



Modeling the aggregated exposure and responses of bowhead whales *Balaena mysticetus* to multiple sources of anthropogenic underwater sound

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ABSTRACT: Potential responses of marine mammals to anthropogenic underwater sound are usually assessed by researchers and regulators on the basis of exposure to a single, relatively loud sound source. However, marine mammals typically receive sounds from multiple, dynamic sources. We developed a method to aggregate modeled sounds from multiple sources and estimate the sound levels received by individuals. To illustrate the method, we modeled the sound fields of 9 sources associated with oil development and estimated the sound received over 47 d by a population of 10 000 simulated bowhead whales *Balaena mysticetus* on their annual migration through the Alaskan Beaufort Sea. Empirical data were sufficient to parameterize simulations of the distribution of individual whales over time and their range of movement patterns. We ran 2 simulations to estimate the sound exposure history and distances traveled by bowhead whales: one in which they could change their movement paths (avert) in response to set levels of sound and one in which they could not avert. When animals could not avert, about 2% of the simulated population was exposed to root mean square (rms) sound pressure levels (SPL) ≥ 180 dB re 1 μ Pa, a level that regulators in the U.S. often associate with injury. When animals could avert from sound levels that regulators often associate with behavioral disturbance (rms SPL > 160 dB re 1 μ Pa), $< 1\%$ of the simulated population was exposed to levels associated with injury. Nevertheless, many simulated bowhead whales received sound levels considerably above ambient throughout their migration. Our method enables estimates of the aggregated level of sound to which populations are exposed over extensive areas and time periods.

KEY WORDS: Arctic · Cumulative effects · Harassment · Incidental take · Marine mammals · Negligible impact

INTRODUCTION

Anthropogenic underwater sounds from activities such as seismic exploration for oil and gas, commercial shipping, and operation of offshore oil platforms and wind turbines can reduce the condition and fitness of marine mammals. In some cases these sounds may cause temporary or permanent loss of hearing or shifts in hearing thresholds (reviewed in Southall et al. 2007). Exposure to chronic and impulsive underwater sounds also can increase levels of stress hormones (Rolland et al. 2012) and change behaviors associated with migration (McDonald et al. 2012), acoustic communication (Holt et al. 2011, Blackwell et al. 2013), diving (Costa et al. 2003, Tyack et al. 2011), and foraging (Goldbogen et al. 2013). Anthropogenic underwater sounds may decrease the ability of marine mammals to recognize either conspecific sounds or sounds used to perceive the environment, find prey, or avoid predators (Ellison et al. 1987a, George et al. 1989, Clark et al. 2009). Underwater sounds associated with certain sonar systems have been implicated as the cause of strandings of multiple species of beaked whales (D'Amico et al. 2009). Additionally, changes in the distribution or behavior of marine mammals in response to anthropogenic sound may reduce the availability of those animals to subsistence hunters in Alaska. The US Marine Mammal Protection Act (MMPA) (16 USC 1361-1423) prohibits activities that will have an 'unmitigable adverse impact' on subsistence uses 'by any Indian, Aleut, or Eskimo who resides in Alaska and who dwells on the coast of the North Pacific Ocean or the Arctic Ocean.'

The anthropogenic underwater sounds detectable at any one location often emanate from multiple sources. However, assessments of the potential responses of marine mammals to anthropogenic sound typically evaluate responses to a single, relatively loud source, such as sonar or an airgun array from a particular ship. Attempts to assess responses to aggregated sounds, or responses to the aggregated biological effects of sound from multiple sources, have seldom progressed beyond a statement of need and, occasionally, qualitative descriptions. For example, the German government examined the potential effects of seismic research on Antarctic marine ecosystems, acknowledging the challenges of rigorously assessing the potential environmental effects of sound (Siebert et al. 2012). Similarly, Moore et al. (2012) summarized major anthropogenic sources of sound in the Arctic Ocean and potential mechanisms by which marine mammals and their prey may respond physiologically or behaviorally. Moore et al. (2012) proposed

that aggregated sound fields from human activities and natural phenomena be mapped and compared to spatial data on the distribution and abundance of marine mammals at different life stages. Another effort supported by federal agencies in the U.S. mapped distributions of cetaceans and anthropogenic underwater sound throughout the country's exclusive economic zone, including the Alaskan Arctic (NOAA 2012). Although these maps illustrated that some species of marine mammals occupy regions with substantial levels of underwater anthropogenic sound, the effort did not document contributions to the acoustic environment from individual sources of sound, the distribution of individuals and populations as the acoustic environment changed over time and space, or the levels of sound to which animals were exposed as their acoustic environment changed.

Biological responses to aggregated sound exposure are highly relevant to decision-making about diverse human activities in marine ecosystems. The National Marine Fisheries Service (NMFS) of the US National Oceanic and Atmospheric Administration (NOAA), which is charged with implementing the MMPA as it applies to pinnipeds and cetaceans, currently considers a received root mean square (rms; essentially an average over the duration of a pulse or over a short, fixed period for non-pulse sounds) sound pressure level (SPL; in dB re 1 μ Pa) ≥ 160 dB for impulsive sounds and ≥ 120 dB for continuous sounds as thresholds beyond which behavioral patterns of pinnipeds and cetaceans may be disturbed by human activities. A non-technical definition of the SPL is the instantaneous level of sound received. NOAA typically considers a received rms SPL ≥ 180 dB as the threshold above which cetaceans may be injured physically. Southall et al. (2007) suggested that these rms SPL-based injury thresholds are not supported by empirical data on the physical effects of sound. They further suggested that cetaceans with sensitivity to low-frequency sounds (e.g. baleen whales) may receive an auditory injury from impulsive sounds with peak SPL ≥ 230 dB or cumulative sound exposure levels (SEL; in dB re 1 μ Pa² s, accumulated over 24 h) > 198 dB, and by non-pulse sounds with cumulative SEL ≥ 215 dB (Southall et al. 2007). The SEL is an integrative measure of the received SPL over a given period of time, e.g. 24 h.

