



Concise review/Le point sur

The different breeding strategies of penguins: A review

*Revue des différentes stratégies de reproduction des manchots*André Ancel^{a,b,*}, Michaël Beaulieu^c, Caroline Gilbert^d^a Université de Strasbourg, IPHC, 23, rue Becquerel, 67087 Strasbourg, France^b CNRS, UMR 7178, 67037 Strasbourg, France^c University Freiburg, Biology I, Hauptstrasse 1, 79104 Freiburg, Germany^d Université Paris-Est, École nationale vétérinaire d'Alfort, UMR 7179 CNRS MNHN, 7, avenue du Général-de-Gaulle, 94704 Maisons-Alfort, France

ARTICLE INFO

Article history:

Received 10 December 2012

Accepted after revision 12 February 2013

Available online 11 March 2013

Keywords:

Southern hemisphere
Breeding range
Mate fidelity
Egg formation
Reproduction
Vulnerability

Mots clés :

Hémisphère Sud
Aire de reproduction
Fidélité
Formation de l'œuf
Reproduction
Vulnérabilité

ABSTRACT

The 18 penguin species are exclusively and widely distributed in the Southern hemisphere, from the Equator to the Antarctic continent, and are thus submitted to various ecological constraints in their reproductive strategy. This results in a high variability in all aspects of the breeding biology of the different species. Although penguins appear primarily adapted for a marine existence, they remain dependent on land for breeding, rearing young, and moulting. Here we describe and compare the breeding cycle of all the penguin species, highlighting the characteristics of each species in terms of breeding range, population status, threats induced by environmental changes, duration of the different phases of the breeding cycle, mate fidelity, body mass, body height, egg mass and duration of egg formation. We also focus on the breeding cycle of the genus *Aptenodytes*, since it largely differs from the breeding cycle of most of the other penguin species.

© 2013 Académie des sciences. Published by Elsevier Masson SAS. All rights reserved.

R É S U M É

Les 18 espèces de manchots vivent exclusivement dans l'hémisphère Sud, de l'Équateur jusqu'en Antarctique et subissent en conséquence différentes contraintes écologiques au cours de leur cycle reproducteur. Cela se traduit par une forte variabilité dans tous les aspects de la biologie de la reproduction chez les différentes espèces. Bien que les manchots soient de prime abord adaptés à une vie marine, ils dépendent néanmoins d'une surface ferme pour se reproduire, élever leur progéniture et muer. Nous décrivons et comparons les cycles reproducteurs de toutes les espèces de manchots, tout en mettant en avant les caractéristiques de chaque espèce, que ce soit la durée de leur cycle reproducteur, le statut de la population et les menaces environnementales qui pèsent sur elle, la durée des différentes phases du cycle reproducteur, la fidélité du couple, la masse corporelle, la taille, la durée et les dimensions des œufs. Nous nous sommes plus particulièrement étendus sur le cycle reproducteur du genre *Aptenodytes*, qui diffère notablement de celui des autres espèces.

© 2013 Académie des sciences. Publié par Elsevier Masson SAS. Tous droits réservés.

* Corresponding author. IPHC-DEPE, UMR 7178 CNRS-Uds, 23, rue Becquerel, 67087 Strasbourg, France.

E-mail address: andre.ancel@iphc.cnrs.fr (A. Ancel).

1. Introduction

Natural selection favours a breeding strategy which, in a given environment, is the most likely to ensure the production of the largest number of young that survive to breed, and the survival of parents until they breed again. The decision to breed is triggered by physiological changes set in motion by changes in the environment [1]. In birds, the timing of reproduction is determined by proximate factors and is often initiated before birds return to their breeding grounds [1,2]. Among these proximate factors, the photoperiod, the temperature, and the availability and the quality of the food required for successful breeding are frequently reported [3–6].

The majority of birds lives in a non-uniform environment (spatially and temporarily) and must breed when the conditions are the most favourable. For instance, in temperate and Polar regions, most birds have to synchronize their breeding schedule with the time of the year that will give their offspring the best chances of survival [7]. This restricted period of time dictated by resource availability leads individuals from the same population to breed synchronously. This breeding synchrony can also be associated with different advantages such as the reduction of the impact of predators and the maximisation of chick survival, by timing fledging when prey abundance is highest [8,9], and/or when climatic conditions are most favourable [10].

Among seabirds, penguins are a distinctive group of flightless, long-lived pelagic seabirds. All the 18 species of penguins are grouped within a single family, the *Spheniscidae*. Within this family, there are six genera and each genus comprises one to eight species (Table 1). Several subspecies of little *Eudyptula minor* (six) and gentoo penguins *Pygoscelis papua* (two) have been identified [11,12] but for the purpose of this review, these penguins are considered as a single species. Some penguin species have been studied in detail, especially the Adélie penguin *Pygoscelis adeliae* (more than 400 papers, see ISI Web of Knowledge), but others, such as the Snares penguin *Eudyptes robustus* and erect-crested penguin *Eudyptes sclateri* have not (less than ten publications, see ISI Web of Knowledge). Consequently, there is less information available for some species. All penguins have a similar body shape and structure. The sexes are outwardly similar in all species, though males are usually heavier, at least at the onset of the breeding season, and larger than females (Table 1). They vary considerably in height, from about 40 to 130 cm, and in body mass, from 1 to 37 kg (little and emperor *Aptenodytes forsteri* penguins, respectively, Table 1). Such a range in body mass is only surpassed by another flightless family: the Struthioniformes, where the smallest species, the Little spotted kiwi *Apteryx owenii*, measures 40 cm and weighs 1.2 kg compared to the largest living ratite, the African ostrich *Struthio camelus*, that stands roughly 3 m tall and weighs 160 kg.

Penguins are highly adapted for marine life and some species spend as little as 20% of the year on land (Fig. 1A). Nevertheless, this relatively short period on land represents one of the most important parts of their life cycle; during which penguins have to obtain a nest-site and mate,

lay egg(s), rear chick(s), and moult. The 18 penguins' species share some breeding characteristics such as breeding synchrony (except for the Galápagos penguins *Spheniscus mendiculus*, [72]) for annual reproduction on land, and alternation of sojourns on land and at sea between partners during the chick-rearing period. An important characteristic of all penguin species is their ability to withstand prolonged period of fasting, on land or sea-ice, during breeding. With the exception of the yellow-eyed penguin *Megadyptes antipodes*, all penguin species breed colonially [73]. In these colonies, birds often breed in high densities, for instance, 1.4 nest.m⁻² in Adélie penguins [74] and up to 10 birds.m⁻² in huddling emperor penguins [16,75], which highly increases the level of social stimulations and interactions [76]. High densities of breeding birds have led to the evolution of a varied and complex repertoire of visual and vocal displays (e.g. [77]). As penguins are widely distributed in the Southern hemisphere, from the Equator to the Antarctic continent, they are submitted to various ecological constraints that will impact the timing of reproduction. For example, for a high latitude species such as the emperor penguin, two proximate factors controlling the onset of breeding are exogenous factors such as day-length [1,78] and the peak of ocean productivity preceding breeding [79]. For the equatorial Galápagos penguin, the onset of breeding is closely linked to mean sea surface temperature [70]. Therefore, because of the variety of environments where penguins live, the examination of the breeding cycle observed across the Sphenisciformes order can give some insights about how environmental conditions can shape a variety of reproductive strategies. The different aspects of the biology of penguins are relatively sparse in the literature although some books attempted to gather these data (e.g., [11,12,80,81]). Therefore, the aim of the present study is to provide researchers working on seabirds with a concise and clear overview of the breeding cycle of the 18 penguin species.

In this review, the breeding cycle of penguins is divided into three stages:

- the pairing period, when breeders come ashore for courtship and mating;
- the incubation period, when mates generally take turns to incubate the egg(s);
- the rearing period, from chick(s) guard to fledging.

We defined the courtship period as the shortest period of time between the early arrival of both sexes on the reproductive site and the laying of the first egg because females may occasionally go to sea before the full clutch is completed (e.g. gentoo penguins, [11], magellanic penguins *Spheniscus magellanicus* [82]). The body mass of the adult birds considered was the mean body mass at the onset of the breeding period, i.e. at their arrival on the breeding site.

2. Geographical range and population status

Penguins are widely distributed in the Southern hemisphere, mainly between 45 and 60°S (Fig. 2) and

Table 1
Some biological characteristics of the 18 penguin species.

