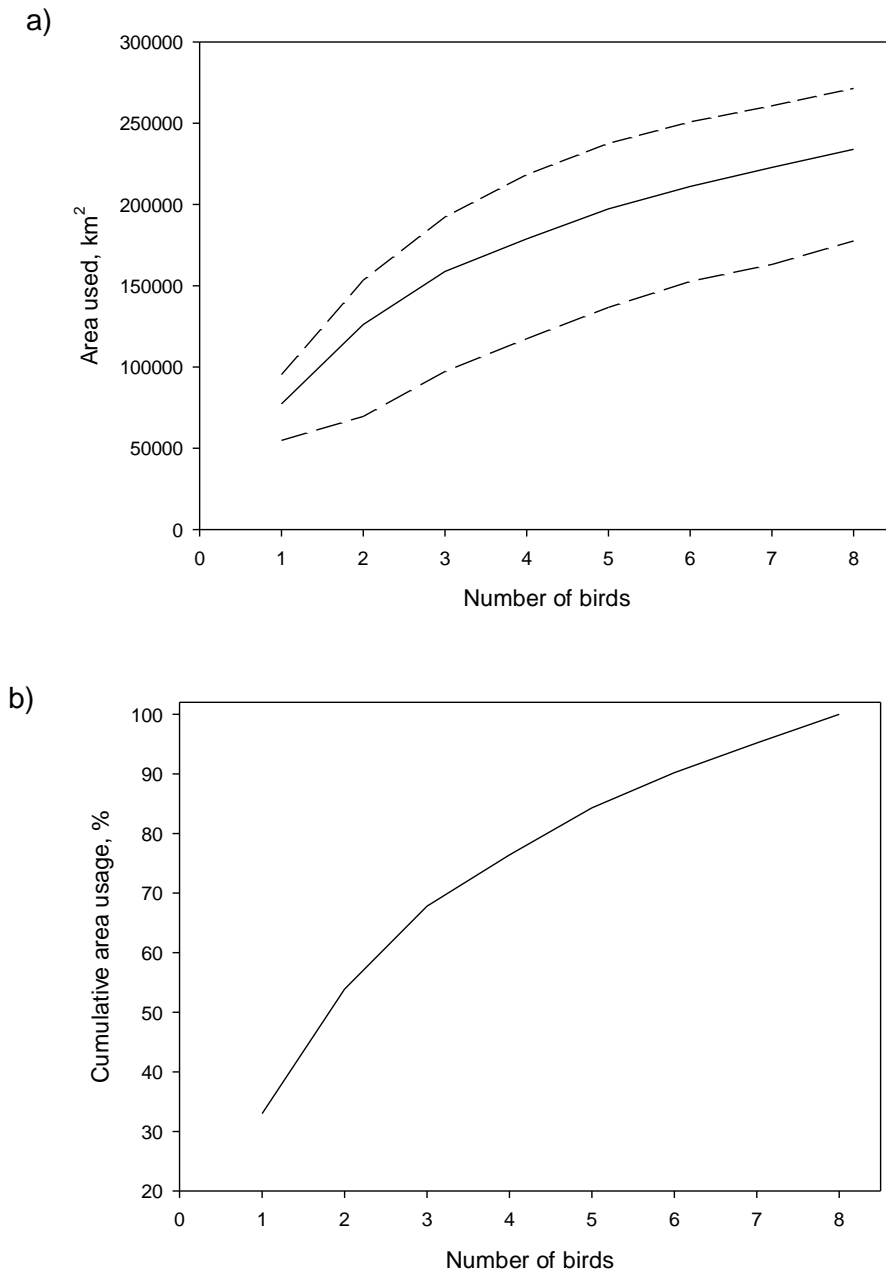


Fig. 38: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from Buchan Ness to Collieston Coast in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

The increase in size of core areas with increasing sample size of birds for Isle of May razorbills is shown on Fig. 39a. The increment in cumulative area size was larger with sample size up to around 8 birds, after which it declined to less than 5% with each additional bird (Fig. 39b). Randomized sample of 5 birds covered 74% of the area identified using all study birds (Fig. 39b).

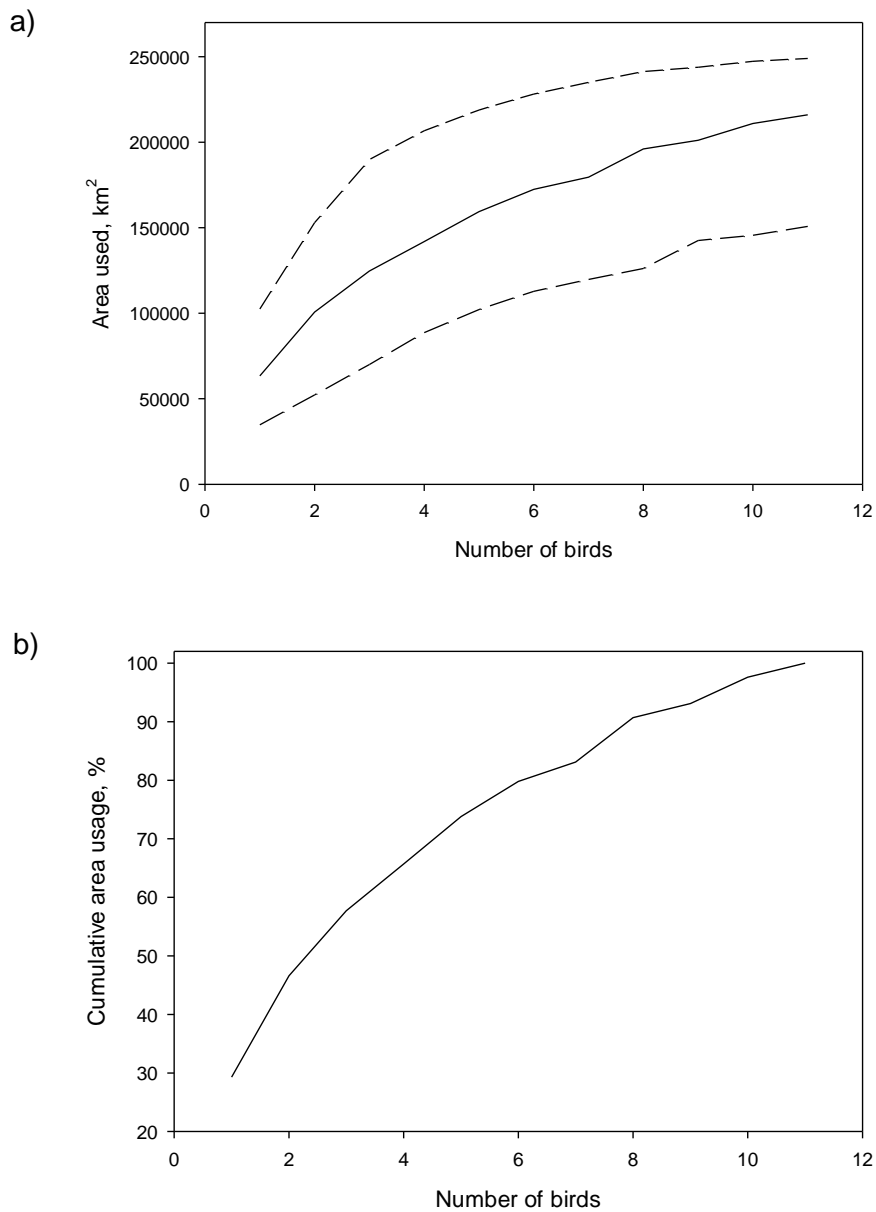


Fig. 39: Relationship between core area used (50% UD contours) and sample size of birds estimated from a resampling procedure in razorbills from Isle of May in 2018-19. a) median area (solid line) and 2.5 and 97.5 percentiles (dashed lines) shown for each randomized sample size; b) cumulative percentage of area used by the population.

3.2 Diet

3.2.1 *Breeding season in 2017*

We obtained 548 diet observations from East Caithness (n = 422 for guillemot; n=126 for razorbill), 389 diet observations from Buchan Ness to Collieston Coast (n = 298 for guillemot; n=91 for razorbill) and 1,270 diet observations from the Isle of May (n = 1,021 for guillemot; n = 249 for razorbill), giving a total sample across all three colonies of 2,207 in 2017.

At all three colonies, the diet of guillemots as presented as Frequency of Occurrence (i.e. the percentage of diet observations that comprise that prey type) was dominated by clupeids (70% East Caithness; 65% Buchan Ness to Collieston Coast, 86% Isle of May). Guillemots at East Caithness brought back more 1+ group sandeels than at the other two colonies (19%, 9% and 5% respectively), and gadoids were less important on the Isle of May (5%, 7% and 1% respectively). 0 group sandeels were unimportant at all three colonies (Table 5).

The diet of razorbills showed marked among-colony variation, such that 0 group sandeels dominated at East Caithness and Buchan Ness to Collieston Coast (98% East Caithness; 93% Buchan Ness to Collieston Coast, 33% Isle of May) whereas clupeids dominated on the Isle of May (1% East Caithness; 2% Buchan Ness to Collieston Coast, 60% Isle of May; Table 5).

3.2.2 *Breeding season in 2018*

We obtained 299 diet observations from East Caithness (n = 262 for guillemot; n=37 for razorbill), 528 diet observations from Buchan Ness to Collieston Coast (n = 481 for guillemot; n = 47 for razorbill) and 1,579 diet observations from the Isle of May (n = 1,382 for guillemot; n = 197 for razorbill), giving a total sample across all three colonies of 2,406 in 2018.

A summary of Frequency of Occurrence of each prey type for the two species at the three colonies can be found in Table 5. At all three colonies, the diet of guillemots was dominated by clupeids (76% at East Caithness; 70% Buchan Ness to Collieston Coast,

77% Isle of May; Table 5). Guillemots on the Isle of May brought back fewer 1+ group sandeels than at the other two colonies (19%, 19% and 6% respectively), and gadoids were more important on the Isle of May (1%, 1% and 5% respectively). 0 group sandeels were unimportant at all three colonies. These results accorded closely with those found in 2017 (Table 5).

The diet of razorbills showed marked among-colony variation such that 0 group sandeels dominated at East Caithness and Buchan Ness to Collieston Coast (95% East Caithness; 98% Buchan Ness to Collieston Coast, 10% Isle of May) whereas clupeids dominated on the Isle of May (5% East Caithness; 2% Buchan Ness to Collieston Coast, 80% Isle of May). No 1+ group sandeels were observed in the diet of razorbills. These results accorded closely with those found in 2017 (Table 5), although the importance of clupeids in the diet of Isle of May razorbills was even more marked in 2018.

3.2.3 Breeding season in 2019

We obtained 284 diet observations from East Caithness (n = 207 for guillemot; n=77 for razorbill), 479 diet observations from Buchan Ness to Collieston Coast (n = 417 for guillemot; n = 62 for razorbill) and 1,061 diet observations from the Isle of May (n = 881 for guillemot; n = 180 for razorbill), giving a total sample across all three colonies of 1,824 in 2019.