The nature and methods of calculating SPL versus cumulative SEL are fundamentally different. However, in most regulatory assessments in the U.S., both SPL and cumulative SEL are treated as static thresholds for the onset of injury or behavioral effects. Data are rarely sufficient to consider dynamic interactions

among an animal, the sound field to which it is exposed at a particular time and place, and the biology of the animal (e.g. its identity, age, sex, and condition). Recent workshops and forums have considered potential methods for estimating the cumulative effects of anthropogenic underwater sound that are more informative than simple thresholds of take (i.e. a single specified level of sound that indicates potential injury or harassment; NOAA 2012). The individual-based Acoustic Integration Model[®] (AIM) (Frankel et al. 2002, Independent System for Peer Review 2006) that we used in this study can evaluate complex cumulative SELs. To date, the AIM has been applied to inform diverse major environmental studies and environmental impact statements that examined potential responses of marine mammals to multiple sources of sound, such as shipping, airguns, and sonar (e.g. Department of the Navy 2001, Gabriele et al. 2010, NSF-USGS 2011).

Clark et al. (2009) examined potential effects on endangered baleen whales of the aggregated sound field generated by one or more commercial ships. They used the aggregated sound field to measure the potential effect of a reduction in communication space, referred to as communication masking. Hatch et al. (2012) calculated communication-masking metrics to evaluate the aggregated effects of sound from 117 commercial ships on a modeled population of 89 North Atlantic right whales *Eubalaena glacialis* off Boston, Massachusetts, over 1 mo. Clark et al. (2009) calculated measures of communication masking for individual modeled whales, whereas Hatch et al. (2012) calculated masking at the level of the group of simulated whales. We built on those previous methods of assessment by quantifying the aggregated contributions to exposure from multiple sources of underwater sound, both continuous and impulsive, associated with the offshore oil and gas industry.

The US National Environmental Policy Act (NEPA) is generally interpreted to require an analysis of 'the impact on the environment which results from the incremental impact of the action when added to other past, present, and reasonably foreseeable future actions' (40 CFR §§ 1500–1508; Council on Environmental Quality 1997). An analysis of the cumulative effects of human activities ideally would encompass the effects not only of sound, but of all potential stressors. Nevertheless, developing a quantitative process to estimate exposure to multiple sound sources could enhance cumulative-effects analyses.

Our objective was to develop a quantitative method for estimating the sound exposure level of marine mammals to multiple sources of continuous and

impulsive anthropogenic sound over weeks to months, and, as a proof of concept, to apply this method to a simulated population of marine mammals. Some species of marine mammals, such as gray whales, have been observed avoiding sound sources in their migratory path (Malme et al. 1984, Buck & Tyack 2000, Ellison et al. 2012). Therefore, we aimed to conduct an initial exploration of how changes in movement paths of modeled animals programmed to avoid certain received levels of sound (i.e. aversion) might affect both sound exposure and travel distance.

METHODS

To develop the modeling method and our simulations, we focused on bowhead whales *Balaena mysticetus* in the Alaskan Beaufort Sea. Bowhead whales have cultural meaning for Alaska natives, and subsistence hunting of the species is protected under the MMPA. The species is also listed as endangered under the US Endangered Species Act. Each autumn, bowhead whales move westward from feeding areas in the Canadian Beaufort Sea through the Alaskan Beaufort Sea and into the Chukchi Sea. This region increasingly is a focus of exploration for and production of oil and gas. We bounded our model in space from about 144–152°W and in time from 1 September through 23 October 2008. These extents encompass most of the autumn migration through the region. We selected 2008 because it was a year of relatively high intensity of seismic exploration in the region (Roth et al. 2012).

As detailed below, we first modeled the sound field generated by each anthropogenic sound source during a defined period of time. Second, we simulated the movements of migrating whales through each of these sound fields. We refer to the simulated whales as animats. Each animat functions as an acoustic dosimeter that records its exposure at 30 s intervals. In 1 run of the simulation model, we assumed that the animats would not or could not change their movement path to avoid a given received level of sound (i.e. would not or could not avert). In another model run, we allowed the animats to avert from certain received sound levels. Third, we recorded the modeled SPL rms history and then calculated the resulting cumulative SEL for each animat from each source. Fourth, we estimated the aggregated exposure from all sources for each modeled whale. Individual animats migrated through the sound fields in 3–4 d. The entire modeled population moved through the area over 47 d.

Acoustic modeling

We compiled data on sounds from oil and gas production, seismic exploration, and vessel traffic from publicly available sources. These sources included a multi-year comprehensive report (Funk et al. 2010), applications for incidental harassment authorizations under the MMPA, and post-exploration season monitoring reports (www.nmfs.noaa.gov/pr/permits/incidental.htm#applications). On the basis of these empirical data, we modeled the sound fields generated by the Northstar and Oooguruk production islands, 1 offshore and 1 nearshore vessel towing a barge ('barge tow'), and 2 offshore and 3 nearshore seismic survey operations (Fig. 1). The production islands and barge

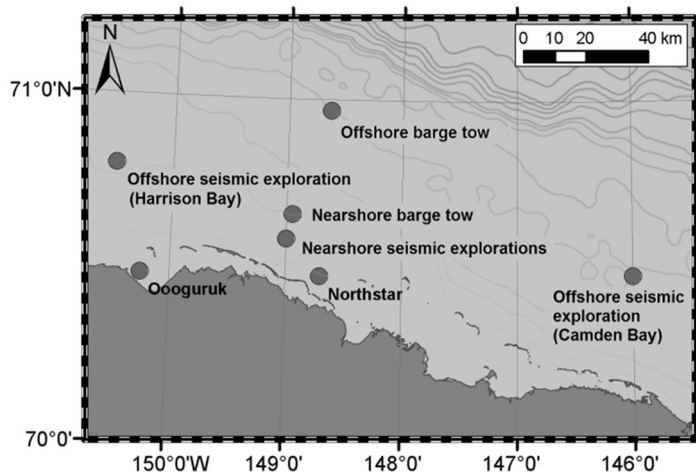


Fig. 1. Locations of sound-producing operations included in the model of the aggregated anthropogenic sound field in the Alaskan Beaufort Sea. Oooguruk and Northstar are production islands. The darker grey toward the south indicates mainland Alaska. Contour lines are in 10 m increments

tows produced sound throughout the assessment period, whereas seismic exploration was episodic and varied among operations (Table 1).