Scientific name (abbreviation) Common name	Breeding range (°S)	IUCN 2012 status	Height (cm)	Mean body mass (kg)		Mate fidelity (%)	Mean egg mass (g)		Egg formation (d)	
				Male	Female		A-egg	B-egg	A-egg	B-egg
<i>Aptenodytes forsteri</i> (Af) Emperor penguin	66–78 [11,12]	NT	100–130 [11,12]	37.3 [15,16]	28.8 [15,16]	15 [31]	445 [16,36–38]		21	
<i>Aptenodytes patagonicus</i> (Ap) King penguin	45–55 [11,12]	LC	85–95 [11,12]	14.5 [17,18]	12.9 [17,18]	29 [17]	307 [11,17,39]		20	
<i>Eudyptes chrysocome</i> (Ec) Northern rockhopper penguin	37–53 [12]	VU	45–58 [11,12]	3.4 [19,20]	3.1 [19,20]	59 [12]	84 [19,40–43]	113 [19,40–43]	16	20
<i>Eudyptes chrysolophus</i> (Ech) Macaroni penguin	46–65 [12]	VU	71 [11,12]	5.0 [12]	5.2 [12]	75 [32,33]	93 [41,44,45]	149 [41,44,45]	16	21
<i>Eudyptes moseleyi</i> (Em) Southern rockhopper penguin	37–40 [13]	EN	45–58 [11,12]	3.4 [19,20]	3.1 [19,20]	59 [12]	84 [19,40–43]	113 [19,40–43]	16	20
<i>Eudyptes pachyrhynchus</i> (Ep) Crested or fiordland penguin	43–48 [11]	VU	55 [11,12]	4.3 [21,22]	3.8 [21,22]	Long [11]	100 [22,46]	118 [22,46]	16	21
<i>Eudyptes robustus</i> (Er) Snares penguin	48 [11,12]	VU	51–61 [11,12]	2.6 [23]	2.5 [23]	Long [23]	90 [23]	117 [23]	16	21
<i>Eudyptes schlegeli</i> (Es) Royal penguin	54 [11]	VU	65–75 [11,12]	5.7 [24]	5.3 [24]	Long [11,12]	100 [11]	159 [11]	16	21
<i>Eudyptes sclateri</i> (Esc) Erect-crested penguin	47–49 [12]	EN	67 [11,12]	6.4 [25]	5.4 [25]		85 [47–49]	151 [47–49]	16	21
<i>Eudyptula minor</i> (Em) Little blue penguin	32–47 [11]	LC	40–45 [11,12]	1.2 [11,26]	1.0 [11,26]	82 [11,12]	54 [11]	53 [11]	15	19
<i>Megadyptes antipodes</i> (Ma) Yellow-eyed penguin	43–52 [11]	EN	66–78 [11,12]	5.3 [27]	4.9 [27]	33 [34]	138 [34]	137 [34]	17	21
<i>Pygoscelis adeliae</i> (Pa) Adélie penguin	54–77 [12]	NT	70 [11,12]	5.6 [28,29]	4.9 [28,29]	80 [11,29,35]	122 [50–52]	114 [50–52]	17	21
<i>Pygoscelis antarctica</i> (Pan) Chinstrap penguin	54–64 [12]	LC	71–76 [11,12]	5.0 [11,29]	4.7 [11,29]	83 [11,12,29]	114 [53]	112 [53]	16	20
<i>Pygoscelis papua</i> (Pp) Gentoo penguin	46–65 [11,12]	NT	75–90 [11,12]	5.6 [12]	5.2 [12]	85 [11,12,29,33]	129 [43,44,54,55]	129 [43,44,54,55]	17	21
<i>Spheniscus demersus</i> (Sd) Jackass or African penguin	24–34 [14]	EN	70 [12]	3.1 [12]	2.8 [12]	62 [12]	107 [56]	105 [56]	16	20
<i>Spheniscus humboldti</i> (Sh) Humboldt penguin	5–42 [12]	VU	65 [12]	5.0 [12]	4.2 [12]		122 [12]	122 [12]	17	21
<i>Spheniscus magellanicus</i> (Sm) Magellanic penguin	29–54 [12]	NT	70 [11,12]	4.9 [30]	4.6 [30]		125 [30,57]	125 [30,57]	17	21
<i>Spheniscus mendiculus</i> (Sme) Galápagos penguin	0–2 [12]	EN	53 [12]	2.1 [12]	1.9 [12]	89 [12]				

Body mass is the mean body mass of birds at the onset of the breeding season. Mate fidelity is from one year to the consecutive year. For egg formation values, see the Material and Methods section. The B-egg formation column represents the time elapsed since the onset of the first yolk development. Abbreviations of the different species are given in brackets after the scientific name of each species. Each reference is indicated by a number in brackets. IUCN 2012 red list codes: EN: endangered; LC: least concern; NT: near threatened; VU: vulnerable.

four penguin species are endemic to their breeding island: Fiordland (or crested) *Eudyptes pachyrhynchus*, Galápagos, royal *Eudyptes schlegeli* and Snares penguins. Penguins represent roughly 90% of bird biomass in the Southern Ocean [12] and *Pygoscelis* species (chinstrap *P. antarctica*, Adélie and gentoo penguins) represent 70% of Antarctic avian biomass [83]. Penguins feed in pelagic cold waters, rich in zooplankton and biomass, where they consume approximately two millions tons of carbon per year [84]. However, they occupy a wide variety of habitats while breeding on land, ranging from the burrows in the volcanic

rocks of the hot Galápagos Islands (Galápagos penguin), the bushes of south Australia and New-Zealand (little penguin), to the ice of the border of the Antarctic continent (emperor, Adélie, chinstrap and gentoo penguins). This results in a high variability in all aspects of the breeding biology of the different penguins' species (Fig. 1A). While most species breed annually, some species, such as African (or jackass) and Galápagos penguins, can have no distinct annual breeding season (*i.e.* breeding season occurs year-round). Because of the short Antarctic summer, breeding is generally much synchronised in Antarctic species, but in

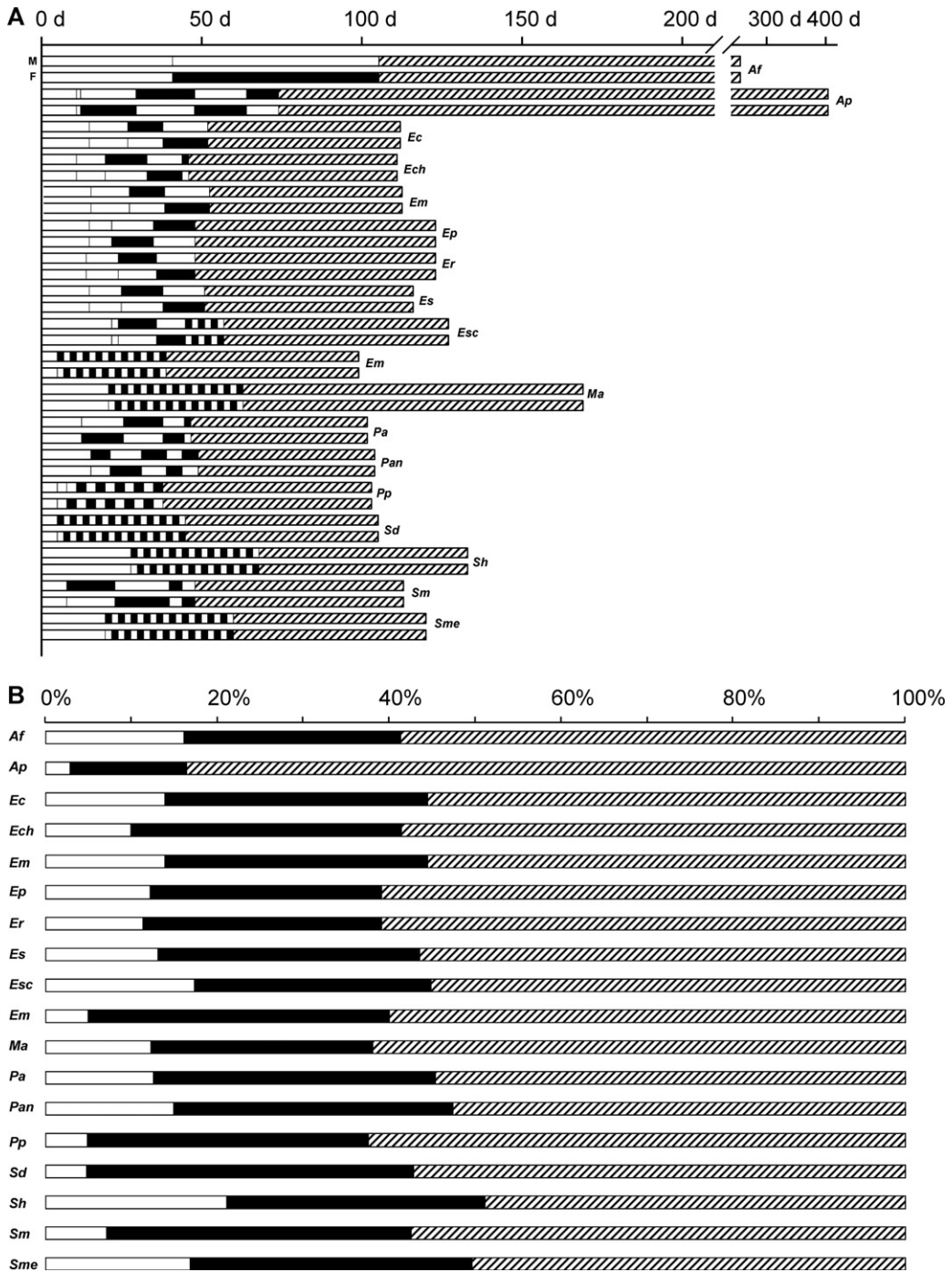


Fig. 1. Breeding cycle of the penguins. White, dark and dashed bars denote for fasting on land, foraging at sea and feeding, chick rearing, respectively. Lettering stands for species (Table 1). A. Breeding cycle of the 18 penguin species and for both sexes (M stands for males and F for females). B. Relative breeding cycle of the 18 penguin species. Mean courtship period is defined as the shortest time between pair formation and the laying of the first egg. Breeding cycle is defined as the time elapsed from the arrival of the penguins on their breeding site to the fledging of the chick(s). References for each species and periods.