A summary of Frequency of Occurrence of each prey type for the two species at the three colonies can be found in Table 5. At all three colonies, the diet of guillemots was dominated by clupeids (59% Frequency of Occurrence at East Caithness; 59% Buchan Ness to Collieston Coast, 87% Isle of May; Table 5). Guillemots on the Isle of May brought back fewer 1+ group sandeels than at the other two colonies (29%, 26% and 4% respectively), and gadoids were somewhat more important at Buchan Ness to Collieston Coast (4%, 7% and 3% respectively). 0 group sandeels were unimportant at all three colonies. These results accorded closely with those found in 2017 and 2018 (Table 5).

The diet of razorbills showed among-colony variation, such that 0 group sandeels dominated at East Caithness and Buchan Ness to Collieston Coast (100% East Caithness; 98% Buchan Ness to Collieston Coast, 69% Isle of May). However, this was a much higher Frequency of Occurrence of this prey type on the Isle of May than in the other two years. Unlike in 2017 and 2018, clupeids didn't dominate at any colony (1% East Caithness; 12% Buchan Ness to Collieston Coast, 16% Isle of May). Few 1+ group sandeels were observed in the diet of razorbills.

To summarise the interannual variation, the diet of guillemots was largely consistent with the dominance of clupeids and negligible importance of 0 group sandeels at all colonies, and greater importance of 1+ group sandeels at East Caithness and Buchan Ness to Collieston coast compared with the Isle of May apparent in all three years. As regards razorbill, consistent across all years was the dominance of 0 group sandeels at East Caithness and Buchan Ness to Collieston coast. However, their diet on the Isle of May differed among years, with clupeids dominant in 2017 and 2018 but 0 group sandeels dominant in 2019.

The overall mean across the three years support these conclusions, with the diet of guillemots dominated by clupeids (69% Frequency of Occurrence at East Caithness; 65% Buchan Ness to Collieston Coast, 82% Isle of May). Guillemots on the Isle of May brought back fewer 1+ group sandeels than at the other two colonies across the three years (21%, 19% and 5% respectively), and gadoids were of similar importance at the three colonies overall (4%, 5% and 3% respectively). The overall mean across the three years for razorbills showed that 0 group sandeels dominated at East Caithness and Buchan Ness to Collieston Coast (98% East Caithness; 96% Buchan Ness to Collieston Coast, 36% Isle of May) whereas clupeids dominated on the Isle of May (2% East Caithness; 5% Buchan Ness to Collieston Coast, 54% Isle of May).

Hywind Scotland Ornithological Monitoring Programme

| 2017 | Prey type | East Caithness | | Buchan Ness to Collieston Coast | | Isle of May | |
|------|------------------|----------------|-----------|---------------------------------|-----------|-------------|-----------|
| | | guillemot | razorbill | guillemot | razorbill | guillemot | razorbill |
| | 0 group sandeel | 0.71 | 97.62 | 0.67 | 93.41 | 0.29 | 32.53 |
| | 1+ group sandeel | 19.43 | 2.38 | 9.06 | 2.20 | 5.09 | 0.00 |
| | Clupeid | 69.91 | 0.79 | 65.44 | 2.20 | 86.09 | 59.84 |
| | Gadoid | 5.21 | 0.00 | 7.05 | 1.10 | 1.27 | 0.00 |
| | Rockling | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Squid | 0.00 | 0.00 | 0.00 | 0.00 | 0.10 | 0.00 |
| | Crustacean | 0.00 | 0.79 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Unidentified | 4.74 | 0.79 | 17.79 | 2.20 | 7.15 | 8.03 |

| 2018 | Prey type | East Caithness | | Buchan Ness to Collieston Coast | | Isle of May | |
|------|------------------|----------------|-----------|---------------------------------|-----------|-------------|-----------|
| | | guillemot | razorbill | guillemot | razorbill | guillemot | razorbill |
| | 0 group sandeel | 0.00 | 94.59 | 1.25 | 97.87 | 0.58 | 9.64 |
| | 1+ group sandeel | 18.70 | 0.00 | 19.13 | 0.00 | 6.22 | 0.00 |
| | Clupeid | 76.34 | 5.41 | 69.65 | 2.13 | 77.21 | 79.70 |
| | Gadoid | 0.76 | 0.00 | 1.66 | 0.00 | 4.78 | 0.00 |
| | Rockling | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 3.05 |
| | Squid | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Crustacean | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Unidentified | 4.20 | 5.41 | 8.32 | 2.13 | 11.79 | 9.64 |

| 2019 | Prey type | East Caithness | | Buchan Ness to Collieston Coast | | Isle of May | |
|------|------------------|----------------|-----------|---------------------------------|-----------|-------------|-----------|
| | | guillemot | razorbill | guillemot | razorbill | guillemot | razorbill |
| | 0 group sandeel | 5.31 | 100.00 | 1.20 | 98.39 | 0.00 | 69.44 |
| | 1+ group sandeel | 29.47 | 0.00 | 26.14 | 1.61 | 3.75 | 0.56 |
| | Clupeid | 59.42 | 1.30 | 58.51 | 1.61 | 87.40 | 16.11 |
| | Gadoid | 3.86 | 0.00 | 6.95 | 0.00 | 3.18 | 1.11 |
| | Rockling | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 1.67 |
| | Squid | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Crustacean | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| | Unidentified | 1.93 | 0.00 | 7.19 | 0.00 | 5.68 | 12.22 |

Table 5: Frequency of Occurrence of each prey type (expressed as percentage of diet observations that comprise that prey type) delivered by guillemots and razorbills at each colony in 2017, 2018 and 2019.

4 Discussion

4.1 Guillemot and razorbill winter distributions

4.1.1 Within- and between-species variation in winter distributions in relation to colony origin

At a large spatial scale, the distribution of guillemots from East Caithness, Buchan Ness to Collieston Coast and the Isle of May was broadly similar, with key wintering areas located around the colonies, off the north-eastern coast of Scotland and in the central and southern North Sea, as previously demonstrated in Isle of May guillemots (Harris et al. 2015; Dunn et al. 2020). In addition, some East Caithness and Isle of May birds migrated eastwards, to the Skagerrak/Kattegat that form the transient zone between the North Sea and the Baltic Sea. As with guillemots, razorbill distributions were broadly similar among colonies at a large spatial scale. The key wintering area of this species was also in the central and southern North Sea, with some birds from East Caithness and Buchan Ness to Collieston Coast migrating to the Skagerrak/Kattegat. Single individuals from both species also used parts of the Norwegian Sea, as shown previously (Harris et al. 2015). These areas are generally known to be highly productive. For example, the waters around the Dogger Bank have increased productivity associated with the existence of tidal mixing frontal systems (Daewel & Schrum 2013). Similarly, the Skagerrak/Kattegat zone supports high concentrations of immature schooling fish (Knijn et al. 1993, Skov et al. 2000), of which herring has been shown to be of key importance as seabird prey (Skov et al. 2000). Thanks to the favourable feeding conditions, these areas host large multispecies aggregations of wintering seabirds (Skov et al. 1995, 2000).

In both species, however, the similarity in distributions of birds from different colonies varied between months, suggesting there were some differences in the location of areas used during the non-breeding season. This may reflect localised within-species segregation driven by competition (Ratcliffe et al. 2014, McFarlane Tranquilla et al. 2015), although competition is likely to be less pronounced in the winter than during the breeding season when the birds are restricted to forage near the colony (Grémillet et al. 2004, Masello et al. 2010, Wakefield et al. 2013). It may also arise in part because of the cost of reaching winter destinations, which will depend to some extent on colony

location (Bogdanova et al. 2017) and may be particularly high in auks due to the high flight costs they incur (Pennycuik 1987, Gaston 2004; Dunn et al. 2020).

Partial overlap was also observed between the wintering areas of guillemots and razorbills from the same colony. This pattern may be linked to differences in diet between the two species as shown previously (Linnebjerg et al. 2013, Glew et al. 2018). Such partial overlap can lead to reduced between-species competition at the wintering grounds, as found in other sympatric species (e.g. Fort et al. 2013, Ratcliffe et al. 2014, McFarlane Tranquilla et al. 2015).

In both species, the birds generally spent the post-breeding moult period (during which they are flightless) relatively close to their colonies, with the exception of some Isle of May guillemots that had moved further north along the east coast of Scotland or to the Skagerrak area prior to moult. This matches previous observations of birds visiting the colony in the autumn (Harris & Wanless 1990). However, there was a substantial overlap in the areas used by the three colonies at this time, suggesting that the large post-breeding aggregations of both species observed in the Hywind Scotland project area (Natural Research Projects Ltd 2015, Statoil 2015) may be formed by birds from all three colonies. We cannot discount the possibility that the area is also used by birds from other breeding colonies that were not part of this study. Our study birds were not sexed so we were unable to determine whether the longer-distance post-breeding movements by some of the Isle of May guillemots were associated with one of the sexes, resulting from differences between males and females in the duration of parental care (Gaston & Jones 1998). As expected, more major movements took place largely during October and February-March. However, among guillemots some birds returned to the colonies already in January. Although razorbills generally wintered away from the colony area, they spent more time in local waters than previously thought. These findings highlight the year-round importance of local areas around the colonies.

4.1.2 *Minimum adequate sample size*

The analyses of minimum adequate sample size of tracked birds for each species at each of the colonies and years all showed the typical non-linear decline in rate of

increase in the size of core area used with increasing sample size. In all cases, the cumulative curves did not reach a horizontal plateau, the point at which adding more birds to the sample would lead to no further increase in the population core wintering area. However, the method we used is relatively conservative since the cumulative area of individual kernel contours is calculated to estimate the size of the population wintering area at each sample size of birds, as opposed to data from all birds within a sample size being pooled and population kernel contours then calculated.

Our results are in line with most tracking studies of seabird at-sea distributions, and reflects the challenge in obtaining the large sample sizes of birds that would be required to achieve a plateau in the cumulative curves (Frederiksen et al. 2012, Fort et al. 2013, Garthe et al. 2016). However, it is encouraging to note that in most cases the shape of the curves indicated that a larger sample size of birds would not have resulted in a dramatically altered size of core wintering areas for these populations. This was particularly the case for guillemots where larger sample sizes were obtained, reflecting the fact that more accessible birds were available and these were easier to recapture than razorbills. As such, the relationship between cumulative area and sample size in this species was approaching a plateau more so than in razorbills.

4.2 Diet variation among colonies and years

We found important differences in diet among the two species and among study colonies. Clupeids dominated the diet of guillemots at all three locations, and 0 group sandeels were unimportant. However, 1+ group sandeels were more important at the two northern colonies than the Isle of May, where clupeids were particularly dominant. The dominance of clupeids in all years suggests consistent availability of this prey across the region over this time period. The long-term study on the Isle of May shows that the dominance of clupeids over 1+ group sandeels has in fact been apparent at this colony since the late 1990s (Wanless et al. 2018). However, our results are in contrast to the only previous study of diet at Buchan Ness to Collieston Coast. In 2006, observers at the colony recorded a low proportion of 1+ group sandeels, as we found, but the principal prey was gadids as opposed to clupeids (Anderson et al. 2014). These differences likely reflect changes in the availability of these two prey types over this 10-

year period. However, it is not possible to verify this because of the lack of direct data on prey distributions.

