From a calf’s perspective: humpback whale nursing behavior on two US feeding grounds

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ABSTRACT

Nursing influences growth rate and overall health of mammals; however, the behavior is difficult to study in wild cetaceans because it occurs below the surface and can thus be misidentified from surface observations. Nursing has been observed in humpback whales on the breeding and calving grounds, but the behavior remains unstudied on the feeding grounds. We instrumented three dependent calves (four total deployments) with combined video and 3D-accelerometer data loggers (CATS) on two United States feeding grounds to document nursing events. Two associated mothers were also tagged to determine if behavior diagnostic of nursing was evident in the mother’s movement. Animal-borne video was manually analyzed and the average duration of successful nursing events was 23 s (±7 sd, n = 11). Nursing occurred at depths between 4.1–64.4 m (along the seafloor) and in close temporal proximity to foraging events by the mothers, but could not be predicted solely by relative positions of mother and calf. When combining all calf deployments, successful nursing was documented eleven times; totaling only 0.3% of 21.0 hours of video. During nursing events, calves had higher overall dynamic body acceleration (ODBA) and increased fluke-stroke rate (FSR) compared to non-nursing segments (Mixed effect models, ODBA: F1,107 = 13.57756, p = 0.0004, FSR: F1,107 = 32.31018, p < 0.0001). In contrast, mothers had lower ODBA and reduced FSR during nursing events compared to non-nursing segments. These data provide the first characterization of accelerometer data of humpback whale nursing confirmed by animal-borne video tags and the first analysis of nursing events on feeding grounds. This is an important step in understanding the energetic consequences of lactation while foraging.

INTRODUCTION

Humpback whale (Megaptera novaeangliae) calves exert different energetic demands on lactating females on the breeding and feeding grounds. The winter breeding grounds are
located in non-productive tropical waters where females give birth and nurse a single calf while fasting (Chittleborough, 1958; Clapham & Mayo, 1987). In contrast, the summer feeding grounds are located in very productive high-latitude regions where nursing continues, but females also forage to replenish their depleted energy stores while the calf begins transitioning from complete reliance on its mother for sustenance to becoming fully self-sufficient (Chittleborough, 1958; Clapham & Mayo, 1987).

The duration and process of a calf transitioning from full dependence to complete independence of its mother (“weaning”) is not fully known. It appears that the majority of mother-calf pairs separate after they leave the feeding grounds during the calf’s first year (Baker, Perry & Herman, 1987; Clapham & Mayo, 1987). However, a minority of pairs separate early while still on the feeding grounds (Baraff & Weinrich, 1993; Steiger & Calambokidis, 2000) and a few even remain together for a second feeding season as mother-yearling pairs (Baraff & Weinrich, 1993; Hammond, Mizroch & Donovan, 1990). Along this gradient, a humpback whale calf’s reliance on its mother for nutrients likely gradually decreases during the process of weaning (Oftedal, 1997), but neither the frequency nor the duration of visually verified humpback whale nursing events on the feeding grounds have been documented.

Nursing behavior is challenging to study in cetaceans. Suckling occurs subsurface and milk transfer is hard to verify, let alone quantify (Mann et al., 2000). Historically, our understanding of cetacean lactation has been based on whaling data and surface observations, but technological advancements have enabled underwater documentation of nursing in humpback whales. Studies in the breeding grounds have used free divers and underwater video to document suckling behavior (Zoidis & Lomac-MacNair, 2017), as well as archival suction cup tags to record subsurface movement behavior of mother-calf pairs during periods inferred to be suckling based on surface observations (Videsen et al., 2017).

Our study used CATS (Customized Animal Tracking Solutions) tags to study humpback whale nursing behavior on two United States (US) feeding grounds. CATS tags (Cade et al., 2016; Goldbogen et al., 2017) combine both the aforementioned technologies into one device, allowing for the recording of accelerometer data during visually-confirmed (via video) nursing events, while removing any possible depth limitations or behavioral effects of the presence of free divers. We thus provide the first account of accelerometer data of humpback whale nursing confirmed by video and the first analysis of nursing events on feeding grounds. Specifically, our goals were to (1) report the duration, frequency, and depth at which successful nursing occurred on two US feeding grounds, where it had not previously been described, and (2) describe accelerometer signals of calves and their mothers during visually verified nursing events and non-nursing segments. We expected (1) the duration, frequency, and depth at which nursing occurred on the feeding grounds to differ from what has been described on the breeding grounds; (2) the majority of the time a calf spent in close proximity to its mother’s ventral side to be spent nursing, and (3) due to a mother’s need to balance her time between foraging and offspring care, that nursing could occur in close temporal proximity to foraging.
MATERIALS & METHODS
Tag deployments
All tagging effort was part of two multi-year projects involving the study of baleen whale foraging behavior and movement. One project was the Stellwagen Bank National Marine Sanctuary tagging project (2004-present), which occurred in the southern Gulf of Maine (GOM) off the coast of Massachusetts. The other project was the Stanford NSF IOS 1656691 study, which conducted tagging of baleen whales in Monterey Bay National Marine Sanctuary in 2017-2018. All operations were conducted under National Marine Fisheries Service permits #18059 and #16111 as well as Institutional Animal Care and Use Committee protocols (Stanford University #A3213-01 and Cascadia Research Collective #CRCAUP-6). We instrumented humpback whales with combined video and 3D-accelerometer archival data loggers (CATS) on both the US East Coast (n = 4, one mother-calf pair and two additional calf deployments, one of which was a second deployment on the calf from the mother-calf pair) and West Coast feeding grounds (n = 2, one mother-calf pair).

