Shellfish Dredging Pushes a Flexible Avian Top Predator out of a Marine Protected Area

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There is a widespread concern about the direct and indirect effects of industrial fisheries; this concern is particularly pertinent for so-called “marine protected areas” (MPAs), which should be safeguarded by national and international law. The intertidal flats of the Dutch Wadden Sea are a State Nature Monument and are protected under the Ramsar convention and the European Union’s Habitat and Birds Directives. Until 2004, the Dutch government granted permission for ~75% of the intertidal flats to be exploited by mechanical dredgers for edible cockles (Cerastoderma edule). Here we show that dredged areas belonged to the limited area of intertidal flats that were of sufficient quality for red knots (Calidris canutus islandica), a long-distance migrant molluscivore specialist, to feed. Dredging led to relatively lower settlement rates of cockles and also reduced their quality (ratio of flesh to shell). From 1998 to 2002, red knots increased gizzard mass to compensate for a gradual loss in shellfish quality, but this compensation was not sufficient and led to decreases in local survival. Therefore, the gradual destruction of the necessary intertidal resources explains both the loss of red knots from the Dutch Wadden Sea and the decline of the European wintering population. This study shows that MPAs that do not provide adequate protection from fishing may fail in their conservation objectives.

Introduction

Worldwide benthic communities of intertidal soft sediments are seriously threatened by bottom-touching fisheries such as dredging [1]. Not only do target species directly decline in numbers because of removal, but target and non-target species are also indirectly affected [2,3]. In turn, predators such as migrant shorebirds, which make a living from benthic invertebrates, are declining in many parts of the world, and a link with commercial fishing activities seems evident [4–6]. Until 2004, three-quarters of the intertidal flats of the Dutch Wadden Sea were open to mechanical dredging for edible cockles, despite the high-level conservation status, despite the concerns about the damaging effects of cockle dredging to intertidal ecosystems, and despite the limited economic value of cockle dredging (a total of only 11 license holders maintained a fleet of just 22 fishing boats [7]).

A direct, immediate effect of dredging is the complete removal of all organisms larger than 19 mm in the 5-cm top layer. Because the sites dredged are usually the most biodiverse (C Kraan, T Piersma, A Dekinga, A Koolhaas, J Van der Meer, unpublished data), dredging may also affect smaller cockles; other bivalves such as blue mussels (Mytilus edulis), Baltic tellins (Macoma balthica), and sandgapers (Mya arenaria); polychaetes; and crustaceans such as shorecrabs (Carcinus maenas). More indirectly and over longer time scales, sediments become coarser after dredging events, leading to reduced settlement success in both cockeans and Baltic tellins [3]. Because grain size affects the feeding performance of bivalve mussels [8], one expects prey condition or quality (defined as flesh-to-shell ratio [9]) to be negatively affected by dredging activities. We may thus expect mechanical dredging to lead to both short- and long-term declines in quantity and quality of a variety of macrobenthic organisms.

Shellfish-eating shorebirds can cope with reductions in both prey density and in prey quality, as long as the reductions are not too large and do not occur simultaneously (Figure 1). Red knots have been shown to flexibly adjust their digestive capacity to food quality so that food processing rates are just sufficient to obtain the daily amount of energy (i.e., flesh) required to maintain energy balance [9,10]. Maintaining a larger digestive system would be costly in terms of maintenance and transport costs [11]. Therefore, a knot’s intake rate is often constrained by a digestive bottleneck [9,12]. This implies that a decline in prey density, although it may hamper maximum rates of prey collection (dictated by so-called “short-term functional responses” such as Holling’s type II disk equation), will often only marginally affect the digestively constrained intake rate over the full low-tide period (Figure 1). Likewise, as the digestive system is flexible and can be adjusted rapidly [13], a decline in food quality can be compensated for by an increase in digestive capacity (Figure 1). By contrast, once both density and quality...
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Figure 1. Conceptualization of Why a Simultaneous Reduction in Density and Quality of the Prey is Detrimental