For razorbills, there were marked among-colony and among-year differences whereby clupeids dominated on the Isle of May in two out of three years, whereas 0 group sandeels dominated at the other two colonies in all three years, but only in 2019 on the Isle of May. The among-colony differences could have an important effect on provisioning of offspring since clupeids are generally larger and more nutritious than 0 group sandeels (Harris et al. 2008), which may not be compensated for by the typically large number of prey items that are delivered with the latter. The relative profitability of the two prey types also depends on the relative costs in obtaining them. It is challenging to ascertain why these differences exist among colonies, and among years for the Isle of May since, as with guillemot diet, direct data on prey distribution are not available. However, the most parsimonious explanation is that razorbills are responding to what prey are available close to the colony, and that clupeids were more available to razorbills on the Isle of May in 2017 and 2018, while in 2019 at that colony and at the other two colonies in all three years there was likely a greater availability of 0 group sandeels.

It is important to note that while the diet sample sizes were sufficient to be confident that they provide a representative summary of diet over the sampling period, that diet may have differed outside these periods e.g. during incubation.

4.3 Conclusions: winter ecology of guillemots and razorbills from the three colonies and implications for offshore renewable developments

Seabirds are long-lived species whose population dynamics are largely determined by adult survival rates. Adult mortality typically occurs outside the breeding season, so conditions at the wintering grounds can have a profound impact on population trajectories (Reynolds et al. 2011). The extent of winter mixing of birds from different breeding populations, and therefore the potential for shared experiences of poor conditions or interactions with human developments, can have important conservation

implications (Frederiksen et al. 2012). Our results show that the three populations of guillemots and razorbills showed a similar overall distribution, with extensive use of the central and southern North Sea and areas around the breeding colonies. However, in both species there were important differences among colonies in the location of hotspots during the non-breeding period. These differences likely reflect the considerable amount of time that birds were spending close to their colonies outside the breeding season, and suggest that they may show different population dynamic responses to environmental conditions operating at local scales.

Offshore renewable developments must consider potential consequences of impacts on protected seabird populations. In the case of guillemots and razorbills, the principal concern is displacement and barrier effects (Masden et al. 2010; Dierschke et al. 2016). These effects could be particularly important in the case of multiple developments, as the potential for cumulative negative impacts increases (e.g. Busch et al. 2013). Our study provides important insights into the year-round space use of these two key species at three important colonies on the east coast of Scotland, and therefore the extent to which birds from multiple colonies may be affected by particular single or multiple developments at different periods of the year.

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7 Appendix: Deployment details

Tables A1a, A1b and A1c provide deployment details at East Caithness, Buchan Ness to Collieston Coast and the Isle of May, respectively, in 2017.

Tables A2a, A2b and A2c provide deployment details at East Caithness, Buchan Ness to Collieston Coast and the Isle of May, respectively, in 2018.

Table A1a: GLS logger deployment details for East Caithness in 2017. Start date and start time denote the date the logger commenced data collection.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Guillemot | 1a | R99313 | 2561 | 11-Jun-17 | 15:37 | 12-Jun-17 | 09:50 | Brooding chick | NA | NA |
| 2 | Guillemot | 1a | R99314 | 2560 | 11-Jun-17 | 15:36 | 12-Jun-17 | 09:56 | Incubating egg | 2018 | Yes |
| 3 | Guillemot | 1b | R99315 | 2556 | 11-Jun-17 | 15:33 | 12-Jun-17 | 10:10 | Brooding chick | 2018 | Yes |
| 4 | Guillemot | 1b | R99316 | 2564 | 11-Jun-17 | 15:38 | 12-Jun-17 | 10:13 | Brooding chick | 2018 | Yes |
| 5 | Guillemot | 1b | R99317 | 2531 | 11-Jun-17 | 15:22 | 12-Jun-17 | 10:15 | Brooding chick | 2018 | No |
| 6 | Guillemot | 1b | R99318 | 2544 | 11-Jun-17 | 15:28 | 12-Jun-17 | 10:18 | Brooding chick | 2018 | Yes |
| 7 | Guillemot | 1b | R99319 | 2542 | 11-Jun-17 | 15:27 | 12-Jun-17 | 10:20 | Brooding chick | 2018 | Yes |
| 8 | Guillemot | 1b | R99320 | 2533 | 11-Jun-17 | 15:23 | 12-Jun-17 | 10:22 | Brooding chick | 2018 | Yes |
| 9 | Guillemot | 1b | T15286 | 2552 | 11-Jun-17 | 15:31 | 12-Jun-17 | 10:25 | Brooding chick | NA | NA |
| 10 | Guillemot | 1b | T15287 | 2535 | 11-Jun-17 | 15:29 | 12-Jun-17 | 10:28 | Brooding chick | NA | NA |
| 11 | Guillemot | 1b | T15288 | 2555 | 11-Jun-17 | 15:32 | 12-Jun-17 | 10:32 | Incubating egg | 2018 | Yes |
| 12 | Guillemot | 1b | T15289 | 2551 | 11-Jun-17 | 15:31 | 12-Jun-17 | 10:35 | Brooding chick | 2018 | Yes |
| 13 | Guillemot | 1b | T15290 | 2554 | 11-Jun-17 | 15:32 | 12-Jun-17 | 10:37 | Brooding chick | 2019 | Yes |
| 14 | Guillemot | 1b | T15291 | 2550 | 11-Jun-17 | 15:30 | 12-Jun-17 | 10:39 | Brooding chick | 2019 | No |
| 15 | Guillemot | 1b | T15292 | 2539 | 11-Jun-17 | 15:26 | 12-Jun-17 | 10:44 | Brooding chick | NA | NA |
| 16 | Guillemot | 1b | T15293 | 2534 | 11-Jun-17 | 15:24 | 12-Jun-17 | 10:46 | Brooding chick | 2018 | Yes |
| 17 | Guillemot | 1b | T15294 | 2545 | 11-Jun-17 | 15:29 | 12-Jun-17 | 10:48 | Brooding chick | 2018 | No |
| 18 | Guillemot | 1b | T15295 | 2538 | 11-Jun-17 | 15:25 | 12-Jun-17 | 10:50 | Incubating egg | 2018 | Yes |
| 19 | Guillemot | 1b | T15296 | 2565 | 11-Jun-17 | 15:38 | 12-Jun-17 | 10:53 | Brooding chick | NA | NA |
| 20 | Guillemot | 1b | T15297 | 2559 | 11-Jun-17 | 15:36 | 12-Jun-17 | 11:11 | Brooding chick | 2018 | Yes |
| 21 | Razorbill | 1b | M78952 | 2641 | 11-Jun-17 | 14:54 | 12-Jun-17 | 10:06 | Brooding chick | NA | NA |
| 22 | Razorbill | 1b | M78954 | 2646 | 11-Jun-17 | 14:57 | 12-Jun-17 | 10:56 | Incubating egg | NA | NA |
| 23 | Razorbill | 1b | M78955 | 2647 | 11-Jun-17 | 14:58 | 12-Jun-17 | 11:00 | Incubating egg | NA | NA |
| 24 | Razorbill | 1b | M78956 | 2645 | 11-Jun-17 | 14:56 | 12-Jun-17 | 11:02 | Brooding chick | NA | NA |
| 25 | Razorbill | 1b | M78957 | 2640 | 11-Jun-17 | 14:54 | 12-Jun-17 | 11:05 | Unknown | 2018 | Yes |
| 26 | Razorbill | 1b | M78958 | 2648 | 11-Jun-17 | 14:58 | 12-Jun-17 | 11:14 | Brooding chick | NA | NA |
| 27 | Razorbill | 1b | M78959 | 2642 | 11-Jun-17 | 14:55 | 12-Jun-17 | 11:21 | Incubating egg | 2019 | Yes |
| 28 | Razorbill | 1b | M78990 | 2643 | 11-Jun-17 | 14:56 | 12-Jun-17 | 15:13 | Brooding chick | 2018 | Yes |
| 29 | Razorbill | 1b | M78991 | 2666 | 11-Jun-17 | 15:06 | 12-Jun-17 | 15:16 | Unknown | NA | NA |
| 30 | Guillemot | 2 | T15298 | 2541 | 11-Jun-17 | 15:27 | 12-Jun-17 | 11:51 | Incubating egg | 2018 | Yes |
| 31 | Guillemot | 2 | T15299 | 2568 | 11-Jun-17 | 15:40 | 12-Jun-17 | 11:54 | Incubating egg | NA | NA |
| 32 | Guillemot | 2 | T15300 | 2529 | 11-Jun-17 | 15:20 | 12-Jun-17 | 11:56 | Brooding chick | 2018 | Yes |
| 33 | Guillemot | 2 | T15301 | 2528 | 11-Jun-17 | 15:19 | 12-Jun-17 | 11:58 | Brooding chick | NA | NA |
| 34 | Guillemot | 2 | T15302 | 2537 | 11-Jun-17 | 15:25 | 12-Jun-17 | 12:00 | Brooding chick | 2018 | Yes |
| 35 | Guillemot | 2 | T15303 | 2543 | 11-Jun-17 | 15:28 | 12-Jun-17 | 12:02 | Incubating egg | 2019 | Yes |
| 36 | Guillemot | 2 | T15304 | 2558 | 11-Jun-17 | 15:34 | 12-Jun-17 | 12:04 | Brooding chick | 2019 | Yes |
| 37 | Guillemot | 2 | T15305 | 2530 | 11-Jun-17 | 15:20 | 12-Jun-17 | 12:06 | Brooding chick | NA | NA |
| 38 | Guillemot | 2 | T15306 | 2557 | 11-Jun-17 | 15:33 | 12-Jun-17 | 12:08 | Brooding chick | NA | NA |
| 39 | Guillemot | 2 | T15307 | 2547 | 11-Jun-17 | 15:29 | 12-Jun-17 | 12:10 | Brooding chick | NA | NA |
| 40 | Guillemot | 2 | T15308 | 2532 | 11-Jun-17 | 15:23 | 12-Jun-17 | 12:14 | Brooding chick | NA | NA |
| 41 | Guillemot | 2 | T15309 | 2548 | 11-Jun-17 | 15:29 | 12-Jun-17 | 12:16 | Brooding chick | NA | NA |
| 42 | Guillemot | 2 | T15310 | 2549 | 11-Jun-17 | 15:30 | 12-Jun-17 | 12:18 | Brooding chick | 2018 | Yes |
| 43 | Guillemot | 2 | T15311 | 2562 | 11-Jun-17 | 15:37 | 12-Jun-17 | 12:20 | Brooding chick | 2018 | Yes |
| 44 | Guillemot | 2 | T15312 | 2540 | 11-Jun-17 | 15:26 | 12-Jun-17 | 12:22 | Brooding chick | NA | NA |
| 45 | Guillemot | 2 | T15313 | 2536 | 11-Jun-17 | 15:24 | 12-Jun-17 | 12:24 | Brooding chick | NA | NA |
| 46 | Guillemot | 2 | T15314 | 2567 | 11-Jun-17 | 15:39 | 12-Jun-17 | 12:26 | Brooding chick | 2019 | Yes |
| 47 | Guillemot | 2 | T15315 | 2566 | 11-Jun-17 | 15:39 | 12-Jun-17 | 12:28 | Brooding chick | 2018 | No |
| 48 | Guillemot | 2 | T15316 | 2553 | 11-Jun-17 | 15:32 | 12-Jun-17 | 12:30 | Brooding chick | NA | NA |
| 49 | Guillemot | 2 | T15317 | 2563 | 11-Jun-17 | 15:37 | 12-Jun-17 | 12:32 | Brooding chick | 2018 | Yes |
| 50 | Razorbill | 3 | M78980 | 2650 | 11-Jun-17 | 15:00 | 12-Jun-17 | 14:12 | Brooding chick | 2018 | Yes |
| 51 | Razorbill | 3 | M78981 | 2657 | 11-Jun-17 | 15:02 | 12-Jun-17 | 14:13 | Brooding chick | 2018 | Yes |
| 52 | Razorbill | 3 | M78982 | 2649 | 11-Jun-17 | 14:59 | 12-Jun-17 | 14:21 | Brooding chick | NA | NA |
| 53 | Razorbill | 3 | M78983 | 2655 | 11-Jun-17 | 15:02 | 12-Jun-17 | 14:24 | Incubating egg | NA | NA |
| 54 | Razorbill | 3 | M78984 | 2662 | 11-Jun-17 | 15:05 | 12-Jun-17 | 14:26 | Brooding chick | 2018 | Yes |
| 55 | Razorbill | 3 | M78985 | 2665 | 11-Jun-17 | 15:06 | 12-Jun-17 | 14:30 | Brooding chick | NA | NA |
| 56 | Razorbill | 3 | M78986 | 2652 | 11-Jun-17 | 15:01 | 12-Jun-17 | 14:37 | Brooding chick | NA | NA |
| 57 | Razorbill | 3 | M78987 | 2663 | 11-Jun-17 | 15:05 | 12-Jun-17 | 14:51 | Brooding chick | NA | NA |
| 58 | Razorbill | 3 | M78988 | 2644 | 11-Jun-17 | 14:56 | 12-Jun-17 | 14:54 | Brooding chick | 2018 | Yes |
| 59 | Razorbill | 3 | M78989 | 2667 | 11-Jun-17 | 15:07 | 12-Jun-17 | 14:58 | Incubating egg | 2018 | Yes |
| 60 | Razorbill | 4 | M72416 | 2656 | 11-Jun-17 | 15:02 | 12-Jun-17 | 10:31 | Brooding chick | NA | NA |
| 61 | Razorbill | 4 | M72417 | 2669 | 11-Jun-17 | 15:07 | 12-Jun-17 | 10:39 | Brooding chick | 2018 | Yes |
| 62 | Razorbill | 4 | M72418 | 2668 | 11-Jun-17 | 15:07 | 12-Jun-17 | 11:00 | Incubating egg | 2018 | Yes |
| 63 | Razorbill | 4 | M72419 | 2658 | 11-Jun-17 | 15:03 | 12-Jun-17 | 11:06 | Incubating egg | NA | NA |
| 64 | Razorbill | 4 | M72421 | 2664 | 11-Jun-17 | 15:05 | 12-Jun-17 | 11:19 | Brooding chick | NA | NA |
| 65 | Razorbill | 4 | M72420 | 2653 | 11-Jun-17 | 15:01 | 12-Jun-17 | 11:25 | Brooding chick | 2018 | Yes |
| 66 | Razorbill | 4 | M72422 | 2661 | 11-Jun-17 | 15:04 | 12-Jun-17 | 11:34 | Brooding chick | 2019 | Yes |
| 67 | Razorbill | 4 | M72423 | 2660 | 11-Jun-17 | 15:04 | 12-Jun-17 | 11:40 | Brooding chick | 2018 | Yes |
| 68 | Razorbill | 4 | M72424 | 2659 | 11-Jun-17 | 15:03 | 12-Jun-17 | 12:13 | Brooding chick | 2018 | Yes |
| 69 | Razorbill | 4 | M72425 | 2654 | 11-Jun-17 | 15:01 | 12-Jun-17 | 12:20 | Incubating egg | NA | NA |
| 70 | Razorbill | 4 | M72426 | 2651 | 11-Jun-17 | 15:00 | 12-Jun-17 | 12:34 | Incubating egg | 2018 | Yes |

