

Effectiveness of removals of the invasive lionfish: how many dives are needed to deplete a reef?

Paolo Usseglio^{1,*}, Jason D. Selwyn^{2,*}, Alan M. Downey-Wall³ and J. Derek Hogan²

- ¹ Fundación In-Nova Castilla la Mancha, Spain
- ² HoBi Lab, Department of Life Sciences, Texas A&M University—Corpus Christi, Corpus Christi, TX, United States
- ³ Marine Science Center, Northeastern University, Nahant, MA, United States
- * These authors contributed equally to this work.

ABSTRACT

Introduced Indo-Pacific red lionfish (*Pterois volitans/miles*) have spread throughout the greater Caribbean and are associated with a number of negative impacts on reef ecosystems. Human interventions, in the form of culling activities, are becoming common to reduce their numbers and mitigate the negative effects associated with the invasion. However, marine managers must often decide how to best allocate limited resources. Previous work has identified the population size thresholds needed to limit the negative impacts of lionfish. Here we develop a framework that allows managers to predict the removal effort required to achieve specific targets (represented as the percent of lionfish remaining on the reef). We found an important trade-off between time spent removing and achieving an increasingly smaller lionfish density. The model used in our suggested framework requires relatively little data to parameterize, allowing its use with already existing data, permitting managers to tailor their culling strategy to maximize efficiency and rate of success.

Subjects Ecology, Marine Biology

Keywords Invasive species, Lionfish, Caribbean, Removal efficiency, *Pterois volitans*, Management prioritization

INTRODUCTION

Indo-Pacific red lionfish (*Pterois volitans/miles*) were introduced to South Florida in the 1980's and have since spread throughout the greater Caribbean showing exponential population growth throughout the invaded