INTRODUCTION

The foraging success of the endangered Hawaiian monk seal Monachus schauinslandi at French Frigate Shoals Atoll (FFS) has been a focus of study for 2 decades (e.g. DeLong et al. 1984, Goodman-Lowe 1998, Parrish et al. 2002, Littnan et al. 2004, Stewart et al. 2006). Although the largest subpopulation of monk seals persists at this atoll, its numbers have steadily declined there, following a decrease initially identified in the late 1980s and early 1990s (Gilmartin et al. 1993, Gilmartin & Eberhardt 1995). Subsequent demographic studies by Ragen & Lavigne (1999) identified juvenile seals as the segment of the population with the lowest survival (Antonelis et al. 2006). Size at weaning was correlated with survival for most cohorts (Craig & Ragen 1999), and the overall poor body condition of juvenile seals has been primarily attributed to starvation rather than disease (Reif et al. 2004, Aguirre et al. 2007).

Recent studies have focused on various aspects of monk seal foraging behavior and have included scat analysis to define diet (Goodman-Lowe 1998, Longenecker et al. 2006), satellite telemetry to define oceanic movement, analysis of dive patterns to describe foraging behavior (Stewart et al. 2006, Parrish & Abernathy 2006), use of seal-mounted video cameras to understand prey selection and habitat use (Parrish et al. 2000, 2002, 2005), and evaluation of oceanographic factors to assess potential influences on prey resources and seal survival (Schmelzer 2000, Antonelis et al. 2003, Baker et al. 2006). There is also a growing appre-
cation that the NWHI (Northwestern Hawaiian Islands) is one of the few remaining apex-dominated marine ecosystems (Sudekum et al. 1991, Friedlander & DeMartini 2002) that has prompted interest in the degree of inter-specific competition for prey between monk seals and large predatory fish of the region. The unusually high number of sharks, jacks, and snappers in the NWHI is associated with the regional absence of fishing and exerts uncommon top-down pressure on the reef ecosystem (Sudekum et al. 1991, Friedlander & DeMartini 2002, Parrish & Boland 2004, DeMartini & Friedlander 2006). Data obtained from diet studies of the monk seal (Goodman-Lowe 1998), sharks (DeCrosta et al. 1984), and jacks (Sudekum et al. 1991) indicate that these predators feed in similar habitats on the same prey types and thus, at times, may compete for the same resources. Inter-specific competition is difficult to document for highly mobile species that are concealed from view underwater. Seal-mounted video cameras such as National Geographic Television’s CRITTERCAM (Marshall 1998) affords one method to address the topic. Eight years of research using CRIT-TERCAMs on monk seals at French Frigate Shoals to identify various aspects of seal foraging behavior has generated a sizable sample of video recordings of interactions between seals and predatory fish. In the present study, we investigate the level of apparent inter-specific competition among seals and predatory fish by defining the frequency of interactions, identifying the size and species involved, and determining the habitat in which the interactions occurred.

MATERIALS AND METHODS

Study site. FFS is a crescent-shaped atoll bounding a lagoon with a barrier reef running along its northern perimeter and diagonally through its center (Fig. 1). The sand islets located along this diagonal barrier reef are the only land that monk seals have within a 60 mile radius on which to haul out. During summer and fall months of 1995 to 2002, 42 monk seals (adults and juveniles) were captured, sedated with diazepam, and instrumented with programmable digital video cameras [CRITTERCAM: National Geographic Missions Program, Wild Insight Venus UTPR [Underwater Timed Picture Recorder]] using 10-min epoxy to glue the units to the dorsal pelage (Parrish et al. 2000). The camera was oriented such that the field of view was forward, with the seal's head just visible at the bottom of the image (Fig. 2). The size of the instrument deployed depended on the size and maturity of the seal (Table 1), with the smaller instruments attached to juvenile seals. The camera units also included very high frequency radio tags and time depth recorders. Such instrumentations have been shown not to compromise seal survival or ability to forage (Baker & Johanos 2002, Littnan et al. 2004). Sedated seals remained awake during the entire procedure and returned to the water unassisted immediately after recovering from the sedative. The entire procedure never exceeded 60 min. Seal haul-out behavior and location was monitored every 3 h throughout each deployment from the northwest end of the atoll at Tern Island, FFS. The cameras were recovered by recapturing each seal; the time between deployment and recapture ranged between 1 and 10 d following instrumentation.