Holling’s Type II functional response describes intake rate (be it flesh or energy) as a function of the density of either poor-quality (black lines) or good-quality (gray lines) prey. Digestive constraint limits shell-mass processing rate and is given for two gizzard sizes for each prey quality (horizontal cut-offs in functional response; digestively unconstrained intake rates continue as dashed lines). By knowing the threshold intake rate needed to avoid starvation (border between gray and white background), one can predict a bird’s starvation chances on the basis of gizzard size and prey quality and density: (1) A small gizzard is sufficient to stay alive when prey is of good quality and occurs in high densities. Going from (1) to (2), prey density is reduced, which does not affect survival as intake rate remains above the critical threshold. Going from (1) to (3), prey quality (flesh-to-shell ratio) is reduced. To maintain a sufficient intake rate, the knot needs to increase its shell-mass processing rate, which requires a gizzard enlargement. Going from (1) to (4), the combined reduction in density and quality makes a gizzard enlargement no longer sufficient (as intake rate is now constrained by prey density), and the bird is bound to starve.

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Results/Discussion

Densities of small cockles remained stable in areas mechanically dredged, whereas they increased by 2.6% per year in undredged areas (general linear model [GLM] with \( n = 271 \) blocks: \( R^2 = 0.02, p = 0.04 \) (Figure 2B). Moreover, quality of small cockles declined by 11.3% per year in dredged areas and remained stable in undredged areas (GLM with \( n = 59 \) blocks: \( R^2 = 0.07, p = 0.04 \) (Figure 2B). These results are consistent with a previous assessment [3] that showed dredged areas to become unattractive areas for cockles to settle, a finding that was explained by dredged sediments losing silt and becoming coarser. In deposit-feeding bivalves such as freshly settled cockles [18,19], coarser sediments may lead to worse feeding conditions [8] and therefore to reduced body condition, which, from the predator’s point of view, equals reduced prey quality [20].

To express changes in carrying capacity in relation to dredging activity, we calculated for each year the percentage of blocks that would yield insufficient intake rates for knots to maintain a positive energy balance (taking 4.8 W as the critical metabolizable energy intake rate [21]). During our study period, the percentage of km² blocks that were too poor for knots to feed increased from 66% in 1998 to 87% in 2002 (GLM with \( n = 5 y; R^2 = 0.80, p = 0.04 \) (Figure 3A). This was entirely due to an increase in previously suitable blocks that were dredged (GLM with \( n = 5 y; R^2 = 0.96, p = 0.003 \)) (Figure 3A, dark gray bars). We calculated that of this 21% increase in unsuitability, 8% was due to a decline in prey densities alone, whereas the remaining 13% was due to the simultaneous decline in density and quality (note that these calculations include all potential prey species, not only cockles). In contrast, the percentage of unsuitable blocks that were never dredged before did not change (GLM with \( n = 5 y; R^2 = 0.02, p = 0.80 \) (Figure 3A, light gray bars). As a consequence of the widespread dredging in the best areas, diet quality declined by 11.7% per year (GLM with \( n = 174 \) dropping samples: \( R^2 = 0.05, p = 0.003 \)) (Figure 3B), a similar rate of decline (11.3%) as shown by the quality of cockles, the knots’ main prey, in dredged areas. To compensate for reductions in diet quality, knots should increase gizzard mass [9,13]. Indeed, in the course of the study period, gizzard mass increased by 3.4% per year (GLM with \( n = 644 \) birds: \( R^2 = 0.02, p = 0.0001 \)) (Figure 3C). This increase in gizzard mass exactly matches our quantitative expectations, because experimental results on shell mass processing rates [9] imply that gizzard mass should vary inversely with the square root of prey quality (i.e., square root of the annual 11.7% decline in prey quality = 3.4%). More specifically, gizzard masses do not deviate from the predicted gizzard masses required to balance the energy budget (predicted from yearly average prey quality values [9,10], \( p > 0.1, n = 484 \)).

Nevertheless, resightings of individually color-banded birds, whose gizzards were measured before release, suggest
that on average, birds not seen in our study area within the year after release had undersized gizzards (\(p < 0.01, n = 362\)), whereas individuals that we did see again had gizzards that enabled them to achieve a balanced daily energy budget (\(p = 0.4\), \(n = 122\)) (Figure 4A). That birds returning from the high-arctic breeding grounds with undersized gizzards [22] do not simply enlarge their gizzard may be explained by time and energy constraints [13,23]. Birds arriving with too small a gizzard may need more time to adjust their gizzard than their fat stores allow them, and therefore they face starvation unless they leave the area (indeed we found that birds with a smaller gizzard had a lower body mass; \(R^2 = 0.06, p < 0.00001\), \(n = 483\)). This implies that the proportion of birds arriving with a gizzard large enough to survive increases as a function of food quality at the arrival destination (Figure 4A and 4B). Indeed, local annual survival rate (calculated from resighting rates of color-banded birds) increased with food quality in the expected direction (Figure 4C). When we estimated the distribution of gizzard masses upon arrival (by selecting only those birds caught in the Wadden Sea during late July) and calculated, for various degrees of gizzard flexibility, the proportion of birds that would survive the critical gizzard-enlargement phase (Figure 4B and 4C), the best fit with the actual survival data was obtained if knots had time to increase their gizzard by 1 g only (Figure 4C).