Table A1b: GLS logger deployment details for Buchan Ness to Collieston Coast SPA in 2017. Start date and start time denote the start of data collection.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Guillemot | 1 | T23784 | 2587 | 13-Jun-17 | 05:22 | 14-Jun-17 | 18:42 | Brooding chick | 2019 | Yes |
| 2 | Guillemot | 1 | T23785 | 2579 | 13-Jun-17 | 05:18 | 14-Jun-17 | 18:46 | Brooding chick | 2018 | Yes |
| 3 | Guillemot | 1 | T23786 | 2585 | 13-Jun-17 | 05:21 | 14-Jun-17 | 18:50 | Incubating egg | 2018 | Yes |
| 4 | Guillemot | 1 | T70810 | 2606 | 13-Jun-17 | 05:31 | 14-Jun-17 | 18:52 | Brooding chick | 2018 | Yes |
| 5 | Guillemot | 1 | T23787 | 2577 | 13-Jun-17 | 05:18 | 14-Jun-17 | 18:54 | Brooding chick | 2018 | Yes |
| 6 | Guillemot | 1 | T23288 | 2575 | 13-Jun-17 | 05:17 | 14-Jun-17 | 18:57 | Brooding chick | NA | NA |
| 7 | Guillemot | 1 | T23789 | 2571 | 13-Jun-17 | 05:15 | 14-Jun-17 | 19:00 | Brooding chick | 2018 | Yes |
| 8 | Guillemot | 1 | T23790 | 2586 | 13-Jun-17 | 05:22 | 14-Jun-17 | 19:02 | Brooding chick | 2018 | Yes |
| 9 | Guillemot | 1 | T70756 | 2590 | 13-Jun-17 | 05:23 | 14-Jun-17 | 19:04 | Brooding chick | NA | NA |
| 10 | Guillemot | 1 | T23791 | 2569 | 13-Jun-17 | 05:14 | 14-Jun-17 | 19:06 | Brooding chick | 2019 | Yes |
| 11 | Guillemot | 1 | T23793 | 2598 | 13-Jun-17 | 05:27 | 14-Jun-17 | 19:09 | Brooding chick | 2018 | Yes |
| 12 | Guillemot | 1 | T23794 | 2594 | 13-Jun-17 | 05:25 | 14-Jun-17 | 19:13 | Brooding chick | 2018 | Yes |
| 13 | Guillemot | 1 | T23795 | 2602 | 13-Jun-17 | 05:29 | 14-Jun-17 | 19:14 | Brooding chick | 2018 | Yes |
| 14 | Guillemot | 1 | T23796 | 2584 | 13-Jun-17 | 05:21 | 14-Jun-17 | 19:16 | Brooding chick | 2019 | Yes |
| 15 | Guillemot | 1 | T70809 | 2608 | 13-Jun-17 | 05:31 | 14-Jun-17 | 19:18 | Brooding chick | NA | NA |
| 16 | Guillemot | 1 | T23761 | 2592 | 13-Jun-17 | 05:24 | 14-Jun-17 | 19:20 | Brooding chick | NA | NA |
| 17 | Guillemot | 1 | T23797 | 2572 | 13-Jun-17 | 05:15 | 14-Jun-17 | 19:22 | Incubating egg | 2018 | Yes |
| 18 | Guillemot | 1 | T23799 | 2599 | 13-Jun-17 | 05:28 | 14-Jun-17 | 19:25 | Brooding chick | NA | NA |
| 19 | Guillemot | 1 | T23798 | 2588 | 13-Jun-17 | 05:22 | 14-Jun-17 | 19:26 | Brooding chick | 2018 | Yes |
| 20 | Guillemot | 1 | T23800 | 2576 | 13-Jun-17 | 05:17 | 14-Jun-17 | 19:28 | Brooding chick | 2018 | Yes |
| 21 | Guillemot | 1 | T70901 | 2605 | 13-Jun-17 | 05:30 | 14-Jun-17 | 19:30 | Brooding chick | 2018 | Yes |
| 22 | Guillemot | 1 | T70902 | 2607 | 13-Jun-17 | 05:31 | 14-Jun-17 | 19:32 | Brooding chick | 2019 | Yes |
| 23 | Guillemot | 1 | T70903 | 2593 | 13-Jun-17 | 05:25 | 14-Jun-17 | 20:05 | Brooding chick | 2018 | Yes |
| 24 | Guillemot | 1 | T70904 | 2580 | 13-Jun-17 | 05:19 | 14-Jun-17 | 20:19 | Brooding chick | 2018 | Yes |
| 25 | Guillemot | 1 | T70906 | 2595 | 13-Jun-17 | 05:26 | 14-Jun-17 | 20:21 | Brooding chick | 2018 | Yes |
| 26 | Guillemot | 1 | T70905 | 2601 | 13-Jun-17 | 05:29 | 14-Jun-17 | 20:23 | Brooding chick | 2018 | Yes |
| 27 | Guillemot | 1 | T70908 | 2581 | 13-Jun-17 | 05:19 | 14-Jun-17 | 20:26 | Brooding chick | NA | NA |
| 28 | Guillemot | 1 | T70907 | 2589 | 13-Jun-17 | 05:23 | 14-Jun-17 | 20:28 | Brooding chick | NA | NA |
| 29 | Guillemot | 1 | T70961 | 2573 | 13-Jun-17 | 05:16 | 14-Jun-17 | 20:29 | Brooding chick | 2018 | Yes |
| 30 | Guillemot | 1 | T70909 | 2578 | 13-Jun-17 | 05:18 | 14-Jun-17 | 20:31 | Brooding chick | NA | NA |
| 31 | Guillemot | 1 | T23778 | 2604 | 13-Jun-17 | 05:30 | 14-Jun-17 | 20:33 | Brooding chick | 2018 | Yes |
| 32 | Guillemot | 1 | T23776 | 2600 | 13-Jun-17 | 05:28 | 14-Jun-17 | 20:34 | Brooding chick | NA | NA |
| 33 | Guillemot | 1 | T70962 | 2597 | 13-Jun-17 | 05:27 | 14-Jun-17 | 20:36 | Brooding chick | 2018 | Yes |
| 34 | Guillemot | 1 | T70910 | 2596 | 13-Jun-17 | 05:26 | 14-Jun-17 | 20:38 | Brooding chick | NA | NA |
| 35 | Guillemot | 1 | T70911 | 2603 | 13-Jun-17 | 05:29 | 14-Jun-17 | 20:39 | Brooding chick | 2018 | Yes |
| 36 | Guillemot | 1 | T70956 | 2582 | 13-Jun-17 | 05:20 | 14-Jun-17 | 20:41 | Brooding chick | 2018 | Yes |
| 37 | Guillemot | 1 | T23764 | 2574 | 13-Jun-17 | 05:16 | 14-Jun-17 | 20:43 | Brooding chick | 2018 | Yes |
| 38 | Guillemot | 1 | T70912 | 2570 | 13-Jun-17 | 05:15 | 14-Jun-17 | 20:45 | Brooding chick | 2018 | Yes |
| 39 | Guillemot | 1 | T70913 | 2591 | 13-Jun-17 | 05:24 | 14-Jun-17 | 20:46 | Brooding chick | NA | NA |
| 40 | Guillemot | 1 | T70914 | 2583 | 13-Jun-17 | 05:20 | 14-Jun-17 | 20:48 | Brooding chick | NA | NA |
| 41 | Razorbill | 2 | M88601 | 2678 | 13-Jun-17 | 05:04 | 15-Jun-17 | 10:02 | Incubating egg | NA | NA |
| 42 | Razorbill | 3 | M88602 | 2687 | 13-Jun-17 | 05:08 | 23-Jun-17 | 19:34 | Unknown | NA | NA |
| 43 | Razorbill | 3 | M88603 | 2688 | 13-Jun-17 | 05:08 | 23-Jun-17 | 19:32 | Brooding chick | 2019 | Yes |
| 44 | Razorbill | 3 | M88604 | 2684 | 13-Jun-17 | 05:06 | 23-Jun-17 | 19:43 | Brooding chick | NA | NA |
| 45 | Razorbill | 3 | M88605 | 2671 | 13-Jun-17 | 05:01 | 23-Jun-17 | 19:54 | Incubating egg | NA | NA |
| 46 | Razorbill | 3 | M88606 | 2676 | 13-Jun-17 | 05:03 | 23-Jun-17 | 19:57 | Brooding chick | NA | NA |
| 47 | Razorbill | 3 | M88607 | 2681 | 13-Jun-17 | 05:05 | 23-Jun-17 | 20:00 | Brooding chick | NA | NA |
| 48 | Razorbill | 3 | M88608 | 2683 | 13-Jun-17 | 05:06 | 23-Jun-17 | 20:05 | Brooding chick | 2019 | Yes |
| 49 | Razorbill | 3 | M88609 | 2679 | 13-Jun-17 | 05:02 | 23-Jun-17 | 20:09 | Brooding chick | 2018 | Yes |
| 50 | Razorbill | 3 | M88610 | 2677 | 13-Jun-17 | 05:04 | 23-Jun-17 | 20:18 | Incubating egg | 2019 | Yes |
| 51 | Razorbill | 3 | M88611 | 2680 | 13-Jun-17 | 05:05 | 23-Jun-17 | 20:25 | Brooding chick | NA | NA |
| 52 | Razorbill | 3 | M88612 | 2675 | 13-Jun-17 | 05:03 | 23-Jun-17 | 20:38 | Brooding chick | 2018 | Yes |
| 53 | Razorbill | 3 | M88613 | 2674 | 13-Jun-17 | 05:02 | 23-Jun-17 | 20:43 | Brooding chick | NA | NA |
| 54 | Razorbill | 3 | M88614 | 2672 | 13-Jun-17 | 05:02 | 23-Jun-17 | 20:54 | Brooding chick | NA | NA |
| 55 | Razorbill | 3 | M88615 | 2689 | 13-Jun-17 | 05:09 | 23-Jun-17 | 21:05 | Brooding chick | NA | NA |
| 56 | Razorbill | 2 | M88616 | 2686 | 13-Jun-17 | 05:07 | 24-Jun-17 | 14:05 | Brooding chick | NA | NA |
| 57 | Razorbill | 4 | M88617 | 2673 | 13-Jun-17 | 05:02 | 24-Jun-17 | 16:02 | Brooding chick | NA | NA |
| 58 | Razorbill | 4 | M88618 | 2682 | 13-Jun-17 | 05:06 | 24-Jun-17 | 16:05 | Incubating egg | NA | NA |
| 59 | Razorbill | 4 | M88619 | 2685 | 13-Jun-17 | 05:07 | 24-Jun-17 | 16:27 | Brooding chick | NA | NA |
| 60 | Razorbill | 4 | M88620 | 2670 | 13-Jun-17 | 05:00 | 24-Jun-17 | 16:41 | Brooding chick | NA | NA |