The CATS tags included 3-axis accelerometers, magnetometers, pressure sensors, and gyroscopes recording data for up to 36 hrs, and one or two light-triggered cameras recording video for up to 8 h (Cade et al., 2016; Goldbogen et al., 2017). Despite the CATS tags’ capacity for video and data collection, the tags are relatively small and lightweight at 680g (tag #20 and 40) and 810 g (tag #30) compared to the estimated size of calves in the feeding ground (likely over 2,000 kg and closer to 5,000 kg) (Geraci & Lounsbury, 2005; Gulland et al., 2008). At most, the tags amount to 0.04% of the calf’s body weight. To attach the non-invasive suction cup tags, a tagging team used seven-meter rigid hull inflatable boats with 6 or 8 m hand-held poles and placed the tag aft of the pectoral flippers but forward of the dorsal fin on the back of each animal. Additionally, researchers recorded the whales’ immediate reaction to tagging. Any indication of aversive behavior by the mother-calf pair resulted in termination of tagging efforts. Similar methods and protocols were used each year, as described by Friedlaender et al. (2009) and Wiley et al. (2011).

Video data analysis
The animal-borne video was manually analyzed to identify nursing events using VLC media player (version 3.0.6; VideoLan Project, Paris, France). All videos used for analysis include the footage from the camera along with corresponding time, depth, speed, jerk, pitch, roll, heading, and timestamp (frame rate of 25 or 30 frames/second and width x length of 2332 p x 1548, 1554 p x 1034 p, or 1920 p x 1182 p, depending on deployment). For the analysis, we only used good quality video, which had sufficient light and clarity to differentiate individuals and body parts beyond the tip of the tagged calf’s rostrum. Only two of the tag deployments contained poor quality video; 31 s of video was excluded from analysis for mn170612-30 and 888s from mn180831-30.

We defined successful nursing events when the tip of a calf’s rostrum contacted a mammary gland of the mother and milk was seen in the water upon release (Fig. 1, Videos S1 and S2). Probable nursing events were noted when the calf’s rostrum made contact with a mammary slit, but milk was not seen at release or the video ended before the calf released. These probable events (n = 4) totaled 67 s and were excluded from the analysis.
since they were inconclusive. Each nursing event commenced when the tip of the calf’s rostrum contacted the mammary gland and ended when the rostrum was no longer in contact. These times were used to determine segments of nursing in the mother’s data by matching the timestamps of successful nursing events of their calves (Fig. 1).

Sensor data analysis
For all tag deployments, accelerometers (dynamic range $\pm 39.2 \text{ m/s}^2$) were sampled at 400 Hz, magnetometers and gyroscopes (dynamic range 1,000 deg/s) were sampled at 50 Hz, and pressure was sampled at 10 Hz. All data were decimated to 10 Hz before further analysis, tag orientation was corrected to whale-frame using periods of known orientation, and animal orientation (pitch, roll, and heading) was calculated using custom-written MATLAB scripts (Cade et al., 2016; Johnson & Tyack, 2003). Animal speed for all deployments was determined using the amplitude of tag vibrations (Cade et al., 2018), and animal depth, pitch, and speed were used to create 3D reconstructions of the animal’s underwater behavior using Trackplot [version 2.3] (Ware et al., 2006).

Dive phases
To determine a kinematic signal for nursing, we first had to compare nursing and non-nursing events within periods of similar diving orientation. Based on data from pressure and orientation sensors from the six tag deployments in this study, we used a visual assessment of dive profiles (breath to breath) to divide dives into three general phases: descending, horizontal, and ascending. We defined the descending phase as the segment starting at the surface and ending once the maximum depth of the dive was reached. The ascending phase was the segment after the whale had already reached the maximum depth of the dive, and had started to decrease its depth towards the surface without increasing the depth by more than 10 m (less than a body length of an adult) at any point along that ascent until terminating at the surface. The horizontal phase was the segment after reaching maximum depth, but prior to the whale starting its ascent toward the surface. All dives had descending and ascending phases, but not all had horizontal phases since a whale can orient towards the surface immediately after reaching maximum depth. Pitch was not used to determine dive phases since the pitch of the calf may have been influenced by the act of nursing. Using these definitions, we categorized each nursing event by the dive phase in which it occurred.

Next, to reduce noise caused by slight differences in tag sensors, we compared nursing to non-nursing events within, not between, each deployment. Fifteen non-nursing segments were randomly selected for each dive phase in which nursing occurred during each deployment to generate baseline non-nursing behavior for that deployment/dive phase. For example, during the mn170612-30 deployment, nursing occurred during descending and horizontal phases; therefore, within the mn170612-30 deployment, fifteen non-nursing segments during descending phases and fifteen non-nursing segments during horizontal phases were randomly selected for analysis. In total, 105 non-nursing segments were chosen for this analysis among the four calf deployments. To accomplish this, we used a random number generator to select random start times during each tag deployment and chose
Figure 1  A multi-media description of two nursing events. (A) The Trackplot pseudotrack of mn170613-20 (mother). The track profile of the tag deployment is located within the gray box above the pseudotrack. The green vertical line marks the location of the whale along the dive profile as well as the depth of the whale on the pseudotrack (57 m). Each horizontal line represents a meter with the larger crosses marking 10 m. The pseudotrack is a 3D-model that, within the program, a viewer can rotate to examine the track of the tagged animal. The dorsal side of the pseudotrack is marked with the blue and light gray chevrons and the ventral side has dark and light gray chevrons. The red and blue triangles represent the upstroke and downstroke of the flukes, respectively. The corresponding dive profile in blue of (B) the calf (mn170613-40) and (C) its mother (mn170613-20) with overall dynamic body acceleration (ODBA) in green and fluke strokes marked as red X’s. We used the number of fluke strokes within each nursing and non-nursing segment divided by the duration of the segment to calculate the fluke-stroke rates (FSR). FSR (fluke-stroke/s) was used in the analysis to account for differences in nursing and non-nursing segment. Nursing events are distinguished with in the yellow boxes with the first event categorized as a descending phase and the second as a horizontal phase. Note the calf’s higher amount of fluke strokes and increased ODBA compared to the mother’s (continued on next page...)