Our intent was not to recreate the precise temporal, spectral, and spatial properties of the anthropogenic sound field but to generate a plausible aggregated sound field. We kept the acoustic model relatively simple and tractable by selecting sources of sound that represented different major classes of activities, including those with the strongest outputs of sound. We further prioritized simplicity and tractability of the acoustic model by assuming that all sound sources were stationary. These simplifications precluded a quantitative estimate or validation of true sound exposure in the region during the simulated period. However, they allowed for realistic, transferable modeling of a dynamic, anthropogenic sound field in which the moving animals were exposed to received sound levels greater than nominal ambient sound, which we considered to be a biologically meaningful scenario.

We estimated 3 acoustic metrics: rms SPL, per-impulse SEL (in dB re $1 \mu\text{Pa}^2 \text{ s}$), and cumulative SEL (see Southall et al. 2007, their Appendix A). We based cumulative SEL on the aggregated exposure of each animal to all sound sources for the duration of its exposure period (e.g. Madsen 2005, Madsen et al. 2006). Estimates of aggregated exposure are dominated by the source producing the highest received level, which typically occurs when the receiver is at or near its closest point of approach to the sound source.

Southall et al. (2007) proposed new criteria for sound exposure that were based on M-weighted SEL (i.e. sound exposure levels adjusted to the hearing curve of a hypothetical marine mammal). We instead used unweighted values of SEL (i.e. measures of sound not adjusted to a hearing curve) to

Table 1. Major anthropogenic sources of sound in the Alaskan Beaufort Sea during the autumn bowhead whale (*Balaena mysticetus*) migration in 2008 (1 September–23 October). $1 \text{ inch}^3 \approx 16.39 \text{ cm}^3$

Activity	Dates of activity	Characterization of activity
Northstar production island	1 Sep–23 Oct	Routine production with vessel support
Construction of Oooguruk production island	1 Sep–23 Oct	Construction with vessel support
Offshore barge tow	1 Sep–23 Oct	Generic barge tow typical of Beaufort Sea
Nearshore barge tow	1 Sep–23 Oct	Generic barge tow typical of Beaufort Sea
Offshore seismic exploration in Harrison Bay	3–12 Sep and 1–9 Oct	~3000 inch^3 airgun array
Offshore seismic exploration in Camden Bay	14–29 Sep	~3000 inch^3 airgun array
Nearshore seismic exploration 1	3–5 and 20–28 Sep	~900 inch^3 airgun array
Nearshore seismic exploration 2	1–28 Sep	~900 inch^3 airgun array
Nearshore seismic exploration 3	6–13 Sep	~450 inch^3 airgun array

calculate cumulative exposures. Use of unweighted values tends to result in higher estimated levels of exposure than use of weighted values, and therefore is likely to be considered by regulators as a more conservative method. On the basis of data on hearing in cetaceans, the most recent draft guidelines for low-frequency weighting (NOAA 2013) proposed modifications to the M-weighting curve, but these modifications have not been validated for general application.

We assumed that during a given 24 h period, each of the airgun arrays used in seismic surveys could be either inactive (silent) or constantly active and firing at 10 s intervals with a 100 ms impulse duration. For these impulsive sources, we modeled the per-impulse SEL and assumed a nominal impulse duration across the sound field of 100 ms (Richardson et al. 1995a) regardless of the distance from the source, with no acoustic energy in the periods between impulses. We calculated rms SPL values as $SPL = SEL - 10 \log_{10}(\text{impulse duration in seconds})$; for a 100 ms impulse, the SPL is $SEL + 10$ dB. Acoustic measurements taken in the highly reverberant shallow waters of the North Slope of Alaska (Guerra et al. 2011) do not support the assumption that pulse dispersion and multipath propagation are functions of range from the source. However, the net result of the simplified relation we used to calculate SPL is an overestimation of rms SPL as distance from the seismic source increases.

We estimated the source levels (per-impulse SEL in 1/3-octave bands) for the airgun arrays used in the seismic surveys (Table 1) on the basis of a full-waveform airgun array source signature model (AASM; MacGillivray 2006). This model is based on the oscillation and radiation of air bubbles released as airguns discharge (Ziolkowski 1970). The model accounts for pressure interaction between bubbles, effects of reflection from the water's surface on pressure, heat transfer from bubbles to the surrounding water, and buoyancy of the bubbles.

We estimated source levels for barge tows and production islands on the basis of underwater measurements from these and comparable activities (e.g. Blackwell et al. 2004). For the barge tow, we used a single, ocean-bound tug in cruise transit mode as the representative source; we obtained its 1/3-octave band source levels from an extensive collection of sound source measurements (Hannay et al. 2004). We assumed that support vessels operating in proximity of production islands would generate the highest levels of sound, and we obtained representative 1/3-octave band source levels for a shallow-

draft utility and crew-transfer vessel transiting at low speed from Hannay et al. (2004). The broadband rms source level (10–1000 Hz) of this vessel was set at 172 dB re 1 μPa @ 1 m (Hannay et al. 2004).

We obtained basin-level bathymetry data from the Geographic Information Network of Alaska (GINA) online database (Lindquist et al. 2004). We derived sound velocity profile curves from the Generalized Digital Environmental Model Variable Resolution (GDEM-V) database published by the US Naval Oceanographic Office (Teague et al. 1990). Given the seabed composition of the regions where the sources were located, we used 2 sets of geo-acoustic parameters in the models: one for Harrison Bay and adjacent regions to the east, where some of the sound-producing activities were located, and one for Camden Bay (Fig. 1). We derived the parameters from the geological properties of the seafloor strata with the method of Hamilton (1980).

