Feather corticosterone reveals effect of moulting conditions in the autumn on subsequent reproductive output and survival in an Arctic migratory bird

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For birds, unpredictable environments during the energetically stressful times of moulting and breeding are expected to have negative fitness effects. Detecting those effects however, might be difficult if individuals modulate their physiology and/or behaviours in ways to minimize short-term fitness costs. Corticosterone in feathers (CORTf) is thought to provide information on total baseline and stress-induced CORT levels at moulting and is an integrated measure of hypothalamic–pituitary–adrenal activity during the time feathers are grown. We predicted that CORTf levels in northern common eider females would relate to subsequent body condition, reproductive success and survival, in a population of eiders nesting in the eastern Canadian Arctic during a capricious period marked by annual avian cholera outbreaks. We collected CORTf data from feathers grown during previous moult in autumn and data on phenology of subsequent reproduction and survival for 242 eider females over 5 years. Using path analyses, we detected a direct relationship between CORTf and arrival date and body condition the following year. CORTf also had negative indirect relationships with both eider reproductive success and survival of eiders during an avian cholera outbreak. This indirect effect was dramatic with a reduction of approximately 30% in subsequent survival of eiders during an avian cholera outbreak when mean CORTf increased by 1 standard deviation. This study highlights the importance of events or processes occurring during moult on subsequent expression of life-history traits and relation to individual fitness, and shows that information from non-destructive sampling of individuals can track carry-over effects across seasons.

1. Introduction
To cope with unpredictable environments, individuals can modulate their physiology and behaviour to minimize short-term fitness costs. Although environmental factors can influence individual condition and fitness directly over the short term, they might be more likely to result in indirect consequences (carry-over effects, COEs) later in life [1,2]. COEs are defined as events or processes that occur in one season and that can affect an individual’s performance in a subsequent period [2]. COEs on the state of individuals can have important repercussions by magnifying or reducing population regulatory processes [3,4]. For migratory species, obtaining relevant metrics of individual state outside the
breeding period is challenging, which makes identifying linkages between conditions experienced at specific locations or habitats and phases of the life cycle also challenging, more so owing to the difficulty in tracking migrants across seasons and locations [2].

Studies of migratory bird species provide examples of COEs. In earlier studies, arrival date on the breeding grounds was shown to be related to factors that occurred prior to the breeding season such as use of high-quality versus marginal wintering habitats [5–7]. The importance of spring body condition on reproduction is another example of a COE reported in several income–capital breeders [8–11]. Recently, an experimental manipulation of greater snow geese (Anser caerulescens) showed that stressful events (captive) during migration affected subsequent reproduction [12]. Despite these examples, COEs are understudied, particularly in the context of novel environmental challenges such as climate change, food web disruption or emerging infectious diseases. Such factors have the potential to either magnify or ameliorate COEs making detection and the subsequent study of COEs and their impacts context-dependent. Also important in the study of COEs is the ease of measurement and reliability of indices by which conditions experienced ‘earlier on’ are assessed and tracked.

In birds, corticosterone (CORT) is the primary glucocorticoid released by activation of the hypothalamic–pituitary–adrenal (HPA) axis in response to a stressor [13]. CORT is responsible for mediating allostasis and promoting foraging and glucocorticoidogenesis [14]; high levels of CORT can be a consequence of exposure to a threat or a result of increasing requirements for energy, movement (locomotion) and/or metabolism [15–17]. The stress response and quantification of CORT in natural populations have become important components of many studies in ecology, physiology and conservation biology [18–22].

Stress hormone levels have been linked to body condition in several species, although the mechanisms are not clear [22]. In upland geese (Chloephaga picta leucoptera), individuals with higher faecal CORT levels had decreased body condition [23]. Experimentally, tree swallow (Tachycineta bicolor) nestlings that received CORT implants showed reduced growth rates compared to controls [24]. Since increased CORT may affect body condition, CORT levels may be linked to arrival date and reproductive success in subsequent seasons.