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lower amount of fluke strokes and decreased ODBA during nursing events. (D) The first nursing event (14:04:13–14:04:48) with an image from the calf’s tag during nursing in which you can see the calf’s nares (i) and the mother’s left flipper (ii). The second nursing event (14:05:40–14:06:07) with a series of images (E–H) showing different stages of the nursing process. (E) The calf approaching to nurse with its nares (i) towards the right of the frame and the mother’s left flipper (ii), hemispherical lobe (iii), and mammary slits (iv) visible. (F) The calf is nursing. The calf’s nares (i) and the mother’s left flipper (ii) and hemispherical lobe (iii) are labeled as reference points. (G) The end of nursing with a visible cloud of milk (i) in the water. The calf’s nares (ii) and the mother’s left flipper (iii) are labeled as reference points. (H) The calf moving away from the mother and the milk cloud expanding (i). The calf’s nares (ii) and the mother’s left flipper (iii) are labeled as reference points. (I) The mother starts a series of bottom-side rolls (14:06:32), an indicator of bottom-feeding (Hain et al., 1995; Ware et al., 2006; Ware et al., 2014). The image is an example of bottom-feeding from the mother of mn20180620-40. The calf was swimming upside down which allowed for a good view of the mother’s open mouth with exposed baleen (i) and plumes of sand (ii) where she had disturbed the substrate. The image was rotated 180° for easier viewing with the calf’s back (iii) and its direction of travel towards the mother’s head.

the first 30-second segment of data after that point that matched the dive phase category needed and did not overlap with any other segment already used in the analysis. We selected 15 non-nursing segments per dive phase/individual for the statistical comparison due to the limited duration of the shortest calf tag deployment. We chose the length of 30 s for non-nursing segments to account for the average length of a nursing event (23 s) plus the standard deviation (7 s). The process to select non-nursing segments was repeated for the same analysis among the mothers; however, only nine segments were used for each dive phase due to sample size limitations.

**Kinematic analyses**

To compare accelerometer data from nursing events and non-nursing segments for each tag deployment, segments of interest were extracted from mother and calf tag data using a custom-written script in MATLAB 2014a (MATLAB, 2014). A total of 11 nursing events from calves, three corresponding nursing events from the mothers, and 132 non-nursing segments (105 from calves and 27 from mothers) were subjectively examined using 400 Hz accelerometer data to see if there was a difference between nursing and non-nursing segments. After these initial observations, we determined overall dynamic body acceleration (ODBA) (Gleiss, Wilson & Shepard, 2011; Wilson et al., 2006) and fluke-stroke rate (FSR) (López et al., 2016) warranted further examination. Therefore, for each of the 146 segments we determined the minimum, maximum, and mean depth, FSR, mean speed, and mean ODBA. All angle means were determined using circular statistics. Comparisons were completed in R (RStudio Team, 2016; R Core Team, 2018) using mixed effects models including the interaction of nursing status and dive phase while taking repeated measures from individuals into consideration (Bache & Wickham, 2014; Pinheiro et al., 2019; Wickham, 2016).

**Mother-calf proximity**

In addition to analyzing accelerometer data, we also determined the percentage of time the calf spent in close proximity (within 4 m) to its mother’s ventral side without nursing. The calf with the highest percentage of time spent nursing (mn170613-40) was selected
for an additional detailed analysis whereby all 4.2 h of good quality video were manually examined. For this video analysis, we scored the amount of time the calf’s rostrum was within 4 m (less than an estimated body length of a calf) of the ventral side of its mother, including the ventral area ranging from the mother’s rostrum to fluke and within the span of her flippers. The total duration of its nursing events was subtracted from the time it spent in close proximity to its mother, resulting in an estimate of the proportion of time the calf spent under its mother without nursing.

**Foraging behavior**

Our final goal was to determine the temporal proximity of nursing and foraging events of the mothers. For this analysis, lunges were identified when video captured the whale opening its mouth with concurrent track data typical of a lunge, an increase in acceleration followed by a rapid decrease (Cade et al., 2016; Goldbogen et al., 2017), as well as fish scales being flushed from the mouth after the lunge. Surface feeding was identified by using calves’ tag video to visualize the mother with a full buccal cavity processing mouthfuls of water and prey after lunges at the surface, a behavior in the GOM known as “dragging” (Dybas, 2018). Bottom-feeding was identified through the calf’s tag video when the mother was seen stirring up the bottom and slowly lunging along the seafloor as well as when bottom side-rolls were evident in the mother’s Trackplot record (Hain et al., 1995; Ware et al., 2006; Ware et al., 2014). All tracks were examined for these foraging behaviors (lunges, dragging, and bottom-feeding) around each nursing event, and the shortest period between foraging and nursing was recorded.