Species	Pairing	Incubation	Chick rearing
<i>Af</i>	[31,58]	[16,31,35,58]	[31,35,58]
<i>Ap</i>	[17,18,59]	[17,18,35,59,63]	[17,18]
<i>Ec</i>	[11,12,20,40]	[12,20]	[12,40]
<i>Ech</i>	[32,44,60]	[32,44,60]	[32]
<i>Em</i>	[11,12,20,40]	[12,20]	[12,40]

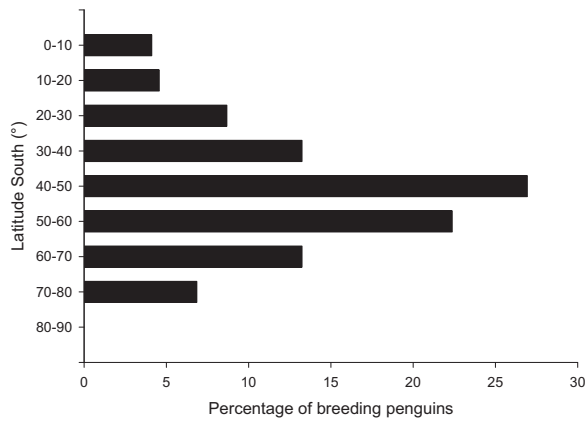


Fig. 2. Repartition of the 18 species of breeding penguins (%) according to the Southern latitudinal gradient.

temperate climates, it can be spread over a longer period of time. For example, the timing of egg-laying greatly varies between years for the little penguin [85].

Penguins exclusively rely on marine food sources that are spatially and temporarily unpredictable. Penguins make use of wide geographical areas in the ocean while foraging and during migrations. Penguins, which are central place foragers during the breeding season, are thus particularly sensitive to variations in ecosystem structure and processes. Of the 18 penguins' species, 13 are considered endangered or threatened (Table 1) and some species are now at their lowest recorded populations: Galápagos, yellow-eyed, and Fiordland have less than 3,000 pairs; Humboldt (*Spheniscus humboldti*), Snares and African have less than 30,000 pairs. Even abundant species like the macaroni (*Eudyptes chrysolophus*) and the rock-hopper (*Eudyptes chrysolophus* and *E. moseleyi*) are in steep decline [86]. On the contrary, the global population trend is stable for Snares and Adélie penguins [86] or is increasing for gentoo [87,88] and king penguins [86]. The status of the emperor penguin might change in the near future. Indeed, thanks to satellite images showing faecal stains on ice, Fretwell and Trathan [89] discovered ten new emperor penguins colonies [90]. Around 80% of the threatened species live on islands, increasing their vulnerability to threats such as introduced predators. Many penguin species face the same four key threats: global climate

change, marine pollution, fisheries mismanagement, and introduced mammalian predators [91].

3. Relative importance of each phase of the breeding cycle: focus on pairing period

Among the 18 penguin species, the duration of the breeding season ranges from 4 to 15 months (Fig. 1A). It is especially long for the two largest penguins (who need more time for their chick to grow) and particularly for the king penguin *Aptenodytes patagonicus*. Emperor penguins' cycle lasts 9 months including a 1.5-month pairing period that is long relative to other penguin species (Fig. 1A; [79]). Indeed, while the courtship period ranges from less than 1 week in little, gentoo, Jackass or African *Spheniscus demersus* and magellanic penguins, it lasts up to 6 weeks in emperor penguins (Fig. 1A). The ratio courtship-breeding duration accounts for 16% in emperor penguin (1.5 month vs. 9 months) whereas it accounts for only 3% in its closest relative, the king penguin (1.5 week vs. 15 months). Considering the long duration of the total breeding cycle in emperor penguins, this ratio is, however, almost of the same magnitude (14–18%) as with other species breeding in Antarctica (Adélie, chinstrap and gentoo penguins; Fig. 1B). Contrary to the emperor penguin, the king penguin does not exhibit a long courtship period but an extended chick-rearing period (84% of the breeding cycle while it represents about 57% in other species; Fig. 1B). In king penguins, breeding overlaps two years (Fig. 1A) which consequently results in a maximum of two chicks being reared within a three year period.

4. Fidelity, sex ratio, and courtship

All penguins are monogamous, mating with only a single partner each year. In addition, most penguin species are territorial and show a moderate to high inter-annual fidelity in breeding (59–89%; Table 1), partners reuniting from year to year on their nest site. Pair bonds can therefore be long-lasting (Table 1) with many birds returning to meet their previous partners at the same breeding site each year. Two *Aptenodytes* species and especially the emperor penguin, however, show a low inter-annual fidelity (15%; Table 1), due to the fact that they do not use a nest, and incubate their egg on their feet

(Continued)

Species	Pairing	Incubation	Chick rearing
<i>Ep</i>	[12,22]	[12,22]	[12]
<i>Er</i>	[11,12,23]	[11,12,23]	[12]
<i>Es</i>	[11,12,24,61]	[24,44,61]	[12]
<i>Esc</i>	[23,42]	[11,12,48]	[12,25]
<i>Em</i>	[62]	[11,62,64,65]	[71]
<i>Ma</i>	[11]	[27,34]	[12]
<i>Pa</i>	[12,29,52,53]	[11,12,35,50,52,66,67]	[12,35]
<i>Pan</i>	[12,29,50,53]	[12,50,53]	[12,53]
<i>Pp</i>	[29]	[11,12,54,68,69]	[69]
<i>Sd</i>	[56]	[14,56]	[56]
<i>Sh</i>	[12]	[12]	[12]
<i>Sm</i>	[30]	[30]	[12]
<i>Sme</i>	[12]	[70]	[12]

[92,93]. Partners cannot therefore reunite themselves from year to year on the nest site and one could hypothesize that these penguins need more time than other species to find and reunite with their previous mate. King penguins, however, show a low inter-annual fidelity (29%; Table 1) despite being highly territorial and occupying distinct nest sites. These birds could easily reunite with their partners provided they return on time but it does not seem to be the case for 70% of them.

A male-biased sex ratio appears to be a characteristic of breeding populations of several penguin species such as yellow-eyed, gentoo, little, Adélie, macaroni penguins [11]. In Adélie and macaroni penguins, males return first [11] to previous year's nest-site to maximize their chances of reuniting with their mates from the previous year. This suggests that the primary "aim" of male penguins returning to the breeding colony would be to retain their old nest-sites and, only secondarily, to reunite with their previous mates. Because in many species there are more males in the population than females, male-male competition for nest-sites, rather than for mates, is an important determinant of breeding opportunity. In contrast, at least for the Pointe Géologie colony (Adélie Land, 140°E-67°S), sex ratio favours female emperor penguins that outnumber males by about 10% [31,92–94]. This might be due to a higher mortality of male breeders after their long winter fast [95]. Because of this unbalanced sex ratio and because mate fidelity is particularly low in this species, competition between females is high to find a male and the earlier a female returns at the onset of the breeding cycle, the higher the probability she will get a mate [31], and the lower the probability that her previous mate will already be paired with another female. Furthermore, the number of male partners available per unpaired female decreases as time passes. Consequently, an early arrival enhances the likelihood for the females to preserve their breeding status by finding an unpaired mate. Unbalanced sex ratio in emperor penguins probably also explains occurrence of polygynous trios (one male with two females) but these groups are temporary, one female usually ejecting the other after a few hours [16,31].

5. From yolk development to fledging

The period of yolk development that precedes ovulation is given by the following equation for non-procellariiform seabirds: $\log t = 0.396 + 0.283 \log \text{egg mass}$ [96] where t stands for time (day) and egg mass is expressed in grams. Once the yolk is fully developed, it is retained within the ovarian follicle for about 6 days before ovulation [97]. The albumen and shell are then added over about 24 hours, following ovulation. In species with two eggs [98–101], development of the second egg (B-egg) is initiated 4 days after the first one (A-egg). Thus, in order to calculate the time elapsed from yolk formation until egg-laying, we added 7 (6+1) and 11 days (4+6+1) for the A-egg and the B-egg, respectively (Table 1).

Incubation refers to the process by which birds lay their egg(s), and to the development of the embryo within the egg. The most vital factors of incubation are the temperature required for embryo development over a specific

period, humidity (which could be problematic in dry environments like the Galápagos Islands or Antarctica), and egg rotation rate [102]. If the air is too dry, the egg will lose too much water, which can make hatching difficult or impossible. As incubation progresses, an avian egg becomes lighter, and the air space within the egg becomes larger owing to the evaporation from the egg. In most penguin species, incubation is divided differently between parents with the male and the female taking turns incubating the egg(s). The emperor penguin represents an exception as in this species, only the male incubates.