Seal dive depths were digitally logged every 10 s throughout the camera deployment. The video cameras were programmed to record segments of images and sound for 6 min each daylight hour (1.5 min were recorded every 15 min, or 3 min every 30 min). A sea water conductivity and depth/pressure switch (<1 m) was used to cease recording when the seal was on the beach or at the surface to maximize collection of images during foraging activity. If the seal surfaced while the camera was still recording, the segment was interrupted and the tape saved for future recording segments. The total surveillance time for the seals ranged from 36 to 72 h depending on the size of the tape cartridge used and the amount of time the seal spent at sea. Five seals were fitted with night vision CRITTERCAMs, and the sampling was limited to evening hours only. The night vision systems were able to ‘see’ the bottom area extending 3 m in front of the seal (Parrish et al. 2002).

Scoring of data. Video images were reviewed and scored into a set of standardized variables entered into a relational database. For each video segment, the depth, type of seal behavior (e.g. swimming, resting, bottom-searching), type of habitat (e.g. atoll, bank, open ocean), feeding events, and the maximum number of predators present were recorded. The forward-facing mount of the camera could not document the presence of ‘following’ predators, and thus the count of fish predators is unavoidably biased low. Side-to-side motion of the seal’s head as it passed over the bottom was clearly swimming behavior. Segments showing the seal motionless on the sea floor, often under a ledge or in a cave, were classified as resting behavior. Bottom-searching behavior was quantified for each segment by scoring the number of times that the seal probed the substrate with its muzzle (standardized for observation time) while moving along the sea floor. The maximum number of predatory fish was the greatest number present on screen at any one time during the video segment. Behaviors of the predatory fish were binned into 4 different classes: ‘escort’ — the fish were clearly swimming in association with the seal,
visible either in front or peripherally; ‘in-face’—the predatory fish nosed in close to the mouth of the seal while the seal probed the bottom for food; ‘feeding’—fish was consuming a prey item; and ‘stealing’—fish either took or attempted to take a prey item from the mouth of the seal.

**Analysis.** Descriptive statistics were used to summarize monk seal encounters with predatory fish. For comparison the numbers of competing fish tallied from the video were standardized by the run time of each image segment. Mean values were used in the comparisons of individual seals to avoid possible effects of consecutively recorded video segments. Data that violated assumptions of homogeneity or were not normally distributed were analyzed using nonparametric techniques (Siegel & Castellan 1988). Statistical significance for all comparisons was set at 0.05. The smallest sample size used in comparisons permitted assessments of significant effects (effect size = 0.50) with a power of 0.66.

**RESULTS**

A total of 3192 video segments were collected comprising 69 h of underwater surveillance with a mean of 1.7 h (±0.9 h SD) per seal. The mean duration of video segments was 83 ± 47 s. Twenty-two percent of the video segments were low-light images collected during crepuscular hours of the daytime video sampling or at night when the night-vision cameras were recording images. Video segments showed that jacks Caranx ignobilis, Seriola dumerili, Caranx melanpygus were the predatory fish species most frequently encountered (mean 0.19 ± 0.55 video segments seal⁻¹) (Fig. 3). The jacks were easily identifiable by their bold movements, often approaching the seals head on. The next most common predator was the large-bodied grey snapper Aprion virens (0.10 ± 0.48 segments seal⁻¹), also easily distinguished. Carcharhinid sharks were the third most common predatory fish (0.02 ± 0.03 video segments seal⁻¹) but were difficult to identify at
the species level, with the exception of white-tip reef shark *Triaenodon obesus*, which were mostly recorded when seals rested in underwater caves. At night, the grey snapper was seen once, sharks were seen in 5 segments and jacks were present in 22 of the sampling segments.

Table 1. CRITTERCAM deployments by year with number of seals (n), seal age, class, sex, camera size, and surveillance parameters.