**RESULTS**

**Tag deployments**

Three calves were successfully tagged (one of the three calves was tagged twice: mn170612-30 and mn170613-40) on US feeding grounds. Determined by estimates in the field, all of the calves were over 5.5 m in length, had healthy body condition, and were over six months old based on the time of year tagging occurred. Mn170612-30 had a mild reaction to tagging (small tail flick) on the first day and no reaction on the second day (mn170613-40), mn180620-40 had a moderate reaction (hard tail flick), and mn180831-30 showed no reaction to tagging. These reactions fell within the same range found on the breeding grounds when placing DTAGS (a similar type of suction-cup tag) on one- to three- month old humpback whale calves (Stimpert et al., 2012). All nursing events occurred between 1100 and 1800 (local time), which was a result of the time of day that the tags were on the whales (0745-1900). With a mean tag deployment duration of 5.22 ± 1.33 hrs, none of the tags remained on the whales overnight to determine how nursing rates or the mother’s behavior may have varied over a 24-h period.

**Nursing duration, frequency, and depth**

When combined, the four deployments provided 21.0 h of good quality video data to detect nursing behavior from the calf’s perspective between 0745 and 1900 local time (Table 1). We measured 11 successful nursing events, with an average duration of 23 s (±7 sd).
Table 1  Tag deployment time and duration compared to the average duration of nursing events and the percentage of time spent nursing. The exact time a tag was attached to an animal was recorded in local time (EDT and PDT). The duration of “tag deployment with video” represents the amount of time video was recorded; however, the amount of “good quality video” was used in the analysis. The amount of “good quality video” was determined by subtracting the total amount of time the video was too poor in quality to be able to verify nursing (caused by low light levels or high turbidity of the water) from the duration of “tag deployment with video”. The average durations of nursing events were calculated using the duration for each nursing event per individual, while the percentage of time spent nursing is the total percentage of time nursing was recorded during the good quality video for each individual.

| Individual ID | Tag on time (local time) | Duration of tag deployment with video | Amount of good quality video | Number of nursing events | Average duration of nursing events | Total time spent nursing | Percentage of time spent nursing |
|---------------|--------------------------|--------------------------------------|------------------------------|--------------------------|-------------------------------|--------------------------|-------------------------------|
| mn170612-30   | 09:07 EDT                | 6.7h                                 | 6.7h                         | 3                        | 21s                           | 64s                      | 0.26%                         |
| mn170613-40   | 13:17 EDT                | 4.2h                                 | 4.2h                         | 3                        | 28s                           | 83s                      | 0.54%                         |
| mn180620-40   | 11:10 EDT                | 7.2h                                 | 7.2h                         | 4                        | 24s                           | 94s                      | 0.36%                         |
| mn180831-30   | 07:45 PDT                | 3.1h                                 | 2.8h                         | 1                        | 13s                           | 13s                      | 0.13%                         |
| Average of all nursing events |                          |                                       |                              |                           | 23s (±7sd)                    |                          | 0.33%                         |

When all calf deployments were combined, only 0.33% (4.2 min) of the compiled good quality video was spent successfully nursing (Table 1). Nursing events occurred during the descending, ascending, and horizontal phases of the calves’ dives at depths ranging from 4.1 to 64.4 m (along the seafloor) (Table 2). Although four of the nursing events took place in the upper 15 m of the water column, the majority (7 of 11) occurred deeper than 18 m, below the threshold of surface-based observations.

**Kinematic signal**

During nursing segments, calves were more active, with higher mean ODBA and increased FSR across all dive phases compared to non-nursing segments (Mixed effect models, ODBA: F1,107 = 13.57756, p = 0.0004, FSR: F1,107 = 32.31018, p < 0.0001, Fig. 2). In contrast, mothers appeared to be less active with lower ODBA and reduced FSR during nursing segments compared to non-nursing dives (Table 2; Fig. 2); however, statistical analyses were not possible due to the small number of nursing segments across dive phases in mothers. The contrasting activity level is visually apparent in a video captured by mn180831-30 of another mother and calf pair swimming past while appearing to nurse (Video S3). Additionally, a range of speeds were found during nursing events and non-nursing segments regardless of dive phase (Table 2). Overall, the average speed of nursing events was 1.4 m/s compared to the non-nursing segment average speed of 1.6 m/s for both mothers and calves. While these average speeds were the same, calves had a slightly larger range of non-nursing segment speeds (1.0–2.8 m/s) than mothers (1.3–2.4 m/s).

**Mother-calf proximity**

During the analysis of mother-calf proximity, we found that the calf, mn170613-40, spent 14.7% (37.4 min) of the good quality video from its deployment in close proximity to its mother’s ventral side. However, we witnessed successful nursing behavior in only 83 s (3.69%) of that time. Contrary to our expectations, the vast majority of the time the calf was under its mother was not spent nursing.
Table 2  Comparison of speed, overall dynamic body acceleration (ODBA), and fluke-stroke rate (FSR) of all nursing events to the mean speed, ODBA and FSR of non-nursing segments for each individual and dive phase. Associated tag data and dive phase category (ascending, descending, or horizontal) of each nursing event for calves (n = 11) and mothers (n = 3). For calves, the mean speed (m/s), ODBA (m/s²), and FSR (fluke-stroke/s) for non-nursing segments were based on the average kinematic data for 15 randomly selected segments without nursing, but had the same dive phase category as the nursing events during each calf’s deployment. For mothers, the mean speed (m/s), ODBA (m/s²), and FSR (fluke-stroke/s) for non-nursing segments are based on the average kinematic data for nine (due to sample size limitations) randomly selected segments without nursing, but had the same dive phase category as the nursing events during each mother’s deployment.