Compared with other birds, penguins lay small eggs (and small egg clutches) relative to their body weight ($2.9 \pm 0.9\%$, $n = 16$, mean \pm SD, Table 1), and eggs of smaller penguin species are proportionately larger (Table 1). The majority of penguins lays two white eggs (except genus *Aptenodytes* laying only one egg), weighing from 55 to 445 g (little and emperor penguins, respectively, Table 1). Within the genus *Eudyptes*, the second egg or B-egg is 20 to 78% larger than the first one (Table 1) and consequently the first chick is smaller than the second one and generally fails to survive [11]. Erect-crested penguins are obligate brood reducers [46] and exhibit the most extreme egg-size dimorphism of any bird: the second egg is up to 100% the size of the first and is the only one to be incubated. It remains unclear why these birds should lay two eggs but only ever rear one chick. The total time necessary for yolk deposition is proportional to female's body mass [97] but full clutch synthesis has the same duration among the 18 penguin species, ca. 3 weeks (Table 1; 20.6 ± 0.7 days, $n = 16$, mean \pm SD). By subtracting the duration of the full clutch completion from the duration of the pairing period, we can report that egg formation begins while females are still at sea (in 13 species), while they arrive at the breeding site (in one species), and while they already are on the breeding site (in three species; Fig. 3). Among the latter, emperor penguins' egg formation begins late, three weeks after their arrival on the colony. Thus, egg formation is not responsible for the long duration of the pairing period, representing only one half of the courtship duration.

Incubation ranges from ca. 35 to 65 days (*Eudyptes* and *Pygoscelis* species, emperor penguins, respectively, Fig. 1B). It is an energetically demanding process, especially in male emperor penguins that can lose up to 45% of their initial

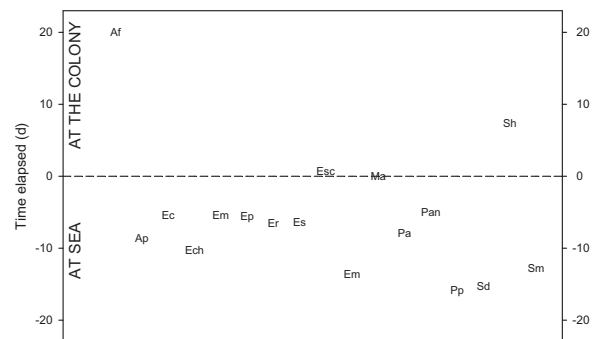


Fig. 3. Time elapsed from arrival at the breeding site (dashed line) and onset of yolk formation for the 17 penguin species among the 18 ones. Lettering stands for penguin species (Table 1).

body weight during this time. Some species begin incubation with the first egg, causing the young to hatch asynchronously; others begin after laying the second egg, thus decreasing hatching asynchrony between siblings [103].

Both parents are involved in parental care. From the moment the egg is hatched, one parent cares for the newly hatched chick(s) while the other forages for food. Penguin chicks are semi-altricial, *i.e.* they need parental care (food, warmth, protection) before becoming independent. At the start of the rearing period, the chicks either sit on their parents feet (emperor and king penguins) or under their bellies, to be kept warm and dry. Young are guarded (guarding stage) by both parents for varying periods of time before forming crèches. Both parents feed the chicks by regurgitation. Nestlings beg for food by pecking adult's bill and/or by singing. Guarding stage may be affected by environmental conditions [102] and when they are not being protected by the adult, the chicks form crèches to keep warm and stay protected. As the chicks grow, their feeding requirements quickly increase, making it difficult for just one of the parents to obtain enough food. Eventually, the chicks are large enough so that both parents can go to sea to gather food for their chick simultaneously. When parental provisioning is low, alloparental feeding (feeding of offspring by adults other than their own parents) sometimes occurred in little [104], emperor [105], king [106] and Adélie penguins [74]. In most species, the chicks gather together in crèches to provide protection both from predators and from the elements. In some species, such as king penguins, crèches can be large with many hundreds of chicks tightly packed together. In other species, such as African penguins, the crèches are smaller (with up to ten chicks coming together) and less dense. Age at fledging ranges from one to eleven months (Fig. 1A) and fledglings are a bit smaller and lighter than adults, except in emperor penguins for which body mass of fledglings is about half of adults' [16]. While they are still on their breeding grounds, chicks have to moult to adorn themselves with a waterproof plumage, which will allow them in turn to go hunting offshore and acquire their food independently. Adults also have to moult like other birds and all penguins replace their feathers each year on their breeding grounds, except for the emperor penguins for which moulting often takes place far away from their breeding grounds [107–109]. Although other birds lose some feathers individually at a time and replace them over a few months, penguin moulting is fast, taking only two (*e.g.*, *Pygoscelis antarctica*, [110]) to five weeks (*e.g.*, genus *Aptenodytes*, [16,111,112]). Moulting is extremely important to penguins as they need to maintain their feathers in perfect condition at all times to insulate their body from environmental conditions at sea or on land. Adults generally moult after the breeding season. Once their chicks have moulted into their own juvenile plumage the adults return to sea for a few weeks to build up their own fat reserves and then come back ashore to moult. During the moult, penguins are no longer waterproof and cannot enter the sea, they can lose up to 50% of their body mass, they are not well insulated and they are vulnerable to predation. Therefore, this is a critical period of time for

penguins during which they have to face the elements and starve until their new set of feathers is ready.

6. Genus *Aptenodytes*

Birds of the genus *Aptenodytes* (*A. forsteri* and *A. patagonicus*) are bigger and taller than other penguin species (Table 1). They do not build a nest and only lay one large egg (on average 445 and 307 g, respectively; Table 1), which is kept on the top of the incubating parents' feet at least for 54 days (Fig. 1B). As for other penguins, parents recognise their chick by voice and young also recognise parents by call [11].

The breeding cycle of the genus *Aptenodytes* largely differs from the breeding cycle of most of the other penguin species. In most penguin species, it takes from 8 to 15 weeks to raise a chick to the juvenile stage (Fig. 1B) but it can take 10 to 13 months for king penguins to fledge their chick [18]. Because of this long chick-rearing period, king penguins only produce two chicks every three years. As a result, 12-month-old chicks cohabit with incubating birds in king penguin colonies. Every year, emperor penguins manage to raise their large chicks more quickly (five months), using a different strategy. Indeed the chicks moult into juvenile plumage while they are still much smaller than their parents. The juveniles then continue to grow out at sea.

The emperor penguin also seems to be an exception among penguins as they begin their breeding cycle when other Antarctic birds have finished theirs. Each year, from late December to March (*i.e.* late summer), emperor penguins disperse into the oceans, travelling and foraging into the waters surrounding the Antarctic continent [109,113]. In March–April, as winter approaches and fast-ice extent grows, all mature emperor penguins move south towards their colonial breeding areas at the border of the Antarctic continent. The breeding cycle of emperors stands in contrast with that of other penguin species (except for the king penguin) by its long duration and by the fact that it takes place in the midst of the severe Antarctic winter, whereas other penguins breed during the short and milder summer season. Indeed, the emperor penguin is one of the few birds for which gonadal growth is coincident with short days, when birds are still at sea. Gonadal steroids are several fold above basal level at the time of arrival on the breeding area suggesting that environmental cues, especially decreasing daylength, decreasing air temperature, and sea-ice formation, stimulate gonadal development and reproduction [1]. Before breeding, emperor penguins forage far away from their breeding grounds [109,114,115]. In the Southern Ocean, a period of high productivity occurs during summer, from October to April, and is followed by a period of low productivity during winter, from May to September [116–118]. Because emperor penguins breed in winter, they have to anticipate their breeding season by accumulating body reserves during high ocean productivity in the previous summer [79]. To our knowledge, this breeding strategy is unique among animals. Furthermore, emperors breed on sea-ice in a few favourable zones that may be a hundred kilometres distant from the open sea or polynias

where they exclusively feed [119]. As a consequence of the distance from the feeding grounds, and because breeding activity competes with feeding, female and male emperors fast for as long as 1.5 and 4 months, respectively [16,31]. For females, the breeding fast comprises only the courtship period, since they leave their single egg to their mate as soon as it is laid and then go back to sea for building up their reserves. For males, the period of fasting includes the courtship and the whole incubation period. To face the austral winter, emperor penguins have to exploit in an optimal way their limited body fuels in order to succeed in their breeding [16,120,121]. This is possible only thanks to their huddling behaviour, which allows them to decrease energy expenditure [120,121].

7. Which environmental changes might affect the breeding cycle of the different penguin species?

The Earth's climate is undergoing rapid warming, which is driving shifts in the distribution and phenology of many plants and animals [122,123]. Among animals, penguins are adapted to live in extreme environments (Fig. 2), but, because each species is restricted to a limited latitudinal range (Table 1), they can be highly sensitive to climate change [124]. Environmental changes are not uniform across regions, with resource increasing in the subantarctic areas and decreasing in Antarctica [123]. Quantifying changes in breeding phenology is important for understanding how populations respond to these changes, especially those resulting from human activities [123].