In birds, CORT is deposited into feathers during growth so that the amount of CORT measured in a feather can provide an index of an individual’s HPA activity during the growth of that feather and provides an integrated measure of CORT [24–26]. Previous studies have demonstrated the utility of measuring CORT in feathers and have shown that feather CORT (CORTf) is related to parental efficiency [26], social signals [27], nest microclimate [28], egg mass [29], cost of reproduction [30] and possibly climatic conditions [31]. Thus, CORTf has the potential to be used as a reliable index to study COEs of events occurring during the moulting period on the subsequent breeding period. In Anatidae, including northern common eiders (Somateria mollissima borealis), all flight feathers are moulted simultaneously once a year in late summer, after breeding. A feather collected in spring, prior to breeding, could thus provide an indication of the energetic demands experienced by the moulting birds approximately nine months earlier.

In addition to increasing energetic and catabolic costs, elevated CORT can also alter feather quality. Elevated levels of CORT in passerines during moulting can affect the rate of feather growth [13,24] and experimental increases in CORT resulted in a decline in feather quality [32]. If eiders with higher CORTf levels have diminished feather quality, this may lower their flight or foraging efficiency during the over-wintering period and result in negative COEs into the breeding season. In common eiders, CORT measured in feathers probably reflects responses to environmental conditions experienced by birds during moulting [31] and may potentially be used as a metric to study COEs of responses to climatic conditions on subsequent reproduction and survival many months later. Furthermore, it may also be used as a tool to examine the relationship between glucocorticoid responses during the molting period and infectious diseases.

Glucocorticoid levels can affect susceptibility to disease in many species, usually through effects on immune function [33] and energy metabolism. Exposure to chronically elevated CORT levels may decrease immune function [34] and increase susceptibility to disease, and even acute stress has been shown to affect survival of eiders in the face of an infectious disease outbreak [35]. Avian cholera (caused by the bacterium Pasteurella multocida) has been a cause of massive annual adult mortality in common eiders nesting at our study site in the low Arctic since 2005 [35,36]. Female eiders do not eat during the approximately 26 day incubation period. Egg laying and incubation are energetically demanding activities that may reduce immune function and future fecundity [37]. Large clutch sizes in eiders are associated with lower survival of female eiders in the face of severe avian cholera outbreaks [36]. Prior to 2005, avian cholera had not been documented in this population of eiders [35], and avian cholera in northern common eiders in the eastern Canadian arctic has previously only been documented in northern Quebec (Canadian Wildlife Health Cooperative, S. Iverson, N. J. Harms 2012, unpublished data).

Breeding success of common eiders is strongly influenced by body condition at time of breeding and by timing of migration [38]. Here, we expand the previous path analyses of Descamps et al. [38] by testing whether HPA activity during moulting could be carried-over approximately nine months later to affect the timing of migration and arrival condition, and have direct or indirect links to reproductive success and survival, in the face of avian cholera outbreaks. The unexpected appearance of annual disease outbreaks in our study colony was the impetus behind our investigation into the potential COEs of events occurring during moulting on the following breeding season.

2. Material and methods

(a) Study area and field methods

Eiders were captured on Mitivik Island (64°02’N, 81°47’W) in the East Bay Migratory Bird Sanctuary, Nunavut, Canada, from 2007 to 2013 [31,32]. Eiders were captured using large mist nets very early in the season when they were flying over the colony; we therefore assumed that capture date was a good proxy of arrival date [39]. At capture, body mass was measured using a Pesola scale (± 2.5 g), and one tail feather (second lateral right feather) was plucked from each individual and stored in an envelope in a dark and dry place until laboratory analyses. All eider flight feathers are moulted simultaneously once a year in late summer, after breeding. Commonly, eider tail feathers are grown during moulting in August–September, following the breeding season [40] and prior to autumn migration. Eiders from the eastern Canadian Arctic winter along the western coast of Greenland and northeastern coast of Labrador, Canada [40,41].