<table>
<thead>
<tr>
<th>Year</th>
<th>n</th>
<th>Seal age</th>
<th>Sex</th>
<th>Camera (kg in air)</th>
<th>Camera size (cm)</th>
<th>Surveillance Sampling</th>
<th>Day/Night</th>
</tr>
</thead>
<tbody>
<tr>
<td>1995–97</td>
<td>5</td>
<td>Adult</td>
<td>M</td>
<td>CCAM (2 kg)</td>
<td>35 × 10</td>
<td>180 s / 30 min</td>
<td>Day</td>
</tr>
<tr>
<td></td>
<td>22</td>
<td>Adult</td>
<td>M</td>
<td>CCAM (2 kg)</td>
<td>35 × 10</td>
<td>90 s / 15 min</td>
<td>Day</td>
</tr>
<tr>
<td>2000</td>
<td>5</td>
<td>Adult</td>
<td>M</td>
<td>CCAM (2.5 kg)</td>
<td>45 × 10</td>
<td>180 s / 30 min</td>
<td>Night</td>
</tr>
<tr>
<td>2002</td>
<td>1</td>
<td>Adult</td>
<td>M</td>
<td>VENUS (0.75 kg)</td>
<td>15 × 15</td>
<td>90 s /15 min</td>
<td>Day</td>
</tr>
<tr>
<td></td>
<td>9</td>
<td>Juvenile</td>
<td>M and F</td>
<td>CCAM (1.08 kg)</td>
<td>25 × 7.5</td>
<td>90 s /15 min</td>
<td>Day</td>
</tr>
</tbody>
</table>

**Patterns in predatory fish encounters**

Overall, 0.0037 ± 0.007 predatory fish were seen per video segment per seal. Their presence in similar numbers and sizes on adjacent video segments suggests that many of the predatory fish seen are the same indi-

![Fig. 2. *Monachus schauinslandi*. Photo of Hawaiian monk seal fitted with a CRITTERCAM. Insets are video captures from the camera showing ‘escort’ of *Caranx ignobilis* (top 2), ‘in face’ behavior of *Seriola dumerili* (bottom right), and a carcharhinid shark (bottom left). The top of the seal’s head is visible at the bottom of each frame.](image-url)
viduals being recorded repeatedly, but this is difficult to verify. When present on video, they averaged $4.5 \pm 6.4$ ind.; the highest number recorded in a single frame was 46. The snappers, jacks and sharks were terminal phase adults (sizes listed in Fig. 3) and often schooled inter-specifically. Most of the video data (total duration 55 h) were collected at the atoll where the seals were instrumented, but some seals ($n = 11$) spent time in the open ocean (7.9 h) and on the summits of neighboring banks ($n = 9; \sim 5.9$ h). The highest densities of predatory fish were recorded accompanying adult seals that were foraging on the neighboring bank summits (Fig. 4). They were clearly attracted to the intense bottom-searching activities near the seal’s head and anterior, where their presence was easily recorded. The degree to which this attraction inflates the numbers of predatory fish on the video was tested in an analysis of covariance (ANCOVA) using location (atoll, bank, open ocean) as a fixed effect and the rate of seal bottom-searching behavior as a covariate. The bottom-searching covariate narrowly missed significance in explaining the numbers of predatory fish ($F (1, 42) = 3.4; p = 0.08; r = 0.11$). A follow-up analysis controlling for bottom-searching behavior showed that the mean numbers of predatory fish were primarily linked to the location variable ($F (2, 42) = 286.3, p < 0.001$), indicating that seals feed at certain sites even though they have high densities of predatory fish. Removing the effect of location, by looking at data only from the banks, revealed the attraction-influence, with the number of predatory fish correlated with seal bottom-searching behavior ($r_s = 0.53, p < 0.001$).

The depths at which seals encountered predatory fish differed from the overall depth distribution of the seals’ movements (Wilcoxon Signed-Rank $Z = -3.9, p < 0.001$) (Fig. 5). All predators were seen at depths shallower than 100 m. Encounters with predators peaked at 60 to 80 m for jacks and snappers (Fig. 5, inset). Seal dive activity at the atoll extended deeper (>200 m) than at the neighboring banks (90 m) and a comparison of seal diving patterns in the overlapping depth range (0 to 90 m) showed that diving patterns between the atoll and bank did not significantly differ (Wilcoxon $Z = -0.14, p = 0.88$). There was also no significant difference in the depth distribution of predatory fish encountered between the atoll and neighboring banks (Wilcoxon $Z = -0.968, p = 0.333$).

During the present study monk seals were never seen engaged in interference competition. Competing predatory fish were recorded on 17% of the video footage for the 42 seals. The prey, bottom-associated fish and invertebrates, were observed burrowed deep in the sand or hiding under rocks and thus presumably otherwise unavailable to the jacks and sharks. A total of 96 feeding events by monk seals were documented: 70 on the slope surrounding the atoll, 25 on the neighboring banks, and 1 from open ocean subphotic depths. In a number of cases it was possible to use the seal’s prey handling to deduce successful capture of the prey items that were blocked from view.