7.1. Climatic changes and resource availability

Detecting and predicting how populations respond to environmental variability are crucial challenges in management and conservation research. This is particularly true for populations at high latitudes, many of which demonstrate changes in population dynamics associated with global warming [125]. Some seabird populations of the Southern Ocean have been responding to climate change for the last three decades and demographic models suggest that projected warming will cause dramatic population changes over the next century [114]. In the Antarctic ecosystem, population dynamics of top predators like penguins may yield important information about how the environment is changing [126]. The phenotypic plasticity of penguins may allow them to continue to exploit their transformed ecological niche and maintain their current distributional ranges. For instance, penguins may vary the timing of breeding in response to changes in environmental conditions [127]. However, palaeoecological records suggest that penguins are more likely to respond by dispersal rather than adaptation [124]. Thus shift in species distribution is likely to be one of the major possible adaptations to changing environmental conditions [114]. This is exemplified by the distributional range of chinstrap [124,126,128], gentoo and Adélie penguins [124,129,130] that has shifted southwards around the Antarctic Peninsula.

However, as each species is limited to a specific latitudinal range, a latitudinal shift may be very limited.

Thus, emperor penguins' colonies north of 70°S are projected to decrease or disappear, and limited growth might occur south of 73°S [131]. These population trends are likely to be related to sea ice conditions [132]. For example, at Pointe Géologie (Adélie Land), distance to the fast-ice edge and its extent are major determinants of emperor penguin breeding success [132]. Therefore, the increased frequency of warm events associated with projected decreases in sea ice extent is likely to reduce population viability [133,134].

Other physical factors than sea-ice can also affect penguin populations. For instance, sea surface temperature consistently drives the foraging behaviour of king penguins, and, according to climate models, the projected warming of surface waters could lead to a gradual southward shift of their more profitable foraging zones [114]. Such a shift would negatively affect the king penguin population, unless penguins develop alternative foraging strategies [114] as to modify their timing of breeding [127].

The Antarctic Peninsula is among the fastest-warming areas on the Earth, with 5–6 °C increases in mean winter air temperatures and associated decreases in winter sea-ice cover [135]. These perturbations have affected the ecosystem profoundly [135]. To respond to these climatic changes, varying the timing of reproduction in response to local environmental conditions is a key factor influencing reproductive success [127]. For example, clutch initiation and hatching dates of royal, Adélie and gentoo penguins occur earlier with warmer temperatures [123,127]. However, these behavioural adjustments may not be sufficient to prevent populations from declining. The "sea-ice hypothesis" proposing that ice-obligate species directly decline because of sea-ice reduction, does not appear to be sufficient to explain why populations of both ice-loving and ice-avoiding penguins have declined significantly [135]. Some researchers argue in favour of an alternative, more robust hypothesis, that attributes both increases and decreases in penguin populations to changes in the abundance of their main prey, Antarctic krill [135]. Indeed, decline of chinstrap penguin populations has been suggested as being related to climate change through a reduction in sea-ice extent during winter and a consequent decline in the abundance of krill in summer during the breeding season [126].

Climate changes can also have more subtle consequences on the foraging behaviour of penguins. For instance, mixed water regimes resulting from storms, result in the dispersion of prey items in the water column. This lack of prey stratification has been described as resulting in reduced foraging efficiency and poor breeding success in the little penguin [136]. Mixed water regimes are currently unusual during the breeding period of little penguins, but are expected to become more frequent due to climate change and may therefore represent an important threat for this species [136].

7.2. Tourism

Antarctica now fuels one of the fastest growing tourism markets in the world with over 30,000 visitors annually

travelling to the continent [137]. Increasing ecotourism activity has led to concerns about the effects of ecotourism on wildlife populations. While some species of penguins habituate to human visits, others exhibit negative effects due to disturbance [138]. Behavioural, physiological, and reproductive parameters might thereby be affected. For example, human presence at the nest site is physiologically stressful for breeding Magellanic penguins that are not accustomed to seeing humans [139]. Indeed, Magellanic penguins in visited areas have higher corticosterone stress responses than penguins in undisturbed areas [138,140]. Moreover, birds exposed to moderate levels of disturbance do not show evidence of habituation over a period of a few years [139]. However, penguins may habituate to humans, as birds that have been exposed to very high levels of human visitation do not respond anymore to human presence as a stressor. Furthermore, Magellanic chicks from tourist-visited colonies do not flee anymore when approached by humans [140], and breeding success is not affected by visitation levels in this species [138]. However, penguin species differ in their sensitivity to human presence. For instance, in contrast to Magellanic and Adélie penguins, yellow-eyed and gentoo penguins show significantly lower breeding success at sites exposed to unregulated tourism compared to areas visited infrequently [130,141]. This may be attributed to the presence of people on beaches that delays post-foraging landing by penguins provisioning their chicks, which may in turn affect the amount of food delivered to the young. Indeed, yellow-eyed chicks from nesting areas with high numbers of tourists have significantly lower fledging weights than chicks from areas with no tourist visitors [142]. Taking into account that the probability of survival is positively associated with mass at fledging, lower fledging weights may have long-term population consequences [142].

8. Conclusion

The present article shows that breeding strategies are diverse and differ between penguin species. However, breeding behaviour can also exhibit some plasticity within each penguin species and particularly when environmental conditions vary (e.g. [102]). More studies simultaneously conducted (1) in several penguin species breeding in the same location and (2) on the same species in different locations/environmental conditions would be useful to highlight how environmental conditions influence breeding strategies in penguins and how penguins can adapt to environmental perturbations.

Many penguin species face the same threats [91]. Marine and coastal ecosystems are undergoing unprecedented alterations in their processes and structure. Penguins are sensitive species impacted by these phenomena. As top predators, they are key constituents of marine ecosystems, and are indicators of the oceanic and coastal ecosystem health. Larger scale ecosystem-based conservation planning and more focused local efforts are needed for the successful conservation of many penguin species.

Disclosure of interest

The authors declare that they have no conflicts of interest concerning this article.

Acknowledgements

We thank Drs C.-A. Bost, A. Chiaradia, P. Dee Boersma, C. Hull, M. Massaro, K. Pütz, Y. Ropert-Coudert, A. Steinfurth and P. Trathan for providing data on the different species of penguins. We also thank Dr. S. Gallon for her help in revising the language. The manuscript benefited from critical comments by anonymous reviewers.

References

- [1] R. Groscolas, M. Jallageas, A. Goldsmith, I. Assenmacher, The endocrine control of reproduction and moult in male and female Emperor (*Aptenodytes forsteri*) and Adélie (*Pygoscelis adeliae*) penguins. I. Annual changes in plasma levels of gonadal steroids and luteinizing hormone, *Gen. Comp. Endocr.* 62 (1986) 43–53.
- [2] R.H. Drent, The timing of birds' breeding seasons: the Perrins hypothesis revisited especially for migrants, *Ardea* 94 (2006) 305–322.
- [3] E.L. Charnov, J.R. Krebs, On clutch size and fitness, *Ibis* 116 (1974) 217–219.
- [4] D. Lack, *Population studies of birds*, Oxford University Press, Oxford, 1966.
- [5] M.M. Lambrechts, J. Blondel, M. Maistre, P. Perret, A single response mechanism is responsible for evolutionary adaptive variation in a bird's laying date, *Proc. Natl Acad. Sci. USA* 94 (1997) 5153–5155.
- [6] P.J. Sharp, Strategies in avian breeding cycles, *Anim. Reprod. Sci.* 42 (1996) 505–513.
- [7] P. Monaghan, J.D. Uttley, M.D. Burns, Effect of changes in food availability on reproductive effort in Arctic Terns *Sterna paradisaea*, *Ardea* 80 (1992) 71–81.
- [8] K. Camphuysen, Post-fledging dispersal of Common Guillemots *Uria aalge* guarding chicks in the North Sea: the effect of predator presence and prey availability at sea, *Ardea* 90 (2002) 103–119.
- [9] G.M. Liddle, Interannual variation in the breeding biology of the Antarctic prion *Pachyptila desolata* at Bird Island, South Georgia, *J. Zool.* 234 (1994) 125–139.
- [10] J.P. Croxall, P.A. Prince, Food, feeding ecology and ecological segregation of seabirds at South Georgia, *Biol. J. Linn. Soc.* 1 (1980) 103–131.
- [11] S. Marchant, P.J. Higgins, *Handbook of Australian, New Zealand and Antarctic Birds*, 1, M. Sharp, Melbourne, 1990.
- [12] T.D. Williams, *The Penguins: Spheniscidae*, Oxford University Press, New York, 1995.
- [13] P. Jouventin, R.J. Cuthbert, R. Ottvall, Genetic isolation and divergence in sexual traits: evidence for the northern rockhopper penguin *Eudyptes moseleyi* being a sibling species, *Mol. Ecol.* 15 (2006) 3413–3423.
- [14] R.W. Rand, The biology of guano-producing seabirds: the distribution, abundance and feeding habits of the Cape penguin, *Spheniscus demersus*, off the south western coast of the Cape Province, *Investig. Rep.* 41 (1960) 1–28.
- [15] R. Groscolas, Changes in plasma lipids during breeding, moulting and starvation in male and female emperor penguins (*Aptenodytes forsteri*), *Physiol. Zool.* 55 (1982) 45–55.
- [16] J. Prévost, *Ecologie du manchot empereur*, in : *Expéditions polaires françaises*, Hermann Press, Paris, 1961.
- [17] A. Barrat, Quelques aspects de la biologie et de l'écologie du manchot royal (*Aptenodytes patagonicus*) des îles Crozet, *CNFRA* 40 (1976) 9–51.
- [18] B. Stonehouse, The king penguin *Aptenodytes patagonicus* of South Georgia I. Breeding behaviour and development, *Sci. Rep. Falkland Islands Dependency Surv.* 23 (1960) 1–81.
- [19] C.L. Hull, M. Hindell, K. Le Mar, P. Scofield, J. Wilson, M.A. Lea, The breeding biology and factors affecting reproductive success in rockhopper penguin *Eudyptes chrysocome* at Macquarie Island, *Polar Biol.* 27 (2004) 711–720.
- [20] J. Warham, The rockhopper penguin, *Eudyptes chrysocome*, at Macquarie Island, *Auk* 80 (1963) 229–256.
- [21] J.O. Murie, L.S. Davis, I.G. McLean, Identifying the sex of Fionland crested penguins by morphometric characters, *Notornis* 38 (1991) 233–238.