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Venous blood samples were collected from a subset of female eiders. Blood samples were collected from the tarsal vein within 3 min of capture, placed into heparinized tubes and centrifuged to harvest plasma. Plasma samples were stored frozen at −20°C in the field and −80°C in the laboratory until analysis for CORT. Although the plasma was collected as part of another study, we used the plasma CORT data to examine the relationship between plasma baseline CORT levels and CORTi. Eiders were banded with a metal band (United States Geological Survey) and two coloured alphanumeric Darvic bands (Pro-Touch, Saskatoon, Canada) [35]. All females were also marked with a unique colour and shape combination of two temporary plastic nasal markers (Junco Inc., Minneapolis, MN, USA) to enable identification of individual birds from a distance. We attached nasal markers with synthetic absorbable suture monofilament (Polydioxanone suture, 2.0 or 3.0 metric; Ethicon, Markham, Canada), so that nasal tags would be shed prior to autumn migration. We restricted the analyses to birds captured during the prelaying period to compare body condition and avoid any effect of egg laying on body mass. To do so, for each year, we included data only from birds caught before the date at which more than 2.5% of the population had started laying [31]. Individuals with known laying dates were subsequently added to the dataset if known laying date was later than capture date (with a buffer of 3 days to account for potential error on laying date estimation). Arrival and laying dates were standardized relative to the median (0—median arrival or laying date in each year). Because body mass alone is a better predictor of condition than mass corrected for body size in this species [38,39], body mass was used as our measure of condition. In 2011, 69 female eiders received subcutaneous corticosterone or sham implants (Innovative Research of America, Sarasota, FL, USA) as part of a separate study, and 44 of the implanted birds were included in this study. Such manipulation could have induced changes in eider reproductive outcomes, which could alter our conclusions. We performed all analyses with and without inclusion of eiders captured in 2011 to assess the robustness of our conclusions. Precise information on reproduction (lay date of the first egg and hatching success—at least one duckling hatched) for all eiders in this study was collected by monitoring nesting birds from eight observation blinds strategically located within the colony. Observation blinds allowed us to monitor over 90% of the eider nesting area [35] while minimizing disturbance to the colony. Final number of ducklings hatched was available for a very limited area [35] while minimizing disturbance to the colony. Final number of ducklings hatched was available for 242 eider females from 2007 to 2011 (table 1). Among the 242 females, only two females were sampled in 2 years representing less than 1% of the data. The two additional measurements of the same individuals were considered as being independent. We developed our path diagram (figure 1) from a similar analysis that tested a condition-dependent optimization model on the same eider population [38]. We expanded the relationships (both direct and indirect) to include CORTi, reproductive success and survival. Arrows in figure 1 indicate relationships between two variables following event chronology (moulting, pre-breeding and breeding). Arrival date and body condition were assessed at the same time so determining causation was not possible. Mortality of female eiders due to avian cholera was detected following nest initiation [35]. Therefore, we investigated whether lay date or nest success could affect the survival of eiders during an avian cholera outbreak.

(b) Corticosterone analysis

CORTi measurements were performed using a previously established protocol [25] that includes a methanol-based extraction followed by analysis of the extracts via radioimmunoassay. This method has been previously used for eider feathers [31] and other avian species [26,28,44,45]. In this study, we assessed the efficiency of methanol extraction by including eider feather samples spiked with a small amount (approx. 5000 CPM) of $^3$H-corticosterone in the extraction. Greater than 92% of the radioactivity was recoverable in the reconstituted samples. Bortolotti et al. [25,27] showed that CORT is deposited into feathers in a time-dependent fashion; therefore, our values are expressed as a function of feather length (pg mm$^{-1}$). All samples were measured in duplicate and were run randomized and blind. Assay variability was determined as the per cent coefficient of variation (CV) resulting from repeated measurement of samples spiked with a known amount of CORT in each assay. The average within-assay variation was 5.4% (range 2–10%), and inter-assay variation was 13.7%. Serial dilution of feather extracts from eider feathers produced displacement curves that were parallel to the standard curves. Hormone analyses were performed at the Department of Biology, University of Saskatchewan (Canada).

Baseline plasma CORT was analysed using a previously validated enzyme-linked immunoassay (Assay Design, Ann Arbor, MI, USA; [46]) run in triplicate at a 1:20 dilution with 1.5% of kit-provided steroid displacement buffer. Each plate was run with a kit-provided standard curve by serially diluting a 200 000 pg ml$^{-1}$ CORT standard and a control of laying hen plasma (Sigma-Aldrich Canada, Oakville, Ontario, Canada). Assay plates were read on a plate reader at a wavelength of 405 nm, and the mean inter- and intra-assay CV across all plates was 7.17% and 6.22%, respectively.