Impact of predatory fish

Escort behavior was the most common behavior noted for the predatory fish taxa (Fig. 6). In almost all cases, predatory fish escorted the seals; however, in rare instances ($n = 3$), seals specifically traveled to a location where jacks or snappers were inspecting the bottom. Video segments with predatory fish were normalized to compare predator encounter rates. *Aprion virens* were encountered at a mean rate of $4 \pm 5.6$ min$^{-1}$; they maintained more than a body length of distance from the seal and never moved near the seal’s head to compete for prey. Jacks were the closest escorts, both as individuals and in schools. Typically, a jack escorting a seal would be less than a body length from the seal’s head (assuming it had not been displaced by other jacks competing for the position closest to the seal). The mean encounter rate for jacks was $3.4 \pm 6.48$ min$^{-1}$. Jacks routinely positioned their mouths within inches of the seal’s nose to maximize their chances of snatching prey items flushed by the bottom prob-
Jacks were routinely observed to capture prey (30 occasions) before the seal could catch it. Sharks were encountered at the lowest rate (mean 0.9 ± 0.5 min⁻¹, but they also tended to trail behind the seals in areas where they were less likely to be recorded by the video. This could mean the magnitude of bias varies among predator species (e.g. the bias is generally greater for sharks than for jacks). Sharks were seen to move in, close to the seal’s mouth, when they were not displaced by the more aggressive jacks. In one segment, a large shark (>2 m) bumped the feeding seal multiple times in an attempt to prompt the seal to drop a fish it was handling.

There were many images of the predatory fish and the seal going after the same prey item, but none of the predatory fish were observed taking a prey item from the seal’s mouth. Comparing the seals’ overall mean foraging success (capture of prey items) with and without predatory fish showed greater feeding success in the presence of predators (Mann Whitney $U = 185386, Z = –13.2, p < 0.001$). The most intense feeding was exhibited by adult seals, who visited the neighboring banks where the highest numbers of predatory fish were observed (3.4 ± 0.8 min⁻¹). The adult seals were clearly capable of obtaining prey, even in the presence of numerous predatory fish. Since juvenile seals did not visit the banks, the comparison was rerun for the mix of adult and juvenile seals at the atoll, where there were fewer predatory fish (mean 0.10 ± 0.05 min⁻¹), and no impact was found (Mann Whitney $U = 75, Z = –3.4, p < 0.01$). A follow-up comparison of prey capture success between the adult and juvenile seals also showed no significant difference (Mann Whitney $U = 91; Wilcoxon W = 497, Z = –1.45, p = 0.226$). Despite these findings it is important to remember that the density of predatory fish was considerably less at the atoll, so the effectiveness of juveniles foraging among competitors is uncertain.
Parrish et al.: Interactions of monk seals and large predatory fish

**DISCUSSION**

**Animal-borne video sampling**

Animal-borne imaging provides an exciting source of data at an unprecedented resolution. It makes possible considerations of habitat type, depth, foraging technique, and the effects of the assemblage of competing predatory fish. It also presents some unique challenges for interpretation and analysis. Using the maximum number of predatory fish recorded in an image frame eliminates recounting of fish within the same video segment, but it is likely some fish were recounted on consecutive video segments. Because predatory fish are attracted to the seal, their presence is typically detected in the first few seconds of the video segment. Longer duration segments provide a greater chance that the camera will capture greater numbers of the predatory fish in a single image frame. In situations with numerous predatory fish the longer the recording time, the greater the number that will be detected, thus improving the abundance count. In cases with few predatory fish the abundance count is less influenced by the recording time. Consequently, the overall probability of capturing predatory fish on video varies with the recording duration and number of predatory fish present. Maximum number, time-of-first-arrival and other image analysis strategies have been effectively used in baited video camera surveys (Ellis & DeMartini 1995). The latter have achieved comparable video data by using standardized orientation and sampling duration in camera deployment. In our case, the attempt to standardize the deployments of animal-borne imaging is further compounded by the dynamic and often unpredictable nature of the monk seals.

![Graph showing number of video segments classified according to type of competitive behavior exhibited by sharks, snappers, and jacks. See 'Materials and methods' for a description of behaviors](image)

**Predatory fish encounters**

Adult seals generally seem indifferent to the jacks, sharks, and snappers, despite the number and proximity of these fish. It is clear that these predatory fish are part of the seal’s foraging landscape. Although the adult seals are larger than most of the competing fish, the juvenile seals are roughly the same size and could have a difficult time capturing prey in the presence of the competitors. To the competing predatory fish, monk seals are an exploitable foraging resource. Snappers, sharks, and especially jacks are quick and more agile than the seals, but they showed much less capacity to detect and flush benthic prey from cover. The seals use their whiskers to brush along the bottom and chase out camouflaged prey (Table 2). They can also dig out wrasses and eels that are buried deep in the sand bottom and they easily flip large rocks (~20 kg) to obtain prey items hiding beneath. The jacks’ awareness of such behavior enables them to swim ahead of the seal and wait near a rock until the seal arrives, moves the rock and flushes prey items from cover.