- [22] J. Warham, The Fiordland crested penguin *Eudyptes pachyrhynchus*, Ibis 116 (1974) 1–27.
- [23] J. Warham, The breeding biology and behaviour of the Snares crested penguin, J.R. Soc. New Zeal. 4 (1974) 63–108.
- [24] R. Carrick, Population ecology of the Australian black-baked magpie, royal penguin, and silver gull. Population ecology of migratory birds, a symposium, US Dept Interior Res. Report 2 (1972) 41–98.
- [25] J. Warham, Aspects of the biology of the erect-crested penguin, *Eudyptes sclateri*, Ardea 60 (1972) 145–184.
- [26] F.C. Kinsky, R.A. Falla, A subspecific revision of the Australian blue penguin (*Eudyptula minor*) in the New Zealand area, Rec. Natl. Museum New Zeal. 1 (1976) 105–126.
- [27] P.J. Moore, Breeding biology of the yellow-eyed penguin *Megadyptes antipodes* on Campbell Island, Emu 92 (1993) 157–162.
- [28] D.G. Ainley, W.B. Emison, Sexual size dimorphism in Adélie penguins, Ibis 114 (1972) 267–271.
- [29] W.Z. Trivelpiece, S.G. Trivelpiece, Courtship period of Adélie, gentoo and chinstrap penguins, in : L.S. Davis, J.T. Darby (Eds.), Penguin biology, Academic Press, San Diego, 1990, pp. 113–127.
- [30] P.D. Boersma, D.L. Stokes, P.M. Yorio, Reproductive variability and historical change of Magellanic penguins (*Spheniscus magellanicus*), in : L.S. Davis, J.T. Darby (Eds.), Penguin biology, Academic Press, San Diego, 1990, pp. 15–43.
- [31] P. Isenmann, Contribution à l'éthologie et à l'écologie du manchot empereur (*Aptenodytes forsteri* Gray) à la colonie de Pointe Géologie (Terre Adélie), L'Oiseau et la R.F.O. 41 (1971) 9–64.
- [32] J.P. Croxall, P.A. Prince, Antarctic seabird and seal monitoring studies, Polar Rec. 19 (1979) 573–595.
- [33] T.D. Williams, S. Rodwell, Annual variation in return rate, mate and nest-site fidelity in breeding gentoo and macaroni penguins, Condor 94 (1992) 636–645.
- [34] L.E. Richdale, A population study of penguins, Oxford University Press, Oxford, 1957.
- [35] P. Isenmann, E.P. Jouventin, Eco-éthologie du Manchot empereur (*Aptenodytes forsteri*) et comparaison avec le Manchot Adélie (*Pygoscelis adeliae*) et le Manchot royal (*Aptenodytes patagonica*), L'Oiseau et la R.F.O. 40 (1970) 136–159.
- [36] T.L. Bucher, G.A. Bartholomew, W.Z. Trivelpiece, N.J. Volkman, Metabolism, growth and activity in Adélie and emperor penguin embryos, Auk 103 (1986) 485–493.
- [37] R. Guillard, J. Prévost, Observations écologiques à la colonie de manchots empereurs de Pointe Géologie (Terre Adélie) en 1963, L'Oiseau et la R.F.O. 34 (1964) 33–51.
- [38] J.L. Mougouin, Observations écologiques à la colonie de Pointe Géologie en 1964, L'Oiseau et la R.F.O. 36 (1966) 166–226.
- [39] R.W. Rand, Note on the birds of Marion Island, Ibis 96 (1954) 173–206.
- [40] M. Poisbleau, L. Demongin, I.J. Strange, H. Otley, P. Quillfeldt, Aspects of the breeding biology of the southern rockhopper penguin *Eudyptes c. chrysocome* and new consideration on the intrinsic capacity of the A-egg, Polar Biol. 31 (2008) 925–932.
- [41] J.C. Stahl, P. Derenne, P. Jouventin, J.L. Mougouin, L. Teulière, H. Weimerskirch, Le cycle reproducteur des gorfous de l'archipel Crozet: *Eudyptes chrysolophus*, le gorfou macaroni et *Eudyptes chrysocome*, le gorfou sauteur, L'Oiseau et la R.F.O. 55 (1985) 27–43.
- [42] J. Warham, Breeding season and sexual dimorphism in rockhopper penguins, Auk 89 (1972) 86–105.
- [43] A.J. Williams, Offspring reduction in macaroni and rockhopper penguins, Auk 97 (1980) 754–759.
- [44] A.M. Gwynn, The egg-laying and incubation periods of rockhopper, macaroni and gentoo penguins, ANARE Report B 1 (1953) 1–29.
- [45] T.D. Williams, J.P. Croxall, Annual variation in breeding biology of macaroni penguins, *Eudyptes chrysolophus*, at Bird Island, South Georgia, J. Zool. 223 (1991) 189–202.
- [46] C.C. St. Clair, Incubation behaviour, brood patch formation and obligate brood reduction in Fiordland crested penguins, Behav. Ecol. Sociobiol. 31 (1992) 409–416.
- [47] C.M. Miskelly, P.W. Carey, Egg laying and egg loss by erect-crested penguin, in : University of Canterbury Antipodes Island Expedition Report, University of Canterbury, New Zealand, 1990, pp. 2–11.
- [48] L.E. Richdale, The erect-crested penguin (*Eudyptes sclateri*), Emu 41 (1941) 25–53.
- [49] L.E. Richdale, Further notes on the erect-crested penguin, Emu 49 (1950) 153–166.
- [50] G.S. Lishman, The food and feeding ecology of Adélie penguins (*Pygoscelis adeliae*) and chinstrap penguins (*Pygoscelis antarctica*) at Signy Island, South Orkney Islands, J. Zool. 205 (1985) 245–263.
- [51] B. Reid, The Adélie penguin (*Pygoscelis adeliae*) egg, New Zeal. J. Sci. 8 (1965) 503–514.
- [52] R.H. Taylor, The Adélie penguin, *Pygoscelis adeliae*, at Cape Royds, Antarctica, Ibis 104 (1962) 176–204.
- [53] G.S. Lishman, The comparative breeding biology of Adélie and chinstrap penguins, *Pygoscelis adeliae* and *Pygoscelis antarctica*, at Signy Island, South Orkney Islands, Ibis 127 (1985) 84–99.
- [54] C.A. Bost, P. Jouventin, The breeding performance of the gentoo penguin *Pygoscelis papua* at the northern edge of its range, Ibis 133 (1991) 14–25.
- [55] A.J. Williams, Aspects of the breeding biology of the gentoo penguin *Pygoscelis papua*, Le Gerfaut 70 (1980) 283–295.
- [56] A.J. Williams, J. Cooper, Aspects of the breeding biology of the jackass penguin, *Spheniscus demersus*. Proceedings of the Fifth Pan-African Ornithological Conference, 1984, pp. 841–853.
- [57] J. Boswall, D. MacIver, The Magellanic penguin *Spheniscus magellanicus*, in : B. Stonehouse (Ed.), The biology of penguins, Macmillan, London, 1975, pp. 271–305.
- [58] Y. Le Maho, Emperor penguin - strategy to live and breed in cold, Am. Sci. 65 (1977) 680–693.
- [59] H. Weimerskirch, J.C. Stahl, P. Jouventin, The breeding biology and population dynamics of king penguins *Aptenodytes patagonica* on the Crozet Islands, Ibis 134 (1992) 107–117.
- [60] J.L. Mougouin, La ponte du gorfou macaroni, *Eudyptes chrysolophus*, de l'Archipel Crozet, L'Oiseau et la R.F.O. 54 (1984) 281–291.
- [61] J. Warham, Aspects of breeding behaviour in the royal penguin, *Eudyptes chrysolophus schlegeli*, Notornis 18 (1971) 91–115.
- [62] A.F. Chiaradia, K.R. Kerry, Daily nest attendance and breeding performance in the little penguin *Eudyptula minor* at Phillip Island, Australia, Mar. Ornithol. 27 (1999) 13–20.
- [63] N.J. Adams, Embryonic metabolism, energy budgets and cost of production of king *Aptenodytes patagonica* and gentoo *Pygoscelis papua* penguin eggs, Comp. Biochem. Phys. A 101 (1992) 497–503.
- [64] P. Dann, J.M. Cullen, Survival, patterns of reproduction and lifetime reproductive output in little blue penguins (*Eudyptula minor*) on Phillip Island, Victoria, Australia, in : L.S. Davis, J.T. Darby (Eds.), Penguin biology, Academic Press, San Diego, 1990, pp. 63–84.
- [65] A. Kemp, P. Dann, Egg size, incubation periods and hatching success of Little Penguins *Eudyptula minor*, Emu 101 (2001) 249–253.
- [66] A. Kato, A. Yoshioka, K. Sato, Foraging behavior of Adélie penguins during incubation period in Lützow-Holm Bay, Polar Biol. 32 (2009) 181–186.
- [67] W.J.L. Sladen, The *Pygoscelis* penguins. 1. Methods of study. 2. The Adélie penguin, Falkland Islands Dependency Survey, Sci. Rep. 17 (1958) 1–97.
- [68] B. Despin, Note préliminaire sur le manchot papou *Pygoscelis papua* de l'île de la Possession (archipel Crozet), L'Oiseau et la R.F.O. 42 (1972) 69–83.
- [69] A. Lescroët, C. Bajzak, C.A. Bost, Breeding ecology of the gentoo penguin *Pygoscelis papua* at Kerguelen Archipelago, Polar Biol. 32 (2009) 1495–1505.
- [70] P.D. Boersma, Breeding patterns of Galápagos penguins as an indicator of oceanographic conditions, Science 200 (1978) 1481–1483.
- [71] A. Chiaradia, I.C.T. Nisbet, Plasticity in parental provisioning and chick growth in Little Penguins *Eudyptula minor* in years of high and low breeding success, Ardea 94 (2006) 257–270.
- [72] F.H. Vargas, S. Harrison, S. Rea, D.W. Macdonald, Biological effects of El Niño on the Galápagos penguin, Biol. Conserv. 127 (2006) 107–114.
- [73] J.F. Cockrem, P.J. Seddon, Annual cycle of sex steroids in the yellow-eyed penguin (*Megadyptes antipodes*) on south-island, New-Zealand, Gen. Comp. Endocr. 94 (1994) 113–121.
- [74] M. Beaulieu, A.M. Thierry, Y. Le Maho, Y. Ropert-coudert, A. Ancel, Alloparental feeding in Adélie penguins: why is it uncommon? J. Ornithol. 150 (2009) 637–643.
- [75] C. Gilbert, G. Robertson, Y. Le Maho, A. Ancel, How do weather conditions affect the huddling behaviour of emperor penguins? Polar Biol. 31 (2008) 163–169.
- [76] V.A. Viblan, V. Valette, M. Kauffmann, N. Malosse, R. Groscolas, Coping with social stress: heart rate responses to agonistic interactions in king penguins, Behav. Ecol. 23 (2012) 1178–1185.
- [77] T. Aubin, P. Jouventin, How to vocally identify kin in a crowd: the penguin model, Adv. Stud. Behav. 31 (2002) 243–277.
- [78] R. Groscolas, J. Lecloup, The endocrine control of reproduction and molt in male and female Emperor (*Aptenodytes forsteri*) and Adélie (*Pygoscelis adeliae*) penguins. II. Annual changes in plasma levels of thyroxine and triiodothyronine, Gen. Comp. Endocr. 63 (1986) 264–274.
- [79] A. Ancel, C. Gilbert, M. Beaulieu, The long engagement of the emperor penguin, Polar Biol. (2013), <http://dx.doi.org/10.1007/s00300-013-1285-9>.
- [80] P. Dann, I. Norman, P. Reilly, The Penguins: Ecology and Management, Dann, Norman, Reilly, Australia, 1995.
- [81] P. Reilly, Penguins of the World, S. Dobney, Oxford University Press, Australia, 1994.