We used the Marine Operations Noise Model (MONM) (Hannay & Racca 2005) to estimate the sound field produced by individual sources. This model computes acoustic-transmission loss in individual frequency bands and solves the acoustic wave equation as a parabolic function as implemented in the US Naval Research Laboratory's Range-dependent Acoustic Model (RAM) (Collins 1993). The MONM uses a complex density approximation (Zhang & Tindle 1995) to account for reductions in water-borne sound caused by transfer of energy from the water column to the seabed in the form of shear waves. We estimated 3-dimensional acoustic propagation over an extensive region by modeling transmission loss in 1/3-octave bands from 10 to 1000 Hz center frequency along a fan of 2-dimensional radial grids at uniform angular increments, each grid evenly spaced in depth and range from the source. Most of the sound intensity for the sources modeled is concentrated in this frequency band. We applied the calculated frequency-dependent transmission loss to the estimated source levels in corresponding bands. For each sound source, we computed the resulting broadband sound levels (10–1000 Hz) on a uniform grid of points over the modeled geographic region of interest. For each of the 7 weeks during the modeled time period, we computed the aggregated received sound level at each animat by summing the received sound energy from each source. Different sources operated each week, and their locations varied (Fig. 2).

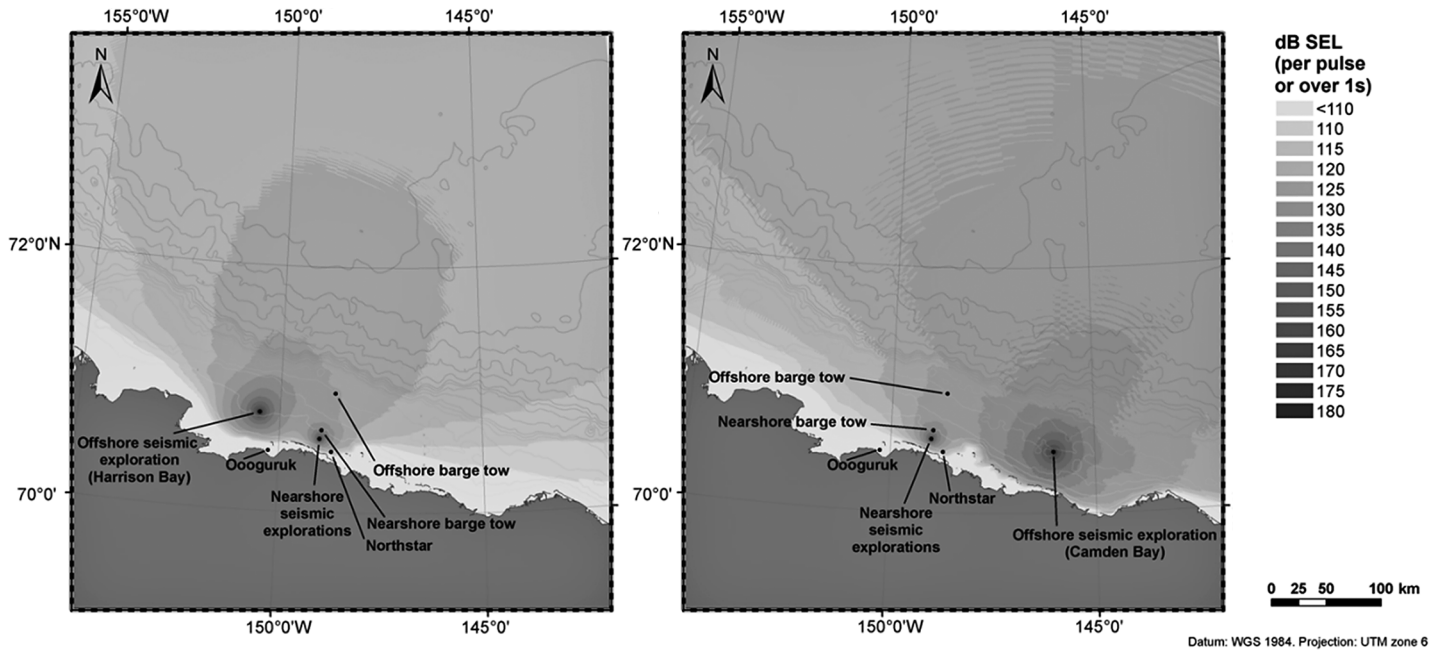


Fig. 2. Two examples of the aggregated acoustic environment (maximum level of sound over the full water column) for periods during which 5 sound sources were operating in the Alaskan Beaufort Sea (left panel, 3–5 September; right panel, 20–28 September). SEL: sound exposure level. For further details see Fig. 1

Modeling movements and exposures

We used the Acoustic Integration Model® (Frankel et al. 2002), a Monte Carlo-based statistical model, to simulate movements of bowhead whales through each of the aggregated sound fields and to calculate each modeled whale’s exposure over time. The AIM is based in part on a model of whale movement and tracking developed for acoustic censuses of bowhead whales (Ellison et al. 1987b).

The parameters in our model of bowhead whale movement and exposure were the local population size for each week during the modeled time period, percentage of the population in the Alaskan Beaufort Sea throughout the migration period, distance of individual animats from the coastline, direction and speed of motion, dive time and dive depth, surface time, water depth, sound exposure level, and aversion or no aversion to sound. Empirical data are limited, but we felt they were sufficient to estimate these parameters as a proof of concept of the method. We did not aim to assess the goodness-of-fit between the outputs of our simulations and field observations.

We extrapolated the size of the bowhead whale population migrating westward in 2008 (14 400) from the estimated population size (10 470) and population growth rate (3.4 %) in 2001 (George et al. 2004). On the basis of traditional and scientific observations in the local region, we estimated that 80 % of the popu-

lation migrated through the Alaskan Beaufort Sea between 1 September and 12 October. Therefore, the population size for our simulations was 11 566. For each modeled week, we designated a percentage of the population as migrating (Table 2).