| Individual ID | Nursing duration (s) | Dive phase | Nursing speed (m/s) | Non-nursing mean speed (m/s) | Nursing ODBA (m/s²) | Non-nursing mean ODBA (m/s²) | Nursing FSR (FS/s) | Non-nursing mean FSR (FS/s) |
|---------------|----------------------|------------|---------------------|-------------------------------|---------------------|-------------------------------|-------------------|----------------------------|
| *Calves*      |                      |            |                     |                               |                     |                               |                   |                            |
| mn180831-30   | 13                   | Ascending  | 1.6                 | 1.9                           | 0.1794              | 0.1289                       | 0.2239            | 0.1816                     |
| mn170612-30   | 15                   | Descending | 1.5                 | 1.9                           | 0.2722              | 0.2761                       | 0.3311            | 0.2680                     |
| mn170613-40   | 35                   | Descending | 1.5                 | 1.3                           | 0.3772              | 0.2209                       | 0.3714            | 0.2370                     |
| mn170613-40   | 20                   | Descending | 1.6                 | 1.3                           | 0.5710              | 0.2209                       | 0.6863            | 0.2370                     |
| mn180620-40   | 27                   | Descending | 1.3                 | 1.5                           | 0.1936              | 0.2048                       | 0.2622            | 0.1927                     |
| mn170612-30   | 31                   | Horizontal | 1.4                 | 1.9                           | 0.2258              | 0.1434                       | 0.2932            | 0.1395                     |
| mn170612-30   | 18                   | Horizontal | 1.4                 | 1.9                           | 0.1938              | 0.1434                       | 0.2825            | 0.1395                     |
| mn170613-40   | 27                   | Horizontal | 1.2                 | 1.2                           | 0.2448              | 0.1497                       | 0.3285            | 0.1395                     |
| mn180620-40   | 22                   | Horizontal | 1.3                 | 1.4                           | 0.1360              | 0.1217                       | 0.2326            | 0.1174                     |
| mn180620-40   | 28                   | Horizontal | 1.2                 | 1.4                           | 0.1499              | 0.1217                       | 0.1079            | 0.1174                     |
| mn180620-40   | 18                   | Horizontal | 1.3                 | 1.4                           | 0.2271              | 0.1217                       | 0.2247            | 0.1174                     |
| *Mothers*     |                      |            |                     |                               |                     |                               |                   |                            |
| mn180831-20   | 13                   | Ascending  | 1.5                 | 2.0                           | 0.0346              | 0.1554                       | 0.0746            | 0.1956                     |
| mn170613-20   | 35                   | Descending | 1.4                 | 1.6                           | 0.0258              | 0.0546                       | 0.0000            | 0.0332                     |
| mn170613-20   | 27                   | Horizontal | 1.2                 | 1.6                           | 0.0311              | 0.2290                       | 0.0000            | 0.1255                     |

**Foraging behavior**

The behavior of the mothers could be determined before and after nine of the 11 nursing events. For all of those nine nursing events, foraging by the mother was detected within 13 s to 12 min of nursing, and was seen in temporal proximity to foraging regardless of the feeding method the mother was using (surface lunge, mid-water lunge, or bottom-feeding). One surface feeding mother in the GOM nursed her calf during the 2.5 min between lunging through two separate bubble-nets. The next day when the same mother was tagged (mn170613-20), she was documented nursing on the descent of a foraging dive and then again along the ocean floor 13 s before she began a series of bottom-side rolls (Fig. 1). The tagged mother in Monterey Bay (mn180831-20) nursed on the dive prior to a foraging dive that contained a single mid-water lunge. In addition to mothers feeding, a tagged calf (mn180831-30) was documented engulfing prey within 10 min of nursing (Video S4).

**DISCUSSION**

This study is the first to successfully capture video of nursing humpback whale calves on the feeding grounds. We obtained valuable video and kinematic data that quantify the
nursing behavior and give new insights into the balance that mothers maintain between foraging and parental care on the feeding grounds.

**Nursing duration, frequency, and depth**

The video and associated accelerometry data here provide the opportunity to compare nursing behavior on the feeding grounds with results from previous studies on the breeding grounds. While on the feeding grounds, calves are learning to forage independently, and mothers are replenishing lost energy stores while still provisioning growing calves. Thus, we would expect nursing to comprise a smaller proportion of time on feeding grounds, but the average duration of successful nursing events on the feeding grounds (23 s ± 7 s, n = 11) was similar to the average duration observed by free divers on breeding grounds (30.6 s ± 16.9 s, n = 5) (Zoidis & Lomac-MacNair, 2017). Additionally, Zoidis & Lomac-MacNair (2017) only documented nursing during 3% of their 199 focal follows that included calves, whereas nursing occurred during all four of our calf tag deployments. This disparity could suggest a difference in frequency of nursing, but more likely the disparity highlights the benefit of animal-borne tags which increase the opportunity to capture infrequent, brief behaviors that can also occur at depths beyond the range of surface observations or free divers.