Color-banded knots that disappeared from our study area may have died or, perhaps more likely for a wide-ranging migrant, emigrated to other areas such as the estuaries in the United Kingdom, where they probably paid a mortality cost due to the extra travel and/or due to uncertainties in the food supply at their new destination (where they also fed on hard-shelled prey, in the past [24,25] and more recently [26,27], and thus faced gizzard-related mortality; note that knots are “forced” to feed on hard-shelled prey as their pressure-sensitive bill tip can only detect hard objects buried in soft sediments [28]). In any case, the declining numbers of knots wintering in the Dutch Wadden Sea [29] can be explained as a response to declining food conditions. Moreover, the 25% decline of the entire northwestern European wintering population between 1997–1998 and 2002–2003 (from ~330,000 to ~250,000) (G Austin, M Van Roomen, B Koks, T Piersma, unpublished data) can be explained by measured decreases in local survival of the Wadden Sea segment of the population during the study period (using the observed local survival rates, we estimated an extra mortality of 58,000 birds over the 5-y period). This study concludes that industrial forms of commercial exploitation of protected marine nature reserves in The Netherlands, by indirectly reducing food resource quality to such extents that changes can no longer be accommodated by adjustments of the digestive system, are directly responsible for the overall population decline of a fully protected shorebird species. This paper thereby adds to growing list of studies [17,30] stressing the uselessness of declaring a marine area as protected whenever the species living in it are seriously affected by the ongoing but regulated human activities.

**Materials and Methods**

**Sampling prey density and quality.** From late July to early September 1998–2002, we sampled macrozoobenthos throughout the western Dutch Wadden Sea in a regular grid (250-m grid intersections; Figure 2A). In total, we visited 2,846 stations, of which the majority (75%) were sampled each year (89% in 4 out of 5 y). Stations were located using handheld global positioning system (GPS) receivers (Garmin 45 and 12; Garmin Corporation, Lenexa, Kansas,
United States) and were either visited during low tide (on foot) or during high tide (by rubber boat). At each station, a sediment core was taken (15-cm diameter; 20-cm deep) that was sieved over a 1-mm mesh. To distinguish accessible prey from prey living beyond the reach of a knot’s bill (4 cm), we sieved the top layer (upper 4 cm) separately from the bottom layer (only the “low-tide samples”). Mudsnails (Hydrobia ulvae) were sampled using a smaller core (7-cm diameter) and a finer mesh (0.5 mm). All potential prey items retained on the sieve were frozen (−20 °C) for later analyses. In the laboratory, items were identified with respect to species and size (to nearest mm; or, in the case of H. ulvae, to nearest 0.5 mm). Size classes were determined to distinguish ingestible prey from prey too large to be swallowed [12]. Methods to determine flesh and shell mass are explained elsewhere [3, 31].

Mechanical dredging and its effects on cockle density and quality. During each of the 5 y studied, mechanical dredging took place after our sampling program; i.e., from early September into December. Exact locations of dredging were known, because for reasons of internal control, every vessel had a GPS-logger onboard [7]. Cumulative seasonal data on dredging locations were available in the form of fine-scaled maps (resolution of 0.1 min latitude by 0.5 min longitude).

We analyzed the effects of mechanical dredging on cockles that were actually available to knots (i.e., <16 mm, because larger ones cannot be swallowed [32]). We pooled sampling stations in 272 blocks measuring 1 km by 1 km each (Figure 2A; many sampling stations did not have cockles available during multiple years making it impossible to study changes in quality at the spatial scale of stations). A block was considered dredged when at least one station was dredged in at least 1 y. Per block, we applied the following GLMs: log10(DENSITY + 1) = CONSTANT + YEAR and log10(QUALITY) = CONSTANT + YEAR. Subsequently, we tested whether significant variation in the coefficients for YEAR could be explained by whether a block was ever dredged or not during 1998–2001 (again using GLM).