Aerial survey data suggested that the density distribution of bowhead whales is a function of water depth and distance offshore (Clarke & Ferguson 2010), with approximately 95 % of the animals in the nearshore region (depths of 20–200 m) and 5 % in the offshore region (depths of 50–2000 m). Therefore, we modeled bowhead whales in the nearshore and off-

Table 2. Dates within each modeled period, number of animats (simulated whales) traveling nearshore (20–200 m water depth) or offshore (50–2000 m water depth) during each period, and percentage of each group (in parentheses) migrating during each period. Modeled animats were placed throughout the 144–152° W area. New animats entered the simulation from the east as the westernmost animats swam out of the modeled area to the west

Modeled period	Nearshore	Offshore
1–7 Sep	2198 (21)	119 (20)
8–13 Sep	2352 (21)	126 (21)
14–19 Sep	2352 (21)	126 (22)
20–26 Sep	1959 (18)	102 (17)
27 Sep–2 Oct	1098 (10)	56 (10)
3–9 Oct	671 (6)	36 (6)
10–16 Oct	350 (3)	21 (4)

shore regions separately, allowing some overlap of the 2 regions at depths of 50–200 m.

We populated each simulation with westward-migrating animats (i.e. animats east of 144°W that moved into the modeled area over time) at the beginning of each 1 wk period. A week is long enough for a westward migrating bowhead whale to traverse the modeled portion of the Alaskan Beaufort Sea.

We assumed that animats began their migration on a generally westerly heading. We allowed each animat's heading to change randomly by $\leq 10^\circ$ every 5 min, always maintaining an attraction to the west, but confining the travel path to the allocated depth contours for the nearshore and offshore distributions (i.e. following the contour of the coast). We allowed the speed of each animat to vary from 4 to 6 km h⁻¹ on the basis of limited data from satellite tags deployed by Mate et al. (2000). The tags recorded a mean speed of 3.8 km h⁻¹ among 8 bowhead whales that were either feeding or migrating, with individual mean speeds of 1.1–5.8 km h⁻¹. One of the tagged bowhead whales (DZ-1) was tracked along its full migration route from Canada to Russia. The mean speed of this whale was 5.0 km h⁻¹. Because calculations of distances between locations assumed straight-line travel, and therefore represent the minimum distance and speed, the speeds reported by Mate et al. (2000) probably underestimated true speeds.

On the basis of satellite tag data, we modeled both the duration and depths of 2 types of dives: relatively short, shallow dives and relatively long, deep dives. Bowhead whales tagged by Krutzikowsky & Mate (2000) spent 60% of their time diving at depths <16 m, 33% of their time at depths from 17 to 96 m, and <3% of their time at depths >96 m, with a maximum dive depth of 352 m (water depth permitting); they spent a mean of 5.5% of their time at or near the surface. We assigned 60% of the nearshore and offshore animals to the short, shallow dive type, with 5–16 m dive depths in both regions, and 40% to the long, deep dive type. Because water depth decreases as distance from the shoreline decreases, we assumed that depths of the longer, deeper dives of nearshore bowhead whales ranged from 17 to 50 m and those of offshore bowhead whales ranged from 17 to 96 m. The satellite tag data also showed that the durations of dives >1 min were 2.6–30.4 min, independent of dive depth, with individual means from 6.9 to 14.1 min and an ensemble mean of 10.4 min. However, when ice cover was <90%, the majority of dives were ≤ 13 min, with a relatively small percentage of longer dives (Fig. 7a in Krutzikowsky & Mate

2000). Because duration and dive depth were independent, we selected durations of 3–10 min for the shorter dives and 8–13 min for the longer dives. These dive durations are consistent with those reported earlier (Dorsey et al. 1989, Richardson et al. 1995b).

As described in the previous subsection, we modeled the exposure of the animats to sound during 7 wk (Table 2), corresponding approximately to the onset and completion of the population's migration period (Table 1). We used the AIM to estimate the instantaneous level of sound (continuous or impulse rms SPL, depending on the source type) received by the animat from each source at 30 s intervals as it migrated westward through the Alaskan Beaufort Sea. In essence, the model created an aggregated acoustic dosimeter record at 30 s increments for each animat.

To illustrate the exposure-modeling process, we calculated the exposure over a 78 h period (the approximate duration of one animal's migration through the Alaskan Beaufort Sea) for a single inshore migrating animat that passed close to both offshore and nearshore seismic operations, the Northstar production island, and one of the barge tows. We used these results to calculate both the maximum SPL and the cumulative SEL for each animat. We compared our modeled estimates of the aggregated received sound levels to the potential range of sound levels to which a marine mammal in this region might be exposed in the absence of anthropogenic sound. To estimate the latter range, we used empirical measurements of non-impulsive ambient sound levels for September 2008 at a location on the shelf slope near the border between the Chukchi and Beaufort Seas (Roth et al. 2012). The Roth et al. (2012) data present broadband (10–1000 Hz) values of 96 dB re 1 μPa^2 for the 10th percentile and 117 dB re 1 μPa^2 for the 90th percentile; these values are higher than those provided in earlier assessments for this region (Urick 1983, Richardson 1995a). Because Roth et al. (2012) monitored sound at a depth of 235 m in a location exposed to sound propagated from full ocean depth, their reported ambient levels potentially could be higher than in the shallow water of the inshore migrating animats.

We ran 2 separate sets of AIM simulations, one in which the whales in the simulated population could not change their movement path to avoid certain sound levels (i.e. avert) and one in which they could avert. In the simulations that did not allow aversion, animats were not programmed to respond to the

sound exposure level, and as a result their migration paths were unaffected by the modeled sound sources. In the second set of simulations, animats were programmed to avert from a sound source as a function of their acoustic environment. The 4 parameters in the aversion function were the SPL threshold at which an animat was assumed to respond, the probability that the animat would respond, the aversion response angle, and the rate at which the animat subsequently evaluated the sound field (SPL) (Table 3). Our selection of aversion parameters for bowhead whales was intended to represent a reasonably expected behavioral response to sound exposure, not to predict response behavior explicitly.