Table A1c: GLS logger deployment details for Isle of May NNR in 2017. Start date and start time denote the start of data collection.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Razorbill | 1 | M82449 | 2614 | 08-Jun-17 | 13:43 | 20-Jun-17 | 20:00 | Brooding chick | 2018 | Yes |
| 2 | Razorbill | 2 | M82440 | 2622 | 08-Jun-17 | 13:47 | 19-Jun-17 | 11:05 | Brooding chick | 2018 | Yes |
| 3 | Razorbill | 3 | M84627 | 2609 | 08-Jun-17 | 13:40 | 14-Jun-17 | 09:00 | Brooding chick | NA | NA |
| 4 | Razorbill | 4 | M84628 | 2611 | 08-Jun-17 | 13:42 | 17-Jun-17 | 10:40 | Brooding chick | NA | NA |
| 5 | Razorbill | 4 | M84629 | 2610 | 08-Jun-17 | 13:41 | 17-Jun-17 | 10:50 | Brooding chick | NA | NA |
| 6 | Razorbill | 4 | M84630 | 2612 | 08-Jun-17 | 13:42 | 17-Jun-17 | 11:02 | Brooding chick | 2018 | Yes |
| 7 | Razorbill | 4 | M84631 | 2613 | 08-Jun-17 | 13:43 | 19-Jun-17 | 10:40 | Brooding chick | NA | NA |
| 8 | Razorbill | 1 | M84632 | 2615 | 08-Jun-17 | 13:44 | 19-Jun-17 | 11:13 | Brooding chick | NA | NA |
| 9 | Razorbill | 1 | M84633 | 2616 | 08-Jun-17 | 13:45 | 19-Jun-17 | 11:23 | Brooding chick | 2018 | Yes |
| 10 | Razorbill | 3 | M84634 | 2617 | 08-Jun-17 | 13:45 | 19-Jun-17 | 11:41 | Brooding chick | NA | NA |
| 11 | Razorbill | 3 | M84635 | 2618 | 08-Jun-17 | 13:45 | 19-Jun-17 | 11:50 | Brooding chick | NA | NA |
| 12 | Razorbill | 5 | M84636 | 2619 | 08-Jun-17 | 13:46 | 20-Jun-17 | 08:45 | Brooding chick | 2018 | Yes |
| 13 | Razorbill | 5 | M84637 | 2620 | 08-Jun-17 | 13:46 | 20-Jun-17 | 09:00 | Brooding chick | NA | NA |
| 14 | Razorbill | 3 | M84640 | 2621 | 08-Jun-17 | 13:46 | 20-Jun-17 | 09:20 | Brooding chick | NA | NA |
| 15 | Razorbill | 2 | M84641 | 2623 | 08-Jun-17 | 13:47 | 20-Jun-17 | 20:30 | Brooding chick | 2018 | Yes |
| 16 | Razorbill | 2 | M84642 | 2624 | 08-Jun-17 | 13:48 | 20-Jun-17 | 20:40 | Brooding chick | NA | NA |
| 17 | Razorbill | 3 | M84643 | 2625 | 08-Jun-17 | 13:48 | 21-Jun-17 | 10:30 | Brooding chick | 2018 | Yes |
| 18 | Razorbill | 6 | M84644 | 2626 | 08-Jun-17 | 13:48 | 21-Jun-17 | 16:40 | Brooding chick | 2018 | Yes |
| 19 | Razorbill | 6 | M84645 | 2627 | 08-Jun-17 | 13:49 | 21-Jun-17 | 16:50 | Brooding chick | NA | NA |
| 20 | Razorbill | 2 | M84646 | 2628 | 08-Jun-17 | 13:49 | 21-Jun-17 | 17:10 | Brooding chick | NA | NA |
| 21 | Razorbill | 7 | M84647 | 2629 | 08-Jun-17 | 13:50 | 21-Jun-17 | 17:30 | Brooding chick | 2018 | Yes |
| 22 | Razorbill | 8 | M84648 | 2630 | 08-Jun-17 | 13:50 | 22-Jun-17 | 08:10 | Brooding chick | NA | NA |
| 23 | Razorbill | 8 | M84649 | 2631 | 08-Jun-17 | 13:50 | 22-Jun-17 | 08:20 | Brooding chick | NA | NA |
| 24 | Razorbill | 8 | M84650 | 2632 | 08-Jun-17 | 13:51 | 22-Jun-17 | 08:30 | Brooding chick | NA | NA |
| 25 | Razorbill | 6 | M84651 | 2633 | 08-Jun-17 | 13:51 | 22-Jun-17 | 08:40 | Brooding chick | NA | NA |
| 26 | Razorbill | 1 | M84652 | 2634 | 08-Jun-17 | 13:52 | 22-Jun-17 | 17:02 | Brooding chick | 2018 | Yes |
| 27 | Razorbill | 1 | M84653 | 2635 | 08-Jun-17 | 13:52 | 22-Jun-17 | 17:17 | Brooding chick | NA | NA |
| 28 | Razorbill | 1 | M84654 | 2636 | 08-Jun-17 | 13:52 | 22-Jun-17 | 17:24 | Brooding chick | 2018 | Yes |
| 29 | Razorbill | 4 | M84655 | 2639 | 08-Jun-17 | 13:53 | 23-Jun-17 | 09:00 | Brooding chick | NA | NA |
| 30 | Razorbill | 6 | M84656 | 2637 | 08-Jun-17 | 13:53 | 23-Jun-17 | 09:30 | Brooding chick | NA | NA |