Suitability for knots. The measurements on (available) prey densities and qualities allowed us to predict for each station an intake rate for a knot with an average-sized gizzard (6-g fresh mass). We did so by applying the so-called “digestive rate model,” a multi-species functional response that takes rates of digestion into account.
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2. Hiddink JG (2003) Effects of suction-dredging for cockles on non-target

1. Jackson JBC, Kirby MX, Berger WH, Bjorndal KA, Botsford LW, et al. (2001)

rate sufficient to maintain energy balance when feeding for 10 h per
day (corresponding to an intake rate of at least 0.5 mg dry flesh mass
s⁻¹), which is realizable energy intake rate of at least 4.8 W; note that the energy contents of 1 g of flesh is more or less constant in
mollusks [34]). This approach is widely used [35,36] and adequately
predicted the distribution of knots in an earlier study [21].

Diet, gizzard sizes, and observed and predicted survival rates. Diet
quality was reconstructed by fecal analyses, following the procedures
outlined by Dekinga and Piersma [37]. Dropping samples were
collected during low tide at sites where flocks of knots had just fed.
Across the 5 y of study, we analyzed 174 dropping samples,
comprising 1–106 droppings (each 150 g average).

In order to estimate gizzard sizes and survival rates, we annually
mistnetted knots from late July to late November, mostly near their
main roost at Richel (an island located midnorth of our study area;
Figure 2A), but occasionally near the isle of Schiermonnikoog (max. 75 km
ENE of Richel). Based on the presence or absence of active wing molt
[15], we determined subspecific identity and selected for the current
analyses islandica individuals only (population numbers and survival
rates are best known in this subspecies). Gizzard masses were estimated
using ultrasonography (Pie 200 ultrasound, Pie Medical Benelux BV,
Maasbracht, the Netherlands; applied to 644 out of the 1,069
individuals selected). This method has been extensively calibrated
[38] and has been successfully applied in earlier studies [9,12,13, 31].
Furthermore, to estimate survival rates, each bird was given a unique
combination of color bands. We used the Cormack-Jolly-Seber model of
the MARK software package [39] to estimate “local survival,” i.e., the
actual survival during the first year after capture minus the (unknown)
fraction of birds that emigrated permanently from our study area.
In the model, annual survival rate was allowed to differ in the first
year after capture and was assumed constant thereafter. This can be
biologically interpreted as a variable proportion of birds becoming
site faithful in each year. In case permanent emigration leads to death,
the extra mortality equals the difference between first-year-after-
capture global survival rate (mean = 0.73; SE = 0.03; calculated over all
resightings, both inside and outside the Wadden Sea) and first-year-
after-capture local survival rate. Assuming that each autumn, 100,000
individuals “try out” the Wadden Sea, this extra mortality in the NW-
European wintering population as the result of emigration can thus be
estimated. Note that we had 4 rather than 5 y of data, because our
color-band program started in 1998, yielding the first survival estimate for 1998–1999.

According to the following procedure, we predicted local survival rate
for various degrees of gizzard flexibility (0–2 g) upon arrival in
our study area (lines in Figure 4C). Based on experimental results on
shell mass processing rates as a function of gizzard size [9], we
calculated the minimal gizzard size required to avoid starvation as a
function of prey quality (line in Figure 4A). Using the observed
distribution of gizzard masses upon arrival (0-g distribution
in Figure 4B, representing knots caught during late July only; n = 218;
mean = 5.92; variance = 2.61), we then calculated the proportion of
birds having a gizzard of at least this critical size, which would be the
proportion of birds able to survive in our study area (47% at the
example prey quality of 0.15 g flesh per g shell in Figure 4). Assuming
that knots upon arrival have the flexibility to increase their gizzard
diameter by an 1-g, we calculated an updated “effective” gizzard
mass distribution upon arrival (i.e., a distribution that shifted by e.g.,
1 g; Figure 4B), yielding an updated proportion of birds with a large
enough gizzard, i.e., able to survive (70% for the +1-g example).
We refer to Van Gils et al. [9,10,20] for more details on modeling gizzard
masses.

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