Of the 4 parameters, probability of aversion had the greatest effect on the behavioral response, and aversion would be the most tractable to validate in the field. Although we set the probability of aversion at $\text{SPL} \geq 160$ dB re 1 μPa at 0.60, the sound level at the animat was reevaluated every 5 min. Accordingly, an animat that continued toward the sound source and did not avert at the first level eventually would be

exposed to $\text{SPL} \geq 160$ dB re 1 μPa and again would have the opportunity to avert. We acknowledge that bowhead whales may respond to received impulsive and continuous sound at $\text{SPL} < 120$ dB re 1 μPa (Richardson et al. 1995a, Davies 1997, Richardson 1999, McDonald et al. 2012). However, we limited aversion responses to received $\text{SPL} \geq 160$ dB re 1 μPa to allow the animats to move through the sound field and to run the model more efficiently. Again, our intent was not to yield outputs that fit field observations, but to develop a transferable method that can be parameterized with data on different activities, locations, and species.

We summarized the exposure for all animats in the modeled populations. We used the single maximum received SPL value for each animat, regardless of source type, as data for both of the aversion options. We binned values in 5 dB increments from 120 to 220 dB re 1 μPa . We calculated the corresponding distribution of cumulative SEL values, computed for each animat by summing the sound exposures from all active sources over all time steps in a run.

Table 3. Aversion parameters for animats (simulated whales) used in models of movement and sound exposure. rms: root mean square; SPL: sound pressure level

Received SPL threshold (rms SPL, dB re 1 μPa)	Probability of aversion	Aversion angle ($^{\circ}$)	Interval between SPL evaluations (s)
≥ 160	0.60	10	300
≥ 170	0.85	20	60
≥ 180	0.99	30	30

RESULTS

Example of individual-level exposure

A single inshore migrating animat (not programmed to change its movements in response to a given sound level) selected from the migrating population illustrates exposure to multiple sources as the animat traveled close to both offshore and nearshore seismic operations, the Northstar production island, and one of the barge tows (Fig. 3) over 78 h. In this case, the large offshore seismic array was the dominant contributor to received sound levels throughout the animat's entire passage, except from about 32–41 h after the simulated migration began (Fig. 4). During those 9 h, the animat passed in close proximity to the nearshore seismic operation, which then dominated the received sound level. The animat's closest point of approach to the offshore seismic source was about 25 km, and occurred about 5 h after the simulation began. The SPL at that point was about 150 dB re 1 μPa . The received SPLs from individual pulses from the offshore seismic source were > 120 dB re 1 μPa throughout the animat's migration. The animat reached its closest range of approach to the nearshore seismic operation—a few kilometers—approximately 36 h after its simulated migration began, and the received SPL of impulses from the nearshore seismic operation peaked at just above

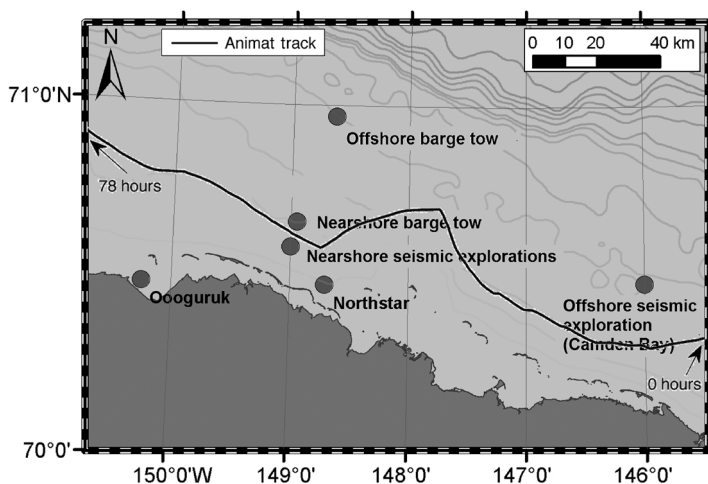


Fig. 3. Example of an animat (simulated whale) track over the 78 h period of its migration through the Alaskan Beaufort Sea (without aversion). For further details see Fig. 1

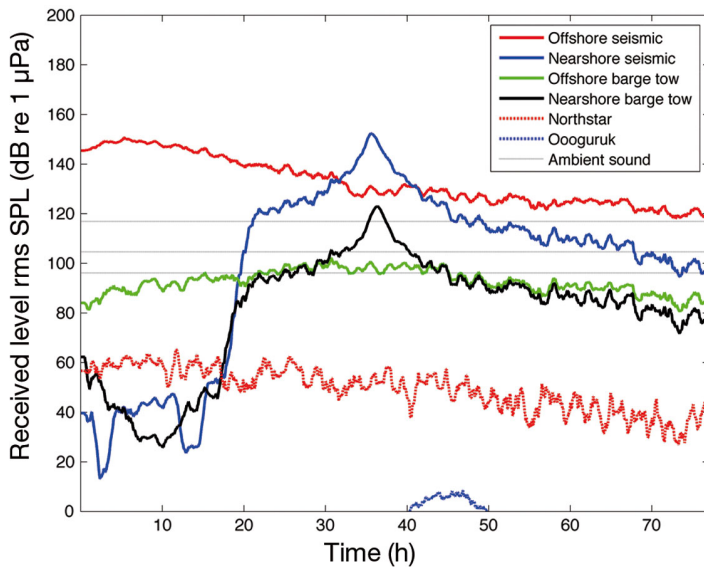


Fig. 4. Instantaneous levels of sound (broadband continuous or impulse root mean square sound pressure level [rms SPL]) received from each source at 30 s intervals by the animat (simulated whale) represented in Fig. 3. Gray horizontal lines denote the median and upper (90%) and lower (10%) bounds of broadband (10–1000 Hz) ambient sound (Roth et al. 2012)

150 dB re 1 μPa . The maximum received SPL levels were about 45 dB higher than the mean ambient band level (10–1000 Hz) estimated by Roth et al. (2012).