Table A2a: geolocator deployment details for East Caithness in 2018. Start date and start time denote the start of data collection.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Guillemot | 1a | R99314 | B3853 | 12-Jun-18 | 19:23 | 17-Jun-18 | 12:04 | Incubating egg | 2019 | Yes |
| 2 | Guillemot | 1b | R99315 | B3850 | 12-Jun-18 | 19:21 | 17-Jun-18 | 14:52 | Unknown | 2019 | Yes |
| 3 | Guillemot | 1b | R99316 | B3818 | 12-Jun-18 | 19:07 | 17-Jun-18 | 10:48 | Incubating egg | 2019 | Yes |
| 4 | Guillemot | 1b | R99317 | B3827 | 12-Jun-18 | 19:13 | 17-Jun-18 | 10:59 | Brooding chick | NA | NA |
| 5 | Guillemot | 1b | R99318 | B3830 | 12-Jun-18 | 19:14 | 18-Jun-18 | 16:25 | Brooding chick | 2019 | Yes |
| 6 | Guillemot | 1b | R99319 | B3832 | 12-Jun-18 | 19:15 | 17-Jun-18 | 11:46 | Brooding chick | 2019 | Yes |
| 7 | Guillemot | 1b | R99320 | B3851 | 12-Jun-18 | 19:22 | 17-Jun-18 | 11:40 | Brooding chick | 2019 | Yes |
| 8 | Guillemot | 1b | T15288 | B3843 | 12-Jun-18 | 19:19 | 17-Jun-18 | 11:13 | Brooding chick | 2019 | Yes |
| 9 | Guillemot | 1b | T15289 | B3823 | 12-Jun-18 | 19:11 | 17-Jun-18 | 11:07 | Incubating egg | NA | NA |
| 10 | Guillemot | 1b | T15293 | B3811 | 12-Jun-18 | 19:06 | 17-Jun-18 | 12:08 | Incubating egg | NA | NA |
| 11 | Guillemot | 1b | T15294 | B3847 | 12-Jun-18 | 19:21 | 17-Jun-18 | 11:18 | Unknown | NA | NA |
| 12 | Guillemot | 1b | T15295 | B3834 | 12-Jun-18 | 19:16 | 17-Jun-18 | 10:52 | Incubating egg | NA | NA |
| 13 | Guillemot | 1b | T15297 | B3824 | 12-Jun-18 | 19:12 | 17-Jun-18 | 11:21 | Brooding chick | 2019 | Yes |
| 14 | Razorbill | 1b | M78957 | BN100 | 13-Jun-18 | 06:56 | 18-Jun-18 | 11:12 | Brooding chick | 2019 | Yes |
| 15 | Razorbill | 1b | M78990 | BN097 | 13-Jun-18 | 04:14 | 17-Jun-18 | 14:48 | Brooding chick | NA | NA |
| 16 | Guillemot | 2 | T15298 | B3829 | 12-Jun-18 | 19:14 | 18-Jun-18 | 15:30 | Brooding chick | NA | NA |
| 17 | Guillemot | 2 | T15300 | B3835 | 12-Jun-18 | 19:16 | 18-Jun-18 | 15:05 | Incubating egg | NA | NA |
| 18 | Guillemot | 2 | T15302 | B3837 | 12-Jun-18 | 19:17 | 17-Jun-18 | 13:37 | Incubating egg | 2019 | Yes |
| 19 | Guillemot | 2 | T15310 | B3840 | 12-Jun-18 | 19:18 | 18-Jun-18 | 13:55 | Incubating egg | NA | NA |
| 20 | Guillemot | 2 | T15311 | B3842 | 12-Jun-18 | 19:18 | 19-Jun-18 | 12:55 | Brooding chick | NA | NA |
| 21 | Guillemot | 2 | T15315 | B3848 | 12-Jun-18 | 19:21 | 18-Jun-18 | 13:41 | Unknown | 2019 | Yes |
| 22 | Guillemot | 2 | T15317 | B3817 | 12-Jun-18 | 19:09 | 18-Jun-18 | 15:50 | Brooding chick | NA | NA |
| 23 | Razorbill | 3 | M78980 | BN055 | 13-Jun-18 | 05:51 | 17-Jun-18 | 13:18 | Brooding chick | 2019 | Yes |
| 24 | Razorbill | 3 | M78981 | BN058 | 13-Jun-18 | 05:53 | 17-Jun-18 | 13:13 | Brooding chick | NA | NA |
| 25 | Razorbill | 3 | M78984 | BN101 | 13-Jun-18 | 07:44 | 19-Jun-18 | 13:17 | Brooding chick | 2019 | Yes |
| 26 | Razorbill | 3 | M78988 | BN84 | 13-Jun-18 | 06:55 | 17-Jun-18 | 14:18 | Unknown | NA | NA |
| 27 | Razorbill | 3 | M78989 | BN91 | 13-Jun-18 | 07:00 | 18-Jun-18 | 12:54 | Brooding chick | NA | NA |
| 28 | Razorbill | Boat | M72417 | BN53 | 13-Jun-18 | 06:49 | 02-Jul-18 | 09:45 | Unknown | NA | NA |
| 29 | Razorbill | Boat | M72418 | BN86 | 13-Jun-18 | 06:57 | 02-Jul-18 | 10:21 | Unknown | NA | NA |
| 30 | Razorbill | Boat | M72420 | BN93 | 13-Jun-18 | 07:01 | 02-Jul-18 | 11:02 | Unknown | NA | NA |
| 31 | Razorbill | Boat | M72423 | BN88 | 13-Jun-18 | 06:58 | 02-Jul-18 | 10:41 | Unknown | NA | NA |
| 32 | Razorbill | Boat | M72424 | BN59 | 13-Jun-18 | 06:53 | 02-Jul-18 | 10:03 | Unknown | 2019 | Yes |
| 33 | Guillemot | 1b | X83633 | B3821 | 12-Jun-18 | 19:10 | 19-Jun-18 | 10:48 | Incubating egg | 2019 | Yes |
| 34 | Guillemot | 1b | X83637 | B3820 | 12-Jun-18 | 19:10 | 19-Jun-18 | 10:53 | Incubating egg | 2019 | Yes |
| 35 | Guillemot | 1b | X83639 | B3852 | 12-Jun-18 | 19:22 | 19-Jun-18 | 11:26 | Incubating egg | 2019 | Yes |
| 36 | Guillemot | 1b | X83664 | B3831 | 12-Jun-18 | 19:14 | 19-Jun-18 | 11:44 | Incubating egg | 2019 | Yes |
| 37 | Guillemot | 1b | X83675 | B3846 | 12-Jun-18 | 19:20 | 19-Jun-18 | 10:43 | Unknown | NA | NA |
| 38 | Guillemot | 1b | X83676 | B3819 | 12-Jun-18 | 19:09 | 19-Jun-18 | 10:59 | Unknown | 2019 | Yes |
| 39 | Guillemot | 1b | X83677 | B3814 | 12-Jun-18 | 19:07 | 19-Jun-18 | 11:04 | Brooding chick | 2019 | Yes |
| 40 | Guillemot | 1b | X83678 | B3844 | 12-Jun-18 | 19:19 | 19-Jun-18 | 11:08 | Brooding chick | 2019 | Yes |
| 41 | Guillemot | 1b | X83679 | B3838 | 12-Jun-18 | 19:17 | 19-Jun-18 | 11:13 | Brooding chick | NA | NA |
| 42 | Guillemot | 1b | X83680 | B3841 | 12-Jun-18 | 19:18 | 19-Jun-18 | 11:17 | Incubating egg | NA | NA |
| 43 | Guillemot | 1b | X83681 | B3815 | 12-Jun-18 | 19:08 | 19-Jun-18 | 11:21 | Brooding chick | 2019 | Yes |
| 44 | Guillemot | 1b | X83682 | B3833 | 12-Jun-18 | 19:15 | 19-Jun-18 | 11:31 | Brooding chick | 2019 | Yes |
| 45 | Guillemot | 1b | X83683 | B3845 | 12-Jun-18 | 19:20 | 19-Jun-18 | 11:35 | Brooding chick | 2019 | Yes |
| 46 | Guillemot | 1b | X83684 | B3822 | 12-Jun-18 | 19:11 | 19-Jun-18 | 11:40 | Brooding chick | NA | NA |
| 47 | Guillemot | 1b | X83685 | B3816 | 12-Jun-18 | 19:08 | 19-Jun-18 | 11:49 | Incubating egg | NA | NA |
| 48 | Guillemot | 2 | X83687 | B3836 | 12-Jun-18 | 19:16 | 19-Jun-18 | 12:41 | Incubating egg | 2019 | Yes |
| 49 | Guillemot | 2 | X83688 | B3812 | 12-Jun-18 | 19:06 | 19-Jun-18 | 12:45 | Incubating egg | NA | NA |
| 50 | Guillemot | 2 | X83689 | B3828 | 12-Jun-18 | 19:13 | 19-Jun-18 | 12:48 | Brooding chick | NA | NA |
| 51 | Guillemot | 2 | X83700 | B3825 | 12-Jun-18 | 19:12 | 19-Jun-18 | 16:06 | Brooding chick | 2019 | Yes |
| 52 | Razorbill | 3 | M89255 | BN57 | 13-Jun-18 | 06:52 | 18-Jun-18 | 11:45 | Brooding chick | NA | NA |
| 53 | Razorbill | 3 | M89256 | BN99 | 13-Jun-18 | 07:05 | 18-Jun-18 | 11:55 | Incubating egg | NA | NA |
| 54 | Razorbill | 3 | M89257 | BN92 | 13-Jun-18 | 07:01 | 18-Jun-18 | 12:05 | Brooding chick | NA | NA |
| 55 | Razorbill | 3 | M89259 | BN98 | 13-Jun-18 | 07:04 | 18-Jun-18 | 12:19 | Brooding chick | NA | NA |
| 56 | Razorbill | 3 | M89260 | BN85 | 13-Jun-18 | 06:56 | 18-Jun-18 | 12:25 | Incubating egg | NA | NA |
| 57 | Razorbill | 3 | M89261 | BN90 | 13-Jun-18 | 06:59 | 18-Jun-18 | 12:35 | Brooding chick | NA | NA |
| 58 | Razorbill | 3 | M89263 | BN52 | 13-Jun-18 | 06:49 | 18-Jun-18 | 13:08 | Brooding chick | NA | NA |
| 59 | Razorbill | 3 | M89264 | BN51 | 13-Jun-18 | 06:48 | 18-Jun-18 | 14:14 | Brooding chick | NA | NA |
| 60 | Razorbill | 3 | M89265 | BN96 | 13-Jun-18 | 07:03 | 18-Jun-18 | 14:20 | Brooding chick | 2019 | Yes |
| 61 | Razorbill | 3 | M89271 | BN56 | 13-Jun-18 | 06:51 | 19-Jun-18 | 13:56 | Unknown | NA | NA |
| 62 | Razorbill | 3 | M89273 | BN95 | 13-Jun-18 | 07:02 | 19-Jun-18 | 14:04 | Incubating egg | NA | NA |
| 63 | Razorbill | 3 | M89274 | BN102 | 13-Jun-18 | 07:08 | 19-Jun-18 | 14:08 | Brooding chick | NA | NA |
| 64 | Razorbill | 3 | M89277 | BN87 | 13-Jun-18 | 06:57 | 19-Jun-18 | 14:26 | Incubating egg | NA | NA |
| 65 | Razorbill | 3 | M89283 | BN83 | 13-Jun-18 | 06:54 | 19-Jun-18 | 14:51 | Unknown | NA | NA |
| 66 | Razorbill | 3 | M89284 | BN89 | 13-Jun-18 | 06:59 | 19-Jun-18 | 14:57 | Brooding chick | NA | NA |
| 67 | Razorbill | 3 | M89254 | BN50 | 13-Jun-18 | 06:47 | 17-Jun-18 | 15:25 | Brooding chick | NA | NA |
| 68 | Razorbill | 3b | M89258 | BN94 | 13-Jun-18 | 07:02 | 18-Jun-18 | 12:13 | Brooding chick | NA | NA |
| 69 | Razorbill | 3b | M89262 | BN54 | 13-Jun-18 | 06:50 | 18-Jun-18 | 12:45 | Brooding chick | NA | NA |