The numbers of predatory fish varied greatly between video segments. Underwater visual surveys in the NWHI (Friedlander & DeMartini 2002, Parrish & Boland 2004, Holzwarth et al. 2006) by divers are the closest data sets we have to compare with the predatory fish abundances recorded on the CRITTERCAM videos. Intuitively, we might think the observations made by divers and videos from instrumented monk seals should differ greatly, but our notions are surely less important than how the predatory fish (jacks, sharks, and snappers) perceive monk seals and similarly sized scuba divers. The predatory fish of the NWHI have had little or no exposure to boats and divers and thus exhibit bold behavior, enhanced by a history of following and competing with monk seals for prey. Treating the CRITTERCAM video segments as inde-
pended underwater visual surveys (search area estimate 50 m² min⁻¹), the mean number of predatory fish recorded by the Crittercam ranges from a value that is consistent with the number of predators reported in diver visual surveys of the atoll to an unrealistically high value on the nearby banks, where the seals foraged most intensively and encountered the highest density of predatory fish (Table 3). The estimate of predatory fish density from the Crittercam will be dependent on the number of predatory fish present and the attraction of the fish to the seal and its foraging activities. The ANCOVA that assessed foraging success by location and bottom searching indicated that the highest encounter rate with predatory fish was primarily a result of location effect (i.e. bank summits) and, to a lesser degree, the attraction to the seals’ intense bottom-searching behavior. It is possible that the abundances of predatory fish are highest in prey-rich patches, and thus seals are confronted with more competitors if they choose to visit prey-rich patches. Competing with predatory fish at the banks and other prey rich locations is an inherent cost that the seals will endure as long as their foraging success exceeds that experienced at other locations with lower prey density.

The absence of predators during the seals’ oceanic transits, or when they foraged deeper than 100 m, suggests the predator aggregations have some spatial fidelity. Regional tracking and tagging studies have shown no evidence of interisland movement for jacks (Tagawa & Tam 2006, Meyer et al. 2007a), snappers (Meyer et al. 2007b), and some sharks (Lowe et al. 2006). The deep water that separates the atoll from the banks and seamounts is a sizable barrier that may discourage predatory reef fish are typically found above the thermocline (Thresher & Colin 1986, Chave & Mundy 1994), so any seals foraging deep may reduce, if not avoid, competition. Surveys of the slope (Kelley & Ikehara 2006) and subphotic depths (Parrish 2006) in the NWHI indicate fewer predatory fish than the numbers seen at shallower depths. One survey of predatory fish abundance at the bank closest to FFS used a subsensible to travel from the 500 m contour to the 60 m summit and encountered only 2 Seriola dumerili deeper than the summit; at the summit, a school of >50 Caranx ignobilis and Aprion virescens were encountered (F. Parrish unpubl. data).

Seals instrumented in satellite telemetry studies have been documented routinely diving below 100 m to forage on the mesophotic slope and in subphotic depths (Stewart et al. 2006). Whether the seals are going to these sites to exploit higher prey density or to avoid aggregations of predatory fish, or both, is unknown. Deep diving by monk seals has been documented since the first trials of telemetry work (DeLong et al. 1984, Schlexer 1984). Thus, monk seals may have always foraged in deepwater habitats below 100 m, but the extent to which such diving has changed over time is not known because there are no other extensive historical dive data sets that can be compared with the recent comprehensive studies (Abernathy 1999, Stewart et al. 2006).