The next-highest received SPL values occurred during the animat's closest approach to the nearshore barge tow as the animat passed between the barge tow and the nearshore seismic source. At this point, the received sound level at the animat reached a maximum SPL for both the nearshore seismic operation (150 dB re 1 μPa) and the nearshore barge tow (125 dB re 1 μPa), the latter at a closest point of approach of 2.4 km. Although the continuous sound exposure from the barge tow for this animat never exceeded the impulse SPL from either of the seismic sources, the level of sound from the barge tow was the highest that reached the animat in the short intervals between airgun impulses. During the 78th hour of this animat's transit, it was exposed to sound levels from some combination of the seismic sources above the 90th percentile ambient in the 10–1000 Hz band (Roth et al. 2012). With respect to the SPL levels from the continuous sources, only those from the 2 barge tows exceeded the 10th percentile ambient levels, and the levels in the same band from the production activities at

Northstar and Oooguruk were well below the 10th percentile ambient. For 83% of the 78 h migration period, the animat received sound levels that were >20 dB re 1 μPa greater than the estimated mean ambient in the 10–1000 Hz band (Roth et al. 2012). These insights would not be evident from traditional reports of a single peak measurement.

Population-level exposure

The proportion of the simulated population that was exposed to SPL >165 dB re 1 μPa decreased markedly when the animats were allowed to avert (Fig. 5a). However, some animats in the population that could avert were still exposed to SPL >165 dB re 1 μPa , probably because they had a finite probability of not averting at each time step and because new sound sources were activated during the migration. The cumulative SEL was considerably lower for a population of animats that could avert than for a population of animats that could not avert (Fig. 5).

We calculated the percentage of animats in each modeled population that were exposed to a range of maximum SPLs (Fig. 6). For example, about 6% of the animats in the population that could not avert

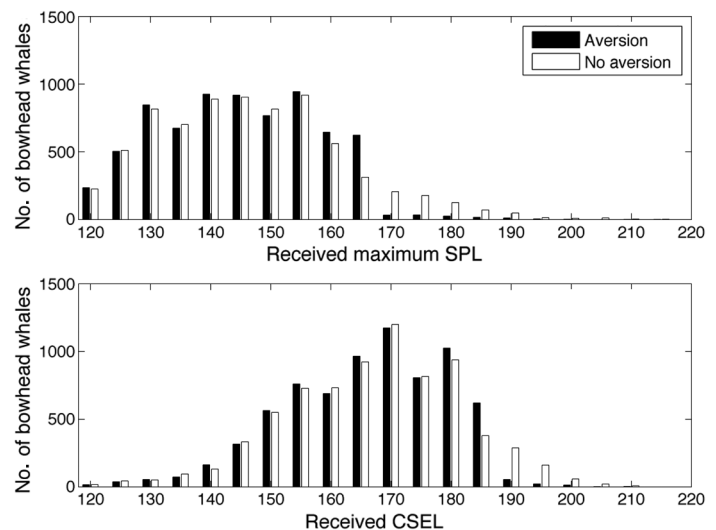


Fig. 5. Aggregated maximum received sound pressure levels (rms SPL; in dB re 1 μPa) (upper panel) and cumulative sound exposure levels (CSEL; in dB re 1 $\mu\text{Pa}^2\text{ s}$) (lower panel) for all animats (simulated whales) during their autumn migration through the Alaskan Beaufort Sea. The CSEL for each animat is the sum of its exposure to sound from all sources over all time steps

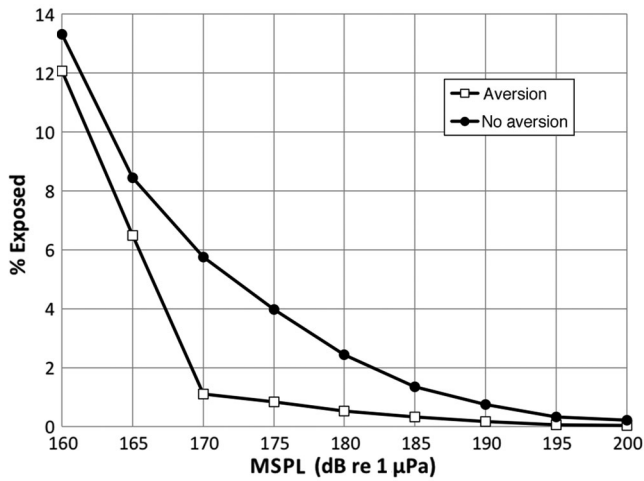


Fig. 6. Percentage of animals (simulated whales) in the population that were exposed to maximum sound pressure levels (MSPL) equal to or greater than given values as a function of whether they averted from those levels

were exposed to SPL ≥ 170 dB re 1 μ Pa. By contrast, about 1% of the animals in the population that could avert were exposed to SPL ≥ 170 dB re 1 μ Pa.

In our models, aversion from any sound source had little effect on travel distance. For example, between 145.5 and 146.5°W, where a seismic operation was positioned, the average distance traveled by animals in the population that could not avert from rms sound levels of ≥ 160 dB re 1 μ Pa was 41 ± 14 km (mean \pm SD). The average distance traveled by animals in the population that could avert was 46 ± 19 km. When we assessed sound exposure along the full migration path (Fig. 1), we found the difference in mean total distance swum by either population was < 1 km. This result is consistent with the fact that the 160 dB SPL threshold for the onset of aversion occurred at a nominal range of 10 km from the largest airgun array. Each animal encountered a large array once during the 78 h mean duration of passage, and only animals that passed within 10 km of an active large array had the potential to avert. Overall, approximately 5% of individuals in the population that could avert changed their movement paths.

DISCUSSION

Our approach differed in 2 ways from conventional acoustic impact assessment analyses. First, we estimated both SPL and cumulative SEL for each migrating animal as it moved through a temporal series of representative, static, aggregated sound fields generated by multiple sources, including ves-

sels and seismic airguns. Second, one of our 2 model runs included the potential for an animal to avert from a sound source with a probability proportional to the received SPL. Altering the aversion parameters could result in substantial changes in the modeled results, especially if the SPLs at which animals averted were reduced. As a result, a higher proportion of animals would be exposed to lower levels of sound. To illustrate, if the 160 dB threshold was lowered to 155 dB and the probability of aversion was set to 50%, then approximately 50% of the animals previously exposed to 155–160 dB would be exposed to < 155 dB. The use of aversion as a modeling method emphasizes the need for more-extensive empirical data on avoidance behaviors. For example, additional sensitivity analyses of our stochastic aversion approach might inform prioritization of empirical data collection on bowhead whales' aversion responses. Such data might be similar to those of Thompson et al. (2013), who used detections of foraging-related sounds to demonstrate sound-level proportional aversion by harbor porpoise *Phocoena phocoena* to airgun activity over an extent similar to that of our case study.