Another study in the breeding grounds used archival motion-sensing tags to study suckling behavior; however, since actual nursing could not be visually verified, Videsen et al. (2017) inferred suckling dives and active dives based on a 1.5 m/s² median trimmed
and normalized minimum specific acceleration (MSA) threshold value determined by surface observations of peduncle dives (Gero & Whitehead, 2007). Therefore, we were restricted in our ability to compare the kinematic values of our “nursing events” to Videsen et al. (2017) “suckling dives” because “suckling dives” by definition only included periods of low MSA values. Acknowledging the differences in methodologies, Videsen et al. (2017) estimated that “tagged neonate humpback whales are in suckling position, and so potentially suckling, on average 20% of the time”; which is only slightly higher than our results for the combination of the percentage of time the calf spent in close proximity to its mother without nursing (14.16%) plus the time it spent nursing (0.54%). Videsen et al. (2017) deepest “suckling dive” was much shallower (19.2 m) than our deepest nursing event (64.4 m); however, it occurred within the Exmouth Gulf in Western Australia where the water depth is less than 25 m. Our deepest recorded nursing dive occurred along the seafloor; therefore, the depth was likely influenced by the topography of the area and the mother’s need to forage along the substrate, more so than the possible physical constraints of the pair’s ability to nurse at even greater depths.

Both Zoidis & Lomac-MacNair (2017) and Videsen et al. (2017) concluded that in the breeding grounds mother-calf pairs remained mostly stationary while nursing, and Videsen et al. (2017) additionally concluded that nursing only occurred during the “bottom phase” of dives (similar to the term “horizontal phase” used in this study). Contrary to these findings, nursing on the feeding grounds during our study occurred while the pair was moving and during descending, ascending, and horizontal phases of dives. The differences between these breeding ground studies and our study suggest that as the year progresses, mother-calf pairs can adapt their behavior based on the abilities of the calves and the needs of the mothers.

**Kinematic signal**

It appears that calves on the feeding grounds must use increased FSR (in turn producing higher ODBA) to sustain a grip on the mammary gland while the mother continues moving forward at an average speed of 1.4 m/s. Since non-nursing segments with lower FSR included times when the calf was in close proximity to the mother’s ventral side without nursing and when the calf was traveling at faster speeds; the higher FSR while nursing must be due to the calf maintaining contact with its mother, rather than just compensating for maternal forward motion. The mechanical act of suckling would likely affect ODBA more than FSR; however, we don’t know how much push/pull the calf needs to exert to produce milk flow to then determine how that motion contributes to the higher ODBA. Compared to the younger calves on the breeding grounds whose mothers may be resting more often; calves on the feeding grounds have to expend more energy during the act of nursing as well as to remain close to their foraging mothers (Tyson et al., 2012) to take advantage of nursing opportunities between their mothers’ foraging efforts.

**Mother-calf proximity**

Contrary to our expectations, mn170613-40 was not nursing for the vast majority of time it spent under its mother. By remaining close to its mother, mn170613-40 may have been
in a better position to take advantage of quick nursing opportunities while its mother was foraging. Alternatively, the calf may have been staying close to learn how to forage or to benefit from its mother’s speed by staying in her slipstream while traveling. On the breeding grounds, Saloma et al. (2018) found that the second most common position of young calves is under their mother (compared to above or alongside), despite no nursing being observed, and speculated that this might increase the social bond and provide a safe location for the calf to rest since it is not able to fully control its buoyancy early in life. Likely, there is a combination of factors influencing the amount of time a calf spends close to its mother’s ventral side on the feeding grounds; although the amount of time a calf spends away from its mother and how that time and distance changes over a season is equally important to explore to better understand the weaning process.

**Foraging behavior**

Lactation is energetically costly, and a female must not only compensate for current milk production but must also gain enough energy to support herself through another breeding season (Chittleborough, 1958). As we expected, the need for females to forage was evident in the tag data as most nursing events occurred in close proximity (12 min or less) to feeding lunges or bottom-side rolls. Based on the low activity level of mothers during nursing events (low ODBA and FSR), the data suggest that nursing did not occur while females were lunging, but our results show that females could modify their behavior to allow nursing for short durations while diving to depth to feed and during periods between lunges.

When calves were initially tagged their mothers were in a foraging state, which may have created a bias towards the calf spending a shorter than average amount of time in close proximity to its mother while she foraged, or possibly a bias towards a longer amount of time as the calf attempted to learn feeding methods. In our study, tagging occurred between mid-June and early-September, at which point calves would have been at least six months old and likely supplementing nursing with foraging effort (Chittleborough, 1958; Oftedal, 1997; Tyson et al., 2012); as suggested by the tag in Monterey Bay (mn180831-30) that documented successful prey engulfment (Video S4). Interestingly, in the field prior to tagging, mn180831-30 was assumed not to be a calf based on initial behavioral observations (successful surface lunges within an aggregation of feeding humpbacks); however, further documentation of size and behavior confirmed it was a dependent calf. Unfortunately, none of the other tags were in the proper location to capture the opening of the mouth or pleat expansion to distinguish whether a calf was mimicking its mother’s behavior or successfully feeding. Therefore, we cannot confirm if mn180831-30’s foraging success is common for its age. To better understand the transition between nursing and foraging, further research is needed to confirm foraging attempts by calves, determine the actual success of those foraging attempts, and determine how the rate of those successes might improve as the season progresses.

**Tag deployment**

 Targeting aggregations of foraging whales was a top priority for both overarching projects in which the data for this study was collected. This preference created a bias towards
mothers in a foraging behavioral state when the calves were initially tagged. Although the
tagged mothers transitioned between different behavioral states during this study, longer
tag deployments would provide a better understanding of mother-calf pair time-budgets
over a 24-hour period. Since our data are biased towards day-time behavior, we also cannot
speak to the nursing rates during the night. Nursing may be more common at night if
there is less opportunity for the mother to forage, although bottom-feeding has been
documented during the night in the GOM (Friedlaender et al., 2009; Parks et al., 2014).
Currently, there are no available video tags that would allow for night vision to confirm
nursing; however, as more data are gathered, a more robust model will be created to search
past and future tag data for periods of low (mothers) and high (calves) ODBA and FSR as
an accelerometer signature for nursing on the feeding grounds.