Table A2b: geolocator deployment details for Buchan Ness to Collieston Coast in 2018. Start date and start time denote the start of data collection.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Guillemot | | T23762 | B3854 | 12-Jun-18 | 19:23 | 19-Jun-18 | 19:53 | Unknown | NA | NA |
| 2 | Guillemot | 1 | T23764 | B3869 | 12-Jun-18 | 19:30 | 23-Jun-18 | 11:40 | Brooding chick | NA | NA |
| 3 | Guillemot | | T23769 | B3879 | 12-Jun-18 | 20:34 | 24-Jun-18 | 18:22 | Brooding chick | NA | NA |
| 4 | Guillemot | | T23771 | B3867 | 12-Jun-18 | 19:29 | 24-Jun-18 | 18:17 | Brooding chick | 2019 | Yes |
| 5 | Guillemot | | T23777 | B3881 | 12-Jun-18 | 20:35 | 24-Jun-18 | 18:19 | Brooding chick | 2019 | Yes |
| 6 | Guillemot | 1 | T23778 | B3872 | 12-Jun-18 | 19:31 | 24-Jun-18 | 16:41 | Brooding chick | NA | NA |
| 7 | Guillemot | 1 | T23785 | B3883 | 12-Jun-18 | 20:36 | 23-Jun-18 | 14:10 | Unknown | 2019 | Yes |
| 8 | Guillemot | 1 | T23786 | B3874 | 12-Jun-18 | 20:32 | 23-Jun-18 | 11:28 | Brooding chick | 2019 | Yes |
| 9 | Guillemot | 1 | T23787 | B3890 | 12-Jun-18 | 20:39 | 23-Jun-18 | 14:16 | Brooding chick | 2019 | Yes |
| 10 | Guillemot | 1 | T23789 | B3858 | 12-Jun-18 | 19:25 | 24-Jun-18 | 17:11 | Brooding chick | NA | NA |
| 11 | Guillemot | 1 | T23790 | B3875 | 12-Jun-18 | 20:33 | 23-Jun-18 | 13:43 | Brooding chick | NA | NA |
| 12 | Guillemot | 1 | T23793 | B3855 | 12-Jun-18 | 19:24 | 23-Jun-18 | 10:49 | Brooding chick | 2019 | Yes |
| 13 | Guillemot | 1 | T23794 | B3861 | 12-Jun-18 | 19:26 | 19-Jun-18 | 19:51 | Unknown | NA | NA |
| 14 | Guillemot | 1 | T23795 | B3862 | 12-Jun-18 | 19:27 | 19-Jun-18 | 19:41 | Brooding chick | 2019 | No |
| 15 | Guillemot | 1 | T23797 | B3868 | 12-Jun-18 | 19:29 | 19-Jun-18 | 19:22 | Brooding chick | 2019 | Yes |
| 16 | Guillemot | 1 | T23798 | B3876 | 12-Jun-18 | 20:33 | 23-Jun-18 | 13:59 | Brooding chick | NA | NA |
| 17 | Guillemot | 1 | T23800 | B3856 | 12-Jun-18 | 19:24 | 19-Jun-18 | 20:00 | Brooding chick | 2019 | Yes |
| 18 | Guillemot | | T70757 | B3887 | 12-Jun-18 | 20:38 | 23-Jun-18 | 12:56 | Brooding chick | 2019 | Yes |
| 19 | Guillemot | 1 | T70810 | B3864 | 12-Jun-18 | 19:27 | 19-Jun-18 | 19:20 | Brooding chick | 2019 | Yes |
| 20 | Guillemot | | T70818 | B3889 | 12-Jun-18 | 20:39 | 24-Jun-18 | 18:13 | Brooding chick | 2019 | Yes |
| 21 | Guillemot | 1 | T70901 | B3857 | 12-Jun-18 | 19:24 | 19-Jun-18 | 20:03 | Unknown | NA | NA |
| 22 | Guillemot | 1 | T70903 | B3880 | 12-Jun-18 | 20:35 | 23-Jun-18 | 13:23 | Brooding chick | 2019 | Yes |
| 23 | Guillemot | 1 | T70904 | B3866 | 12-Jun-18 | 19:28 | 19-Jun-18 | 19:45 | Brooding chick | 2019 | Yes |
| 24 | Guillemot | 1 | T70905 | B3860 | 12-Jun-18 | 19:26 | 19-Jun-18 | 19:49 | Unknown | 2019 | Yes |
| 25 | Guillemot | 1 | T70906 | B3888 | 12-Jun-18 | 20:38 | 24-Jun-18 | 17:37 | Brooding chick | 2019 | Yes |
| 26 | Guillemot | 1 | T70911 | B3877 | 12-Jun-18 | 20:33 | 23-Jun-18 | 13:35 | Brooding chick | NA | NA |
| 27 | Guillemot | | T70915 | B3878 | 12-Jun-18 | 20:34 | 23-Jun-18 | 11:42 | Brooding chick | 2019 | Yes |
| 28 | Guillemot | | T70916 | B3882 | 12-Jun-18 | 19:36 | 24-Jun-18 | 16:53 | Brooding chick | NA | NA |
| 29 | Guillemot | | T70917 | B3863 | 12-Jun-18 | 19:27 | 24-Jun-18 | 17:33 | Brooding chick | 2019 | Yes |
| 30 | Guillemot | | T70918 | B3865 | 12-Jun-18 | 19:28 | 24-Jun-18 | 18:06 | Brooding chick | 2019 | Yes |
| 31 | Guillemot | | T70919 | B3870 | 12-Jun-18 | 19:30 | 24-Jun-18 | 18:10 | Brooding chick | NA | NA |
| 32 | Guillemot | | T70920 | B3893 | 12-Jun-18 | 20:40 | 24-Jun-18 | 18:20 | Brooding chick | NA | NA |
| 33 | Guillemot | | T70921 | B3886 | 12-Jun-18 | 20:37 | 24-Jun-18 | 18:25 | Brooding chick | 2019 | Yes |
| 34 | Guillemot | | T70922 | B3891 | 12-Jun-18 | 20:39 | 24-Jun-18 | 18:27 | Brooding chick | NA | NA |
| 35 | Guillemot | | T70923 | B3884 | 12-Jun-18 | 20:36 | 24-Jun-18 | 18:29 | Brooding chick | NA | NA |
| 36 | Guillemot | 1 | T70956 | B3873 | 12-Jun-18 | 19:32 | 23-Jun-18 | 11:59 | Brooding chick | 2019 | Yes |
| 37 | Guillemot | | T70958 | B3885 | 12-Jun-18 | 20:37 | 24-Jun-18 | 18:31 | Brooding chick | 2019 | Yes |
| 38 | Guillemot | | T70960 | B3892 | 12-Jun-18 | 20:40 | 24-Jun-18 | 18:08 | Brooding chick | NA | NA |
| 39 | Guillemot | 1 | T70961 | B3871 | 12-Jun-18 | 19:31 | 19-Jun-18 | 20:16 | Brooding chick | 2019 | Yes |
| 40 | Guillemot | 1 | T70962 | B3859 | 12-Jun-18 | 19:25 | 23-Jun-18 | 11:06 | Brooding chick | NA | NA |
| 41 | Razorbill | 3 | M88609 | BN41 | 13-Jun-18 | 05:59 | 24-Jun-18 | 20:38 | Brooding chick | NA | NA |
| 42 | Razorbill | 4 | M88621 | BN48 | 13-Jun-18 | 06:04 | 24-Jun-18 | 19:07 | Brooding chick | NA | NA |
| 43 | Razorbill | 4 | M88622 | BN39 | 13-Jun-18 | 05:57 | 24-Jun-18 | 19:27 | Brooding chick | NA | NA |
| 44 | Razorbill | 4 | M88623 | BN36 | 13-Jun-18 | 05:55 | 24-Jun-18 | 19:30 | Brooding chick | 2019 | Yes |
| 45 | Razorbill | 3 | M88624 | BN49 | 13-Jun-18 | 06:04 | 24-Jun-18 | 20:38 | Brooding chick | NA | NA |
| 46 | Razorbill | 4 | M88625 | BN30 | 13-Jun-18 | 05:50 | 01-Jul-18 | 17:58 | Brooding chick | NA | NA |
| 47 | Razorbill | 3 | M88626 | BN31 | 13-Jun-18 | 05:51 | 01-Jul-18 | 19:05 | Brooding chick | NA | NA |
| 48 | Razorbill | 3 | M88627 | BN32 | 13-Jun-18 | 05:52 | 01-Jul-18 | 19:30 | Brooding chick | NA | NA |
| 49 | Razorbill | 3 | M88628 | BN33 | 13-Jun-18 | 05:53 | 01-Jul-18 | 20:03 | Brooding chick | NA | NA |
| 50 | Razorbill | 4 | M88629 | BN34 | 13-Jun-18 | 05:54 | 08-Jul-18 | 14:10 | Brooding chick | NA | NA |
| 51 | Razorbill | 4 | M88630 | BN45 | 13-Jun-18 | 06:01 | 17-Jul-18 | 12:00 | Brooding chick | NA | NA |
| 52 | Razorbill | 4 | M88631 | BN37 | 13-Jun-18 | 05:56 | 17-Jul-18 | 12:49 | Brooding chick | 2019 | Yes |
| 53 | Razorbill | 4 | M88632 | BN47 | 13-Jun-18 | 06:03 | 17-Jul-18 | 13:04 | Brooding chick | 2019 | Yes |
| 54 | Razorbill | 4 | M88633 | BN44 | 13-Jun-18 | 06:01 | 17-Jul-18 | 13:22 | Brooding chick | NA | NA |
| 55 | Razorbill | 4 | M88634 | BN42 | 13-Jun-18 | 06:00 | 17-Jul-18 | 14:56 | Brooding chick | NA | NA |
| 56 | Razorbill | 4 | M88635 | BN35 | 13-Jun-18 | 05:54 | 19-Jul-18 | 13:50 | Brooding chick | NA | NA |
| 57 | Razorbill | 4 | M88636 | BN40 | 13-Jun-18 | 05:58 | 19-Jul-18 | 15:05 | Brooding chick | NA | NA |
| 58 | Razorbill | 4 | M88637 | BN46 | 13-Jun-18 | 06:02 | 19-Jul-18 | 15:42 | Brooding chick | 2019 | Yes |
| 59 | Razorbill | 3 | M88638 | BN38 | 13-Jun-18 | 05:57 | 19-Jul-18 | 17:55 | Unknown | 2019 | Yes |

Table A2c: geolocator deployment details for Isle of May in 2018. Start date and start time denote the start of data collection. Deployments on guillemots were carried out as part of the SEATRACK project.