Understanding the effects of sound exposure will be improved by analyzing dynamic interactions between animals and sound fields or biological variables that affect how animals respond to exposure (Ellison et al. 2012). On the basis of our results, we emphasize the potential increase in information that can be gained not only by modeling the spatial and spectral variability in the sound field produced by multiple sources, but also by broadening the assessment to explicitly quantify the time-varying dynamics of the interaction between animals and the acoustic environment over ecologically meaningful durations.

Extended periods of chronic exposure to impulsive sound at levels above ambient, but < 160 dB re 1 μ Pa SPL, are not considered in current assessments of regulatory take, although exposures to continuous sounds > 120 dB re 1 μ Pa SPL are considered as regulatory takes. Measures of exposure such as signal-to-noise ratio, sensation level, or perceived sound level are not typically addressed in environmental studies, although they are now recognized as biologically relevant to both behavioral responses and chronic effects of sound (Ellison et al. 2012). Chronic exposure at levels substantially above ambient may inform spatial planning, and is likely to have some effect on bowhead whales and other species of whales, perhaps by masking of acoustic communication signals or navigational cues (Ellison et al. 1987a,

George et al. 1989, Clark & Ellison 2004, Clark et al. 2009, Hatch et al. 2012).

Our model treated impulsive and continuous sounds in the same way—that is, by estimating the maximum level in every 30 s interval. In reality, sounds from seismic airgun arrays are sequential discrete impulses, with a single array producing an impulse at intervals of about 8 to 15 s. Continuous sources, if present and included in the aggregate sound level, maintain elevated sound levels during the interstitial periods between impulses. Furthermore, the timing of the impulse from the large and the small seismic sources may be staggered, so sound is generated between each source's impulses. Propagation of acoustic energy from impulsive sources also raises sound levels between seismic impulses through boundary reverberation and multipath propagation effects (Guerra et al. 2011). Thus, sounds originating from a single source potentially arrive at a receiver via different pathways and at different times. Reverberation from the ocean floor near airgun activity may be substantially above ambient sound levels in extremely shallow nearshore environments such as those we modeled, and reverberation and ambient sound levels may be elevated further when multiple seismic operations are being conducted in the same region. Although increases in sound levels between impulses typically are below current regulatory thresholds, they raise background levels and potentially could hinder the ability of animals to communicate or to acoustically sense their environment.

Biological effects of aversion

The tendency of bowhead whales to avert from areas in which industrial sound sources operate was noted by Richardson et al. (1985, 1995a) and in numerous reports (see Schick & Urban 2000). Although it is logical to assume that animals averting from sound sources swim greater distances than those that do not, the extent to which aversion affects the overall migration distance or duration of migration is unknown. Additionally, the biological effects of increased migration distances are not known. Our analysis was essentially a parametric exploration of the possible movement response of bowhead whales to a given sound level. Our estimated input parameter values of aversion behavior were bounded by 160 and 180 dB re 1 μ Pa SPL, the long-established NOAA assumptions of levels that affect behavior and cause injury. This range of reference values thus is

well-understood by industry, researchers, and regulators. More substantive estimates, particularly for initial aversion that might occur at lower SPLs, will require empirical studies similar to those reported by Thompson et al. (2013).

We also recognize that some, perhaps a majority, of bowhead whales migrating in autumn stop to feed. In that case, aversion could lead to 2 energetic costs, the cost of movement itself and the cost of a reduction in food consumption. The total energetic cost would depend on the spatial and temporal extent of the sound source and the proportion of the population affected. Alternatively, if animals feed in an area with high sound levels without averting, then the probability of biologically harmful levels of exposure might increase.

Transferability and applications

Our approach allowed us to assess the feasibility of simulating the migration of a population of >10 000 bowhead whales over a migration period of >2 mo. We encountered no fundamental or intrinsic challenges to estimating aggregated acoustic exposure over extensive areas and time periods. One could adapt the model to include a greater number and variety of sound sources or apply it to different species, longer durations of simulation, or location- and season-specific life stages such as cow-calf pairs or juveniles. It would be tractable to model sound fields created by reverberation and multipath arrivals. Estimating the loss of acoustic communication space as a result of masking would require knowledge of vocalization rates, frequency band, source level, and the change in background sound spectrum due to both natural changes in ambient sound and the introduction of anthropogenic sources (see Hatch et al. 2012). By including mobile sound sources such as shipping traffic, the animat aversion model could also be adapted for proximity to vessel traffic (Ellison et al. 2012, Pirodda et al. 2015).

Our estimates of the instantaneous sound level to which individuals were exposed over time from multiple sources suggested that the presence of multiple sound sources might not substantially increase the maximum SPL to which most individuals in a given population are exposed, but may affect the duration of exposures above a given level. However, inferences might differ at higher modeled levels of industrial activity, such as a greater number of concurrent seismic surveys. Additionally, the cumulative effects of exposure to multiple sound sources may be more

relevant at the population level on a chronic basis than at the individual level on an acute basis, particularly if the exposure includes increases in other anthropogenic activities, such as commercial shipping and other types of vessel traffic.

Although the model presented here is relatively simple, it can be modified to yield increasingly sophisticated representations of the exposure of individual animals and populations to dynamic combinations of continuous and impulsive sounds, whether anthropogenic or natural, and can be parameterized on the basis of traditional ecological knowledge or research on behaviors or behavioral responses. Accordingly, the method improves understanding of the cumulative sounds to which marine mammals are exposed. When coupled with information on bioenergetics, communication, potential habituation to certain sources of sound, and environmental covariates (e.g. distributions of prey), such analyses could begin to reduce some of the considerable uncertainty as to whether sound exposure changes population viability and whether alternative mitigation strategies may be effective. Quantitative assessments of population-level effects of sound are becoming a higher priority given increases in human access to and levels of activity in the Arctic, coupled with increased scrutiny of potential effects of sound on marine mammals.

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