CONCLUSIONS

This study is the first time the duration, relative frequency, and depth of confirmed nursing
events have been described for humpback whales on their feeding grounds. It is apparent
that females can nurse their calves between foraging events as they balance their needs and
the needs of their calf. Many studies rely on surface behavior (e.g., the calf alternating sides
while surfacing on either side of their mother and diving in towards their mother’s pectoral
flipper or peduncle) (Clapham & Mayo, 1987; Smultea et al., 2017; Videsen et al., 2017) to
identify probable nursing behavior; however, during our study the majority of the time
the calf spent close to the mother’s ventral side was not spent nursing. Therefore, caution
should be used when using surface behaviors to quantify nursing behavior on feeding
grounds since it likely overestimates the amount of time a mother-calf pair is nursing.
Lastly, mother-calf pairs are able to coordinate their behavior on the feeding grounds,
allowing calves to remain in close proximity to their mother, and even nurse, while she is
engaged in higher overall activity states, such as foraging.

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Author Contributions
• Jennifer E. Tackaberry and David E. Cade conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, and approved the final draft.
• Jeremy A. Goldbogen, David N. Wiley and Ari S. Friedlaender conceived and designed the experiments, performed the experiments, authored or reviewed drafts of the paper, and approved the final draft.
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REFERENCES

Bache SM, Wickham H. 2014. A forward-pipe operator for r. Available at https://cran.r-project.org/web/packages/magrittr/index.html.

Baker CS, Perry A, Herman LM. 1987. Reproductive histories of female humpback whales megaptera novaeangliae in the North Pacific. Marine Ecology Progress Series 41:103–114 DOI 10.3354/meps041103.

Baraff L, Weinrich MT. 1993. Separation of humpback whale mothers and calves on a feeding ground in early autumn. Marine Mammal Science 9(4):431–434 DOI 10.1111/j.1748-7692.1993.tb00476.x.

Cade DE, Barr KR, Calambokidis J, Friedlaender AS, Goldbogen JA. 2018. Determining forward speed from accelerometer jiggle in aquatic environments. The Journal of Experimental Biology 221(2):jeb170449 DOI 10.1242/jeb.170449.

Cade DE, Friedlaender AS, Calambokidis J, Goldbogen JA. 2016. Kinematic diversity in rorqual whale feeding report kinematic diversity in rorqual whale feeding mechanisms. Current Biology 26(19):2617–2624 DOI 10.1016/j.cub.2016.07.037.

Chittleborough RG. 1958. The breeding cycle of the female humpback whale, megaptera nodosa (Bonnaterre). Marine and Freshwater Research 9(1):1–18 DOI 10.1071/MF9580001.

Clapham PJ, Mayo CA. 1987. Reproduction and recruitment of individually identified humpback whales, megaptera novaeangliae, observed in Massachusetts Bay, 1979–1985. Canadian Journal of Zoology 65(12):2853–2863 DOI 10.1139/z87-434.

Dybas CL. 2018. Over the hump: beleaguered in whaling days, humpback whales chart a new course in the Gulf of Maine. Oceanography 31(3):6–11 DOI 10.5670/oceanog.2018.314.

Friedlaender AS, Hazen EL, Nowacek DP, Halpin PN, Ware C, Weinrich MT, Hurst T, Wiley D. 2009. Diel changes in humpback whale megaptera novaengliae feeding behavior in response to sand lance ammodytes spp. behavior and distribution. Marine Ecology Progress Series 395:91–100 DOI 10.3354/meps08003.

Geraci JR, Lounsbury VJ. 2005. Marine mammals ashore: a field guide for strandings. 2nd ed. United States: National Aquarium in Baltimore.

Gero S, Whitehead H. 2007. Suckling behavior in sperm whale calves: observations and hypotheses. Marine Mammal Science 23(2):398–413 DOI 10.1111/j.1748-7692.2007.00113.x.

Gleiss AC, Wilson RP, Shepard ELC. 2011. Making overall dynamic body acceleration work: on the theory of acceleration as a proxy for energy expenditure. Methods in Ecology and Evolution 2(1):23–33 DOI 10.1111/j.2041-210X.2010.00057.x.

Goldbogen JA, Cade DE, Boersma AT, Calambokidis J, Kahane-Rapport SR, Segre PS, Stimpert AK, Friedlaender AS. 2017. Using digital tags with integrated video and
inertial sensors to study moving morphology and associated function in large aquatic vertebrates. *The Anatomical Record* **300**(11):1935–1941 DOI 10.1002/ar.23650.

**Gulland FMD, Nutter FB, Dixon K, Calambokidis J, Schorr G, Barlow J, Rowles T, Wilkin S, Spradlin T, Gage I, Mulso J, Reichmuth C, Moore M, Smith J, Folken P, Hanser SF, Jang S, Scott Baker C. 2008.** Health assessment, antibiotic treatment, and behavioral responses to herding efforts of a cow-calf pair of humpback whales (megaptera novaeangliae) in the Sacramento River Delta, California. *Aquatic Mammals* **34**(2):182–192 DOI 10.1578/AM.34.2.2008.182.

**Hain JHW, Ellis SL, Kenney RD, Clapham PJ, Gray BK, Weinrich MT, Babb IG. 1995.** Apparent bottom feeding by humpback whales on Stellwagen Bank. *Marine Mammal Science* **11**(4):464–479 DOI 10.1111/j.1748-7692.1995.tb00670.x.