| n | Species | Site | BTO ring | Logger ID | Start date | Start time | Deployment date | Deployment Time | Nest Status | Year retrieved | Deployment successful |
|----|-----------|------|----------|-----------|------------|------------|-----------------|-----------------|----------------|----------------|-----------------------|
| 1 | Guillemot | 3 | R76644 | B3472 | 16-Apr-18 | 14:50 | 27-Jun-18 | 12:00 | Brooding chick | NA | NA |
| 2 | Guillemot | 3 | R03499 | B3473 | 16-Apr-18 | 14:51 | 27-Jun-18 | 12:00 | Brooding chick | 2019 | Yes |
| 3 | Guillemot | 3 | Y09073 | B3474 | 16-Apr-18 | 14:51 | 27-Jun-18 | 12:00 | Brooding chick | 2019 | Yes |
| 4 | Guillemot | 9 | R76260 | B3475 | 16-Apr-18 | 14:52 | 27-Jun-18 | 12:00 | Brooding chick | 2019 | Yes |
| 5 | Guillemot | 9 | R76638 | B3476 | 16-Apr-18 | 14:52 | 27-Jun-18 | 12:00 | Brooding chick | 2019 | Yes |
| 6 | Guillemot | 3 | X80054 | B3477 | 16-Apr-18 | 14:53 | 01-Jul-18 | 11:00 | Brooding chick | 2019 | Yes |
| 7 | Guillemot | 3 | Y09105 | B3478 | 16-Apr-18 | 14:53 | 01-Jul-18 | 11:00 | Brooding chick | NA | NA |
| 8 | Guillemot | 3 | R02615 | B3479 | 16-Apr-18 | 14:54 | 01-Jul-18 | 11:00 | Brooding chick | 2019 | Yes |
| 9 | Guillemot | 9 | R76090 | B3480 | 16-Apr-18 | 14:54 | 04-Jul-18 | 11:00 | Brooding chick | NA | NA |
| 10 | Guillemot | 9 | Y09072 | B3481 | 16-Apr-18 | 14:55 | 04-Jul-18 | 11:00 | Brooding chick | NA | NA |
| 11 | Guillemot | 9 | X80140 | B3482 | 16-Apr-18 | 14:55 | 04-Jul-18 | 15:00 | Brooding chick | 2019 | Yes |
| 12 | Guillemot | 9 | X53431 | B3483 | 16-Apr-18 | 14:56 | 05-Jul-18 | 09:00 | Brooding chick | NA | NA |
| 13 | Guillemot | 9 | Y09008 | B3484 | 16-Apr-18 | 14:56 | 05-Jul-18 | 10:30 | Brooding chick | 2019 | Yes |
| 14 | Guillemot | 4 | Y09124 | B3485 | 16-Apr-18 | 14:57 | 05-Jul-18 | 10:30 | Brooding chick | NA | NA |
| 15 | Guillemot | 4 | R01694 | B3486 | 16-Apr-18 | 14:57 | 05-Jul-18 | 10:30 | Brooding chick | 2019 | Yes |
| 16 | Guillemot | 4 | R76319 | B3487 | 16-Apr-18 | 14:58 | 06-Jul-18 | 10:30 | Brooding chick | NA | NA |
| 17 | Guillemot | 4 | Y09127 | B3488 | 16-Apr-18 | 14:58 | 05-Jul-18 | 15:00 | Brooding chick | NA | NA |
| 18 | Guillemot | 3 | R76263 | B3489 | 16-Apr-18 | 14:58 | 01-Jul-18 | 11:00 | Brooding chick | NA | NA |
| 19 | Guillemot | 3 | Y09126 | B3490 | 16-Apr-18 | 14:59 | 06-Jul-18 | 09:00 | Brooding chick | NA | NA |
| 20 | Guillemot | 9 | R02610 | B3996 | 25-Jun-18 | 18:42 | 05-Jul-18 | 09:00 | Brooding chick | 2019 | Yes |
| 21 | Guillemot | 9 | Y09106 | B3997 | 25-Jun-18 | 18:42 | 04-Jul-18 | 15:00 | Brooding chick | NA | NA |
| 22 | Guillemot | 9 | Y09010 | B3998 | 25-Jun-18 | 18:43 | 04-Jul-18 | 15:00 | Brooding chick | 2019 | Yes |
| 23 | Guillemot | 3 | R03179 | B3999 | 25-Jun-18 | 18:43 | 04-Jul-18 | 15:00 | Brooding chick | 2019 | Yes |
| 24 | Guillemot | 9 | Y09104 | B4000 | 25-Jun-18 | 18:44 | 04-Jul-18 | 15:00 | Brooding chick | NA | NA |
| 25 | Guillemot | 9 | Y09107 | B4001 | 25-Jun-18 | 18:44 | 04-Jul-18 | 15:00 | Brooding chick | 2019 | Yes |
| 26 | Guillemot | 9 | Y09118 | B4002 | 25-Jun-18 | 18:45 | 05-Jul-18 | 10:30 | Brooding chick | 2019 | Yes |
| 27 | Guillemot | 9 | Y09109 | B4003 | 25-Jun-18 | 18:45 | 05-Jul-18 | 09:00 | Brooding chick | NA | NA |
| 28 | Guillemot | 9 | R76657 | B4004 | 25-Jun-18 | 18:48 | 05-Jul-18 | 09:00 | Brooding chick | 2019 | Yes |
| 29 | Guillemot | 9 | Y09121 | B4005 | 25-Jun-18 | 18:48 | 05-Jul-18 | 10:30 | Brooding chick | NA | NA |
| 30 | Guillemot | 9 | R23421 | B4006 | 25-Jun-18 | 18:49 | 05-Jul-18 | 09:00 | Brooding chick | 2019 | Yes |
| 31 | Guillemot | 9 | Y09108 | B4007 | 25-Jun-18 | 18:49 | 05-Jul-18 | 09:00 | Brooding chick | NA | NA |
| 32 | Guillemot | 9 | Y09119 | B4010 | 26-Jun-18 | 13:28 | 05-Jul-18 | 10:30 | Brooding chick | 2019 | Yes |
| 33 | Guillemot | 4 | Y09125 | B4011 | 26-Jun-18 | 13:30 | 05-Jul-18 | 10:30 | Brooding chick | 2019 | Yes |
| 34 | Guillemot | 9 | Y09120 | B4013 | 26-Jun-18 | 13:31 | 05-Jul-18 | 10:30 | Brooding chick | NA | NA |
| 35 | Razorbill | 2 | M82440 | BN000 | 13-Jun-18 | 05:24 | 26-Jun-18 | 07:30 | Brooding chick | NA | NA |
| 36 | Razorbill | 2 | M84641 | BN001 | 13-Jun-18 | 05:25 | 26-Jun-18 | 07:50 | Brooding chick | NA | NA |
| 37 | Razorbill | 2 | M82448 | BN002 | 13-Jun-18 | 05:26 | 26-Jun-18 | 08:05 | Brooding chick | 2019 | Yes |
| 38 | Razorbill | 4 | M84674 | BN003 | 13-Jun-18 | 05:27 | 27-Jun-18 | 07:00 | Brooding chick | 2019 | Yes |
| 39 | Razorbill | 2 | M84675 | BN004 | 13-Jun-18 | 05:28 | 27-Jun-18 | 07:20 | Brooding chick | NA | NA |
| 40 | Razorbill | 3 | M84676 | BN005 | 13-Jun-18 | 05:28 | 27-Jun-18 | 07:50 | Brooding chick | 2019 | Yes |
| 41 | Razorbill | 8 | M84677 | BN006 | 13-Jun-18 | 05:29 | 27-Jun-18 | 08:10 | Brooding chick | NA | NA |
| 42 | Razorbill | 1 | M84652 | BN007 | 13-Jun-18 | 05:30 | 27-Jun-18 | 16:30 | Brooding chick | NA | NA |
| 43 | Razorbill | 1 | M82449 | BN008 | 13-Jun-18 | 05:31 | 27-Jun-18 | 17:00 | Brooding chick | 2019 | Yes |
| 44 | Razorbill | 3 | M84643 | BN009 | 13-Jun-18 | 05:31 | 27-Jun-18 | 17:20 | Brooding chick | 2019 | Yes |
| 45 | Razorbill | 10 | M84678 | BN010 | 13-Jun-18 | 05:32 | 28-Jun-18 | 07:40 | Brooding chick | NA | NA |
| 46 | Razorbill | 2 | M84679 | BN011 | 13-Jun-18 | 05:33 | 28-Jun-18 | 08:10 | Brooding chick | NA | NA |
| 47 | Razorbill | 3 | M84680 | BN012 | 13-Jun-18 | 05:33 | 28-Jun-18 | 08:30 | Brooding chick | NA | NA |
| 48 | Razorbill | 6 | M84644 | BN013 | 13-Jun-18 | 05:34 | 28-Jun-18 | 09:00 | Brooding chick | NA | NA |
| 49 | Razorbill | 5 | M84636 | BN014 | 13-Jun-18 | 05:35 | 28-Jun-18 | 09:20 | Brooding chick | 2019 | Yes |
| 50 | Razorbill | 6 | M84681 | BN015 | 13-Jun-18 | 05:36 | 28-Jun-18 | 12:00 | Brooding chick | 2019 | Yes |
| 51 | Razorbill | 4 | M84682 | BN016 | 13-Jun-18 | 05:37 | 28-Jun-18 | 17:00 | Brooding chick | NA | NA |
| 52 | Razorbill | 2 | M84683 | BN017 | 13-Jun-18 | 05:37 | 28-Jun-18 | 20:20 | Brooding chick | NA | NA |
| 53 | Razorbill | 10 | M84684 | BN018 | 13-Jun-18 | 05:38 | 28-Jun-18 | 20:55 | Brooding chick | 2019 | Yes |
| 54 | Razorbill | 8 | M84685 | BN019 | 13-Jun-18 | 05:39 | 29-Jun-18 | 09:00 | Brooding chick | NA | NA |
| 55 | Razorbill | 7 | M84647 | BN020 | 13-Jun-18 | 05:40 | 29-Jun-18 | 09:40 | Brooding chick | 2019 | Yes |
| 56 | Razorbill | 1 | M84654 | BN021 | 13-Jun-18 | 05:40 | 29-Jun-18 | 09:50 | Incubating egg | NA | NA |
| 57 | Razorbill | 4 | M84686 | BN022 | 13-Jun-18 | 05:41 | 29-Jun-18 | 10:01 | Unknown | NA | NA |
| 58 | Razorbill | 10 | M51471 | BN023 | 13-Jun-18 | 05:42 | 29-Jun-18 | 10:15 | Brooding chick | NA | NA |
| 59 | Razorbill | 6 | M84687 | BN024 | 13-Jun-18 | 05:45 | 30-Jun-18 | 08:40 | Brooding chick | 2019 | Yes |
| 60 | Razorbill | 11 | M84688 | BN025 | 13-Jun-18 | 05:46 | 30-Jun-18 | 09:00 | Brooding chick | NA | NA |
| 61 | Razorbill | 10 | M84689 | BN026 | 13-Jun-18 | 05:47 | 30-Jun-18 | 11:15 | Brooding chick | 2019 | Yes |
| 62 | Razorbill | 1 | M84633 | BN027 | 13-Jun-18 | 05:48 | 30-Jun-18 | 11:13 | Incubating egg | NA | NA |
| 63 | Razorbill | 3 | M84690 | BN028 | 13-Jun-18 | 05:49 | 30-Jun-18 | 12:14 | Brooding chick | NA | NA |
| 64 | Razorbill | 3 | M84691 | BN029 | 13-Jun-18 | 05:49 | 30-Jun-18 | 14:10 | Brooding chick | NA | NA |



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