(c) Path analyses

The importance of HPA activity levels during pre-migratory moulting period on reproduction and survival was assessed using path analysis, a special form of structural equation model [47], following Shipley [48]. The principle of the method is to specify how the variables are linked together in terms of direct and indirect effects or relationships. Information on CORTi, condition at arrival, arrival date, reproduction (laying date, reproductive success) and survival was available for 242 eider females from 2007 to 2011 (table 1). Among the 242 females, only two females were sampled in 2 years representing less than 1% of the data. The two additional measurements of the same individuals were considered as being independent. We developed our path diagram (figure 1) from a similar analysis that tested a condition-dependent optimization model on the same eider population [38]. We expanded the relationships (both direct and indirect) to include CORTi, reproductive success and survival. Arrows in figure 1 indicate relationships between two variables following event chronology (moulting, pre-breeding and breeding). Arrival date and body condition were assessed at the same time so determining causation was not possible. Mortality of female eiders due to avian cholera was detected following nest initiation [35]. Therefore, we investigated whether lay date or nest success could affect the survival of eiders during an avian cholera outbreak.

The fit of a generalized multi-level path model was assessed using the concept of d-sep (directional separation) tests [47]. A d-sep test represents a test of the statistical independence between two variables. If two variables are d-separated relative
to a set of variables $Z$ in a directed graph, then they are independent conditional on $Z$ in all probability distributions such a graph can represent (see Shipley [48] for more details). Shipley [47] shows that for each acyclic path model, there is a subset of independence tests referred to as a ‘minimum basis set’ that account for all possible independence relationships (or claims). Model fit is evaluated using a set of $(k)$ mutually independent claims of probabilistic independence that must be true if the structure of the hypothesized model path is correct. The null hypothesis is that the proposed correlational structure of the model does not differ from the observed correlational structure in the data, and therefore $p \leq 0.05$ indicates the proposed causal structure is incorrect [47]. To calculate path coefficients, each variable was standardized ((value – average)/s.d.) such that path coefficients represent standardized partial regression coefficients, or the standard deviation change in $y$ when $x$ is increased or decreased by 1 s.d. [47].

Shipley [48] showed how the d-sep test can be combined with generalized linear mixed models. We followed detailed instructions provided in Shipley [48] using the packages nlmle and lme4 in R [49]. We used linear mixed models (using terms CORTf, arrival date, body condition, laying date, hatching success and survival) to regress each variable on its direct causes. A random Year effect was included in each model. The random effect accounted from 0.46 to 5.71% of the deviance explained depending on the dependent variable considered.

### 3. Results

The correlational structure of our path model (figure 2) was consistent with the correlational structure of the data (seven tests of probabilistic independence; Fisher’s $C_{14} = 8.82$, $p = 0.84$; implied independencies did not differ from those observed). The model defined in figure 1 provided a strong fit to the data as indicated by the high $p$-values (null probabilities) of the goodness-of-fit tests (table 2). Partial regression slope of CORTf was not different from zero in all claims revealing no direct effect of CORTf on lay date, reproductive success or survival. CORTf was significantly different among years ($F_{2,236} = 11.43; p < 0.001$). Arrival date on the breeding colony was positively associated with CORTf (figure 3a; $\beta = 0.26 \pm 0.13$ (s.e.); $t_{236} = 2.07; p = 0.04$). Lower body condition (body mass) at arrival was related to higher levels of CORTf (figure 3b; $\beta = -22.56 \pm 6.09$; $t_{236} = -3.71; p < 0.001$). There was no direct relationship between CORTf and lay date ($t_{236} = 1.50; p = 0.14$), reproductive success ($t_{236} = -1.38; p = 0.17$) or survival eiders during the avian cholera outbreak ($t_{236} = -0.16; p = 0.87$). We found that body condition increased over time during the pre-breeding period ($\beta = 6.78 \pm 3.14; t_{236} = 2.16; p = 0.03$), so that birds arriving later were in better body condition. As expected, arrival date was positively linked to lay date and birds that arrived earlier laid earlier, and eiders that arrived in better body condition were in better pre-laying condition.

### Table 1. Description of model variables by year for female northern common eiders captured on East Bay Island, Nunavut, Canada.