**Hammond PS, Mizroch SA, Donovan GP. 1990.** International whaling commission individual recognition of cetaceans: use of photo-identification and other techniques to estimate population parameters edited by Report of the International Whaling Commission Special Issue 12 Cambridge 1990. *International Whaling Commission Individual* 12:171–175.

**Johnson MP, Tyack PL. 2003.** A digital acoustic recording tag for measuring the response of wild marine mammals to sound. *IEEE Journal of Oceanic Engineering* **28**(1):3–12 DOI 10.1109/JOE.2002.808212.

**López LMM, De Soto NA, Miller P, Johnson M. 2016.** Tracking the kinematics of caudal-oscillatory swimming: a comparison of two on-animal sensing methods. *The Journal of Experimental Biology* **219**(14):2103–2109 DOI 10.1242/jeb.136242.

**Mann J, Connor RC, Tyack PL, Whitehead H (eds.) 2000.** *Cetacean societies, field studies of dolphin and whales*. Chicago: University of Chicago Press.

**MATLAB. 2014.** MATLAB and Statistics Toolbox Release 2014a. Version 8.3.0.532. Natick: The MathWorks, Inc. Available at https://www.mathworks.com/products/matlab.html?s_tid=hp_ff_p_matlab.

**Oftedal OT. 1997.** Lactation in whales and dolphins: evidence of divergence between baleen- and toothed-species. *Journal of Mammary Gland Biology and Neoplasia* **2**(3):205–230 DOI 10.1023/A:1026328203526.

**Parks SE, Cusano DA, Stimpert AK, Weinrich MT, Friedlaender AS, Wiley DN. 2014.** Evidence for acoustic communication among bottom foraging humpback whales. *Scientific Reports* **4:**1–7 DOI 10.1038/srep07508.

**Pinheiro J, Bates D, DebRoy S, Sarkar D. 2019.** Nlme: linear and nonlinear mixed effects models. Available at https://cran.r-project.org/package=nlme.

**R Core Team. 2018.** R: a language and environment for statistical computing. Vienna: R Foundation for Statistical Computing. Available at https://www.r-project.org/.

**RStudio Team. 2016.** RStudio: integrated development environment for R. Boston: RStudio, Inc. Available at http://www.rstudio.com/.

**Saloma AT, Marchesseau S, Charrier I, Andrianarimisa A, Antogiorgi E, Adam O. 2018.** Do the new-born calves of humpback whales (megaptera novaeangliae) have a preference to position themselves at the side of their mother? *Western Indian Ocean Journal of Marine Science Special Issue* 1:1–9.
Smultea MA, Fertl D, Bacon CE, Moore MR, James VR, Würsig B. 2017. Cetacean mother-calf behavior observed from a small aircraft off Southern California. *Animal Behavior and Cognition* 4(1):1–23 DOI 10.12966/abc.01.02.2017.

Steiger GH, Calambokidis J. 2000. Reproductive rates of humpback whales off California. *Marine Mammal Science* 16(1):220–239 DOI 10.1111/j.1748-7692.2000.tb00914.x.

Stimpert AK, Mattila D, Nosal EM, Au WWL. 2012. Tagging young humpback whale calves: methodology and diving behavior. *Endangered Species Research* 19(1):11–17 DOI 10.3354/esr00456.

Tyson RB, Friedlaender AS, Ware C, Stimpert AK, Nowacek DP. 2012. Synchronous mother and calf foraging behaviour in humpback whales megaptera novaeangliae: insights from multi-sensor suction cup tags. *Marine Ecology Progress Series* 457:209–220 DOI 10.3354/meps09708.

Videsen SKA, Bejder L, Johnson M, Madsen PT. 2017. High suckling rates and acoustic crypsis of humpback whale neonates maximise potential for mother–calf energy transfer. *Functional Ecology* 31(8):1561–1573 DOI 10.1111/1365-2435.12871.

Ware C, Arsenault R, Plumlee M, Wiley D. 2006. Visualizing the underwater behavior of humpback whales. *IEEE Computer Graphics and Applications* 26(4):14–18.

Ware C, Wiley DN, Friedlaender AS, Weinrich M, Hazen EL, Bocconcelli A, Parks SE, Stimpert AK, Thompson MA, Abernathy K. 2014. Bottom side-roll feeding by humpback whales (megaptera novaeangliae) in the Southern Gulf of Maine, U.S.A. *Marine Mammal Science* 30(2):494–511 DOI 10.1111/mms.12053.

Wickham H. 2016. Ggplot2: elegant graphics for data analysis. New York: Springer-Verlag. Available at https://ggplot2.tidyverse.org .

Wiley D, Ware C, Bocconcelli A, Cholewiak D, Friedlaender A, Thompson M, Weinrich M. 2011. Underwater components of humpback whale bubble-net feeding behaviour. *Behaviour* 148(5–6):575–602 DOI 10.1163/000579511X570893.

Wilson RP, White CR, Quintana F, Halsey LG, Liebsch N, Martin GR, Butler PJ. 2006. Moving towards acceleration for estimates of activity-specific metabolic rate in free-living animals: the case of the cormorant. *Journal of Animal Ecology* 75(5):1081–1090 DOI 10.1111/j.1365-2656.2006.01127.x.

Zoidis AM, Lomac-MacNair KS. 2017. A note on suckling behavior and laterality in nursing humpback whale calves from underwater observations. *Animals* 7(7):1–12.