- [82] P. Yorio, P.D. Boersma, Causes of nest desertion during incubation in the magellanic penguin (*Spheniscus-magellanicus*), *Condor* 96 (1994) 1076–1083.
- [83] W.Z. Trivelpiece, S.G. Trivelpiece, N.J. Volkman, Ecological segregation of Adélie, gentoo and chinstrap penguins at King George Island, Antarctica, *Ecology* 68 (1987) 351–361.
- [84] E.J. Woehler, Consumption of Southern Ocean marine resources by penguins, in : P. Dann, I. Norman, P. Reilly (Eds.), *The penguins: Ecology and Management*, Surrey Beatty & Sons Pty Limited, Chipping Norton, 1995, pp. 266–294.
- [85] S. Robinson, A. Chiaradia, M.A. Hindell, The effect of body condition on the timing and success of breeding in Little Penguins *Eudyptula minor*, *Ibis* 147 (2005) 483–489.
- [86] IUCN 2012. IUCN Red list of threatened species, version 2012.2 [www.iucnredlist.org].
- [87] A.P. Clausen, N. Huin, Status and numerical trends of King, Gentoo, and Rockhopper Penguins breeding in the Falkland Islands, *Waterbirds* 26 (2003) 389–402.
- [88] M.I. Ghys, A.R. Rey, A. Schiavini, Population trend and breeding biology of Gentoo penguin in Martillo Island, Tierra Del Fuego, Argentina, *Waterbirds* 31 (2008) 625–631.
- [89] P.T. Fretwell, P.N. Trathan, Penguins from space: faecal stains reveal the location of emperor penguin colonies, *Global Ecol. Biogeogr.* 18 (2009) 1–10.
- [90] P.T. Fretwell, M.A. LaRue, P. Morin, G.L. Kooyman, B. Wienecke, N. Ratcliffe, A.J. Fox, A.H. Fleming, C. Porter, P.N. Trathan, An emperor penguin population estimate: the first global, synoptic survey of a species from space, *PLoS One* 7 (2012) e33751.
- [91] C. Barbraud, V. Rolland, S. Jenouvrier, M. Nevoux, K. Delord, H. Weimerskirch, Effects of climate change and fisheries bycatch on Southern Ocean seabirds: a review, *Mar. Ecol. Prog. Ser.* 454 (2012) 285–307.
- [92] J. Bried, F. Jiguet, P. Jouventin, Why do Aptenodytes penguins have high divorce rates? *Auk* 116 (1999) 504–512.
- [93] P. Jouventin, Comportement et structure sociale chez le manchot empereur, *La Terre et la Vie* 25 (1971) 510–586.
- [94] S. Jenouvrier, C. Barbraud, H. Weimerskirch, Long-term contrasted responses to climate of two Antarctic seabirds species, *Ecology* 86 (2005) 2889–2903.
- [95] C. Barbraud, H. Weimerskirch, Emperor penguins and climate change, *Nature* 411 (2001) 183–186.
- [96] L.B. Astheimer, C.R. Grau, A comparison of yolk growth rates in seabird eggs, *Ibis* 132 (1989) 380–394.
- [97] L.B. Astheimer, C.R. Grau, The timing and energetic consequences of egg formation in the Adélie penguin, *Condor* 87 (1985) 256–268.
- [98] C.J. Scholten, Choice of nest-site and mate in Humboldt penguins (*Spheniscus humboldti*), *Spheniscid*, *Penguin Newslett.* 5 (1992) 3–13.
- [99] E.B. Spurr, Behaviour of the Adélie penguin chick, *Condor* 77 (1975) 272–280.
- [100] A.J. Williams, The clutch size of macaroni and rockhopper penguins, *Emu* 81 (1981) 87–90.
- [101] A.J. Williams, The laying interval and incubation period of rockhopper and macaroni penguins, *Ostrich* 52 (1981) 226–229.
- [102] M. Beaulieu, A.M. Thierry, Y. Handrich, Y. Le Maho, S. Massemin, A. Ancel, Adverse effects of instrumentation in incubating Adélie penguins, *Polar Biol.* 33 (2010) 485–492.
- [103] J. Moreno, L.M. Carrascal, J.J. Sanz, J.A. Amat, J.J. Cuervo, Hatching asynchrony, sibling hierarchies and brood reduction in the chinstrap penguin *Pygoscelis Antarctica*, *Polar Biol.* 14 (1994) 21–30.
- [104] B.C. Wienecke, Adoption of chicks by Little Penguins *Eudyptula minor* on Penguin Island, Western Australia, *Emu* 95 (1995) 119–122.
- [105] P. Jouventin, C. Barbraud, M. Rubin, Adoption in the emperor penguin, *Aptenodytes forsteri*, *Anim. Behav.* 50 (1995) 1023–1029.
- [106] N. Lecomte, G. Kuntz, N. Lambert, J.P. Gendner, Y. Handrich, Y. Le Maho, C.A. Bost, Alloparental feeding in the king penguin, *Anim. Behav.* 71 (2006) 457–462.
- [107] G.L. Kooyman, E.C. Hunke, S.F. Ackley, R.P. Van Dam, G. Robertson, Moulting of the emperor penguin: travel, location, and habitat selection, *Mar. Ecol. Prog. Ser.* 204 (2000) 269–277.
- [108] B. Wienecke, R. Kirkwood, G. Robertson, Pre-moult foraging trips and moult locations of Emperor penguins at the Mawson Coast, *Polar Biol.* 27 (2004) 83–91.
- [109] I. Zimmer, R.P. Wilson, C. Gilbert, M. Beaulieu, A. Ancel, J. Plötz, Foraging movements of emperor penguins at Pointe Géologie, Antarctica, *Polar Biol.* 31 (2008) 229–243.
- [110] D.G. Ainley, R.C. Wood, W.L.J. Sladen, Bird life at Cape Crozier, Ross Island, *Wilson Bull.* 90 (1978) 492–510.
- [111] R. Groscolas, Study of molt fasting followed by an experimental forced fasting in emperor penguin *Aptenodytes forsteri* - relationship between feather growth, body-weight loss, body-temperature and plasma fuel levels, *Comp. Biochem. Phys. A* 61 (1978) 287–295.
- [112] Y. Le Maho, P. Delclitte, J. Chatonnet, Thermoregulation in fasting emperor penguins under natural conditions, *Am. J. Physiol.* 231 (1976) 913–922.
- [113] G.L. Kooyman, T.G. Kooyman, M. Horning, C.A. Kooyman, Penguin dispersal after fledging, *Nature* 383 (1996) 397.
- [114] C. Peron, H. Weimerskirch, C.A. Bost, Projected poleward shift of king penguins' (*Aptenodytes patagonicus*) foraging range at the Crozet Islands, southern Indian Ocean, *Proc. R. Soc. B-Biol. Sci.* 279 (2012) 2515–2523.
- [115] A. Scheffer, C.A. Bost, P.N. Trathan, Frontal zones, temperature gradient and depth characterize the foraging habitat of king penguins at South Georgia, *Mar. Ecol. Prog. Ser.* 465 (2012) 281–297.
- [116] P. Foxton, The distribution of the standing crop of zooplankton in the southern ocean, *Disc. Rep.* 28 (1956) 191–236.
- [117] T.J. Hart, Phytoplankton periodicity in Antarctic surface waters, *Disc. Rep.* 21 (1942) 261–356.
- [118] E.E. Koczyńska, N. Savoye, F. Dehairs, D. Cardinal, M. Elskens, Spring phytoplankton assemblages in the Southern Ocean between Australia and Antarctica, *Polar Biol.* 31 (2007) 77–88.
- [119] A. Ancel, G.L. Kooyman, P.J. Ponganis, J.P. Gendner, J. Lignon, X. Mestre, N. Huin, P.H. Thorson, P. Robisson, Y. Le Maho, Foraging behaviour of emperor penguins as a resource detector in winter and summer, *Nature* 360 (1992) 336–339.
- [120] A. Ancel, H. Visser, Y. Handrich, D. Masman, Y. Le Maho, Energy saving in huddling penguins, *Nature* 385 (1997) 336–339.
- [121] C. Gilbert, Y. Le Maho, M. Perret, A. Ancel, Body temperature changes induced by huddling in breeding male emperor penguins, *Am. J. Physiol. Reg. I* 292 (2007) R176–R185.
- [122] B.B. Hansen, V. Grøtan, R. Aanes, B.-E. Sæther, A. Stien, E. Fuglei, R.A. Ims, N.G. Yoccoz, A.Ø. Pedersen, Climate events synchronize the dynamics of a resident vertebrate community in the high arctic, *Science* 339 (2013) 313–315.
- [123] M.A. Hindell, C.J.A. Bradshaw, B.W. Brook, D.A. Fordham, K. Kerry, C. Hull, C.R. McMahon, Long-term breeding phenology shift in royal penguins, *Ecol. Evol.* 2 (2012) 1563–1571.
- [124] J. Forcada, P.N. Trathan, Penguin responses to climate change in the Southern Ocean, *Global Change Biol.* 15 (2009) 1618–1630.
- [125] A.M.M. Baylis, A.F. Zuur, P. Brickle, P.A. Pistorius, Climate as a driver of population variability in breeding Gentoo Penguins *Pygoscelis papua* at the Falkland Islands, *Ibis* 154 (2012) 30–41.
- [126] A. Barbosa, J. Benzal, A. De Leon, J. Moreno, Population decline of chinstrap penguins (*Pygoscelis antarctica*) on Deception Island, South Shetlands, Antarctica, *Polar Biol.* 35 (2012) 1453–1457.
- [127] J.T. Hinke, M.J. Polito, C.S. Reiss, S.G. Trivelpiece, W.Z. Trivelpiece, Flexible reproductive timing can buffer reproductive success of *Pygoscelis* spp. penguins in the Antarctic Peninsula region, *Mar. Ecol. Prog. Ser.* 454 (2012) 91–104.
- [128] H.J. Lynch, J. Heather, R. Naveen, P.N. Trathan, W.F. Fagan, Spatially integrated assessment reveals widespread changes in penguin populations on the Antarctic Peninsula, *Ecology* 93 (2012) 1367–1377.
- [129] A.R. Carlini, N.R. Coria, M.M. Santos, J. Negrete, M.A. Juarez, G.A. Daneri, Responses of *Pygoscelis adeliae* and *P. papua* populations to environmental changes at Isla 25 de Mayo (King George Island), *Polar Biol.* 32 (2009) 1427–1433.
- [130] H.J. Lynch, W.F. Fagan, R. Naveen, Population trends and reproductive success at a frequently visited penguin colony on the western Antarctic Peninsula, *Polar Biol.* 33 (2010) 493–503.
- [131] D. Ainley, J. Russell, S. Jenouvrier, E. Woehler, P.O. Lyver, W.R. Fraser, G.L. Kooyman, Antarctic penguin response to habitat change as Earth's troposphere reaches 2 degrees C above preindustrial levels, *Ecol. Monogr.* 80 (2010) 49–66.
- [132] R.A. Massom, K. Hill, C. Barbraud, N. Adams, A. Ancel, L. Emmerson, M.J. Pook, Fast ice distribution in Adélie Land, East Antarctica: inter-annual variability and implications for emperor penguins *Aptenodytes forsteri*, *Mar. Ecol. Prog. Ser.* 374 (2009) 243–257.
- [133] S. Jenouvrier, H. Caswell, C. Barbraud, M. Holland, J. Stroeve, H. Weimerskirch, Demographic models and IPCC climate projections predict the decline of an emperor penguin population, *Proc. Natl. Acad. Sci. USA* 106 (2009) 1844–1847.
- [134] S. Jenouvrier, M. Holland, J. Stroeve, C. Barbraud, H. Weimerskirch, M. Serreze, H. Caswell, Effects of climate change on an emperor penguin population: analysis of coupled demographic and climate models, *Global Change Biol.* 18 (2012) 2756–2770.
- [135] W.Z. Trivelpiece, J.T. Hinke, A.K. Miller, C.S. Reiss, S.G. Trivelpiece, G.M. Watters, Variability in krill biomass links harvesting and climate warming to penguin population changes in Antarctica, *Proc. Natl. Acad. Sci. USA* 108 (2011) 7625–7628.
- [136] Y. Ropert-Coudert, A. Kato, A. Chiaradia, Impact of small-scale environmental perturbations on local marine food resources: a case study