Impact of predatory reef fish

Not all of the competitors exerted equal impact on the seals’ foraging efforts. Snappers showed interest in the seals’ activities but did not actively pursue the same prey item as the seals. Their strategy was to capture other prey items flushed from cover by the seals’ activities. Jacks and sharks were seen pursuing the same prey items that the seals were targeting. The lack of instances of prey being taken from the seals by predatory fish indicates that the seals are adept at handling prey once the prey items have been caught, but they are also threatened with competition at the point of prey capture. There were 30 instances where prey items flushed from cover by the seals were consumed by the competing predatory fish before the seals could obtain them. Had the competitors not been present, this prey would have been available to the monk seals. Numerous images show adult seals mitigating loss of prey by deftly tipping large rocks and slipping their heads underneath to eat the small fish hiding under the rock, while the competing jacks crowd around the edges of the rock trying unsuccessfully to get at the

<table>
<thead>
<tr>
<th>Survey type</th>
<th>Location</th>
<th>Depth (m)</th>
<th>Density (no. ha⁻¹)</th>
<th>Predator survey in the NWHI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Belt transect</td>
<td>Atoll</td>
<td>&lt;20</td>
<td>132</td>
<td>Friedlander &amp; DeMartini (2002)</td>
</tr>
<tr>
<td>Towed-diver</td>
<td>Atoll</td>
<td>&lt;20</td>
<td>53</td>
<td>Holzwarth et al. (2006)*</td>
</tr>
<tr>
<td>survey</td>
<td>Bank</td>
<td>&lt;30</td>
<td>13.6</td>
<td></td>
</tr>
<tr>
<td>Crittercam</td>
<td>Atoll</td>
<td>0–100</td>
<td>39</td>
<td>FFS monk seals</td>
</tr>
</tbody>
</table>

*Mean and SD estimated from the respective publication graphic
prey. This competition was most evident for adult male seals feeding among predatory fish aggregations at the summits of neighboring banks. Juvenile seals with CRITTERCAMs encountered fewer predatory fish, probably because most stayed at FFS Atoll, where observed predatory fish densities were lower than at the bank summits (Fig. 4). Only 1 juvenile seal left the atoll for the neighboring banks, and all of its recorded video was spent at depths deeper than 100 m. Juveniles do not have the body mass of adults and thus are less effective at digging or flipping over large rock fragments to obtain prey (Parrish et al. 2005); consequently, their exposed prey may be more available to competing predatory fish.

The degree of competition that has historically occurred between monk seals and predatory fish is unknown. It is difficult to determine whether prey types have changed or if competition has intensified. Food limitations have been proposed as a result of oceanographic regime shifts that have lowered regional productivity (Polovina et al. 1994, Antonelis et al. 2003, Baker et al. 2007). If true, low productivity may have intensified competition between seals and predatory fish for limited prey. Given the large population of predatory fish in the NWHI and their dietary overlap with the seal, an increase in competition could impact the seals. The magnitude of the impact would depend on the population sizes of predatory fish, and there are few population estimates for NWHI. The most comprehensive, in situ diver estimates of predatory fish densities (for sharks and jacks pooled) across the NWHI range from 30 to 140 predators ha⁻¹ with FFS in the middle (50 ha⁻¹; Holzwarth et al. 2006). It is unknown to what degree these densities were inflated because the predatory fish were attracted to the survey divers. At this point, all population estimates of predatory fish should be regarded with caution. Future studies will need to investigate how historic commercial fishing may have influenced the competition level in the seals’ foraging landscape. It is possible that fishing activities have served to both reduce and intensify competition between seals and assemblages of predatory fish. For example, fishing of jacks and other reef fish in the NWHI occurred at varying levels from the 1920s to the 1950s (Uchida 1979) before the shallow reefs of the region were made refuges (mid 1970s) to protect wildlife, including monk seals. The absence of directed fishing for jacks and sharks in the following decades has maintained predatory fish populations at high levels. Populations of predatory fish may also have grown due to ingestion of discarded undersize lobsters or bait from the region’s 20 yr lobster trap fishery (F. Parrish pers. obs.). This fishery was closed in 2000 (DiNardo & Moffitt 2007), ending the fishery discards and perhaps increasing competition between predatory fish and monk seals. Currently, a very limited bottomfish fishery removes Aprion virescens with other members of the deep slope bottomfish complex; and occasionally, jacks and sharks are caught as bycatch.

The fishing effort in the NWHI does not compare to the intense fishing pressure exerted broadly across the marine ecosystem of the main Hawaiian Islands, which has reduced the observed biomass of sharks and jacks seen on diver surveys to less than a tenth of that reported for NWHI surveys (DeMartini & Friedlander 2006). The removal of predators from the main Hawaiian Islands may reduce competition between seals and predatory fish, and could explain why main Hawaiian Island seals are in excellent body condition and seals in the protected NWHI are emaciated (Antonelis et al. 2006). Future research should address the causes of these patterns and the possibility that inter-specific competition is impacting the survivorship of juvenile seals.

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