<table>
<thead>
<tr>
<th>Year</th>
<th>n common eiders</th>
<th>CORTf (pg mm$^{-1}$) (mean ± s.d.)</th>
<th>body condition (kg) (mean ± s.d.)</th>
<th>Julian arrival date (mean ± s.d.)</th>
<th>Julian lay date (mean ± s.d.)</th>
<th>reproductive success (%)</th>
<th>% eiders survived outbreak</th>
</tr>
</thead>
<tbody>
<tr>
<td>2007</td>
<td>106</td>
<td>5.44 ± 2.00</td>
<td>2.191 ± 0.165</td>
<td>172.7 ± 3.2</td>
<td>183.7 ± 5.2</td>
<td>42.5</td>
<td>83</td>
</tr>
<tr>
<td>2008</td>
<td>38</td>
<td>4.33 ± 1.23</td>
<td>2.218 ± 0.132</td>
<td>171.2 ± 3.3</td>
<td>179.3 ± 5.5</td>
<td>44.7</td>
<td>73.7</td>
</tr>
<tr>
<td>2009</td>
<td>44</td>
<td>4.05 ± 1.23</td>
<td>2.225 ± 0.167</td>
<td>176.4 ± 4.2</td>
<td>184.8 ± 5.7</td>
<td>20.5</td>
<td>95.5</td>
</tr>
<tr>
<td>2010</td>
<td>10</td>
<td>3.58 ± 0.97</td>
<td>2.213 ± 0.164</td>
<td>166.4 ± 4.9</td>
<td>176.6 ± 5.2</td>
<td>50.0</td>
<td>100.0</td>
</tr>
<tr>
<td>2011</td>
<td>44</td>
<td>5.82 ± 1.61</td>
<td>2.255 ± 0.206</td>
<td>170.0 ± 2.5</td>
<td>179.8 ± 4.5</td>
<td>50.0</td>
<td>100.0</td>
</tr>
</tbody>
</table>
Table 2. Test of conditional independence implied by the path diagram (Figure 1). \((X; Y) \mid Z\) means that variables \(X\) and \(Y\) are independent conditional of variable \(Z\) (i.e. if \(Z\) is held constant, variation in \(X\) does not imply variation in \(Y\)). The associated mixed model used to test the independence claims is \(Y \sim Z + X + 1\mid \text{Year}\), where \(\text{Year}\) represents a random effect. The variable whose partial regression includes zero is \(X1\) for all claims. Variables: \(X1\) (CORTf), \(X2\) (Arrival date), \(X3\) (Body condition), \(X4\) (Laying date), \(X5\) (Hatching success) and \(X6\) (Survival of an avian cholera outbreak).

<table>
<thead>
<tr>
<th>d-sep claim of independence</th>
<th>mixed model</th>
<th>partial regression slope (s.e.)</th>
<th>null probability</th>
<th>(distribution)</th>
</tr>
</thead>
<tbody>
<tr>
<td>((X1,X4)\mid (X2,X3))</td>
<td>(X4 \sim X1 + X2 + X3 + {\text{Year}})</td>
<td>0.072 (0.176)</td>
<td>0.68</td>
<td>(normal)</td>
</tr>
<tr>
<td>((X1,X5)\mid (X4))</td>
<td>(X5 \sim X4 + X1 + {\text{Year}})</td>
<td>-0.046 (0.090)</td>
<td>0.61</td>
<td>(binomial)</td>
</tr>
<tr>
<td>((X2,X5)\mid (X1,X4))</td>
<td>(X5 \sim X4 + X1 + X2 + {\text{Year}})</td>
<td>0.014 (0.056)</td>
<td>0.89</td>
<td>(binomial)</td>
</tr>
<tr>
<td>((X1,X6)\mid (X4))</td>
<td>(X6 \sim X4 + X1 + {\text{Year}})</td>
<td>0.014 (0.039)</td>
<td>0.71</td>
<td>(binomial)</td>
</tr>
<tr>
<td>((X2,X6)\mid (X1,X4))</td>
<td>(X6 \sim X4 + X1 + X2 + {\text{Year}})</td>
<td>-0.074 (0.069)</td>
<td>0.28</td>
<td>(binomial)</td>
</tr>
<tr>
<td>((X3,X6)\mid (X1,X4))</td>
<td>(X6 \sim X4 + X1 + X3 + {\text{Year}})</td>
<td>0.010 (0.039)</td>
<td>0.79</td>
<td>(binomial)</td>
</tr>
<tr>
<td>((X4,X6)\mid (X5))</td>
<td>(X6 \sim X4 + X5 + {\text{Year}})</td>
<td>0.057 (0.046)</td>
<td>0.21</td>
<td>(binomial)</td>
</tr>
</tbody>
</table>

Figure 2. Standardized path coefficients in hypothesized structural model. Bold numbers are standardized beta coefficients with s.e in brackets (\(n = 242\)).