- of a predator, the little penguin, Proc. R. Soc. B-Biol. Sci. 276 (2009) 4105–4109.
- [137] R.B. Powell, M.T.J. Brownlee, S.R. Kellert, S.H. Ham, From awe to satisfaction: immediate affective responses to the Antarctic tourism experience, Polar Rec. 48 (2012) 145–156.
- [138] C. Villanueva, B.G. Walker, M. Bertellotti, A matter of history: effects of tourism on physiology, behaviour and breeding parameters in Magellanic Penguins (*Spheniscus magellanicus*) at two colonies in Argentina, J. Ornithol. 153 (2012) 219–228.
- [139] G.S. Fowler, Behavioral and hormonal responses of Magellanic penguins (*Spheniscus magellanicus*) to tourism and nest site visitation, Biol. Conserv. 90 (1999) 143–149.
- [140] B.G. Walker, P.D. Boersma, J.C. Wingfield, Physiological and behavioral differences in Magellanic Penguin chicks in undisturbed and tourist-visited locations of a colony, Conserv. Biol. 5 (2005) 1571–1577.
- [141] U. Ellenberg, A.N. Setiawan, A. Cree, D.M. Houston, P.J. Seddon, Elevated hormonal stress response and reduced reproductive output in yellow-eyed penguins exposed to unregulated tourism, Gen. Comp. Endocr. 152 (2007) 54–63.
- [142] M.R. McClung, P.J. Seddon, M. Massaro, A.N. Setiawan, Nature-based tourism impacts on yellow-eyed penguins *Megadyptes antipodes*: does unregulated visitor access affect fledging weight and juvenile survival? Biol. Conserv. 119 (2004) 279–285.