4. Discussion

Our results provide evidence that energetic management during the moulting period, reflected by CORT levels in feathers, can be carried over to the subsequent breeding season and affect reproductive success and survival. Using path analyses, we detected a direct relationship between CORTf levels during moult and body condition and arrival date the following year, and an indirect negative relationship between reproductive success and survival. The magnitude of the indirect relationship between CORTf and fitness parameters was important, with a decrease of approximately 0.25 s.d. from the mean of reproductive success and survival for every increase of 1 s.d. of CORTf. The importance of arrival date and condition on reproductive success was expected given that the path analysis developed here is an extension of Descamps et al. [38], who found similar relationships in accordance with the condition-dependent optimization model [9,50]. As per causal pathways drawn from the optimization model [38,50] birds can adjust their reproductive decisions as a function of their arrival date and body condition at arrival. Our study is probably unique in showing that both these variables can be significantly related to CORTf levels, which may reflect a level of energetic management experienced during the previous year. Increases in energetic challenges or response to stressors experienced during the time of moult in autumn can ultimately have significant fitness consequences, indirectly affecting both reproduction and survival in the following breeding season.

Another possible explanation is that CORTf values reflect basal CORT levels of individuals regardless of the time period considered. This hypothesis would gain credence if baseline CORT during the breeding or pre-breeding period could predict relative fitness of individuals [51–54]. However, we found no evidence for any relationship between CORTf...
Figure 3. Relationship between feather corticosterone (CORTf), arrival date (a) and body condition (b) in female eiders. The model presented here controls for the other covariates (either mean arrival date or condition). Mean and s.e. are provided. Dot sizes are proportional to log (n). The fitted mixed linear model (black line) and its confidence interval at 95% (grey line) are shown.

This agrees with previous studies from our research group [36,62] suggesting that breeding decision, reproductive investment and the duration of exposure to disease at the nesting colony are key factors explaining survival of eiders facing avian cholera outbreaks.

5. Conclusion

Our work emphasizes the importance of determining how events are linked throughout the annual cycle to better understand population dynamics of migratory animals. Our approach also highlights the importance of energetic management challenges outside the breeding period (possibly generated by climatic variability) that can have subsequent carry-over effects on reproduction and survival during outbreaks of avian cholera, an emerging disease in arctic-nesting common eiders. Little is known about the molting period for many bird species, including eiders [63], thus our results shed some light on a relatively unknown stage of the annual cycle. Combining information that can be gained from non-destructively sampling a single feather, including stable isotopes [64–66], coloration [5] or physiological analysis such as hormone levels [98] can contribute to tracking COEs across seasons. Furthermore, considering both direct and indirect pathways may be required to understand relationships among spatio-temporally distinct events affecting individual fitness.

Ethics statement. This study adhered to guidelines of the Canadian Council on Animal Care, and all protocols were reviewed and approved by the University Committee on Animal Care and Supply—Animal Research Ethics Board of the University of Saskatchewan (Protocol no. 20100063 to C.S.), the University Committee on Animal Care of the University of Windsor (Protocol no. 11–06 to O.P.L.), Environment Canada’s Animal Care Committee (Protocol no: EC-PN-07–008 (2007), EC-PN-08–026 to EC-PN-11–026 (2008–2011) to H.G.G.).

Data accessibility. Common eider body condition, arrival date, body condition, feather corticosterone, survival and reproductive data: Dryad doi.org/10.5061/dryad.rp30d.

Acknowledgements. Many thanks to H. Hennin, H. Remenda, W. van Dijk, S. Cabezas, V. Fachal, J. McLeod and T. Marchant for assistance in the laboratory, and thanks to the numerous students and researchers, who assisted us in the field from 2007 to 2011. We thank S. Descamps, R. G. Clark, N. G. Yoccoz and an anonymous reviewer, and G. Fairhurst for helpful comments on the methods and manuscript.

Funding statement. Funding for this work was provided by the National Science and Engineering Council of Canada, Environment Canada, Duck Unlimited Institute for Wetland and Waterfowl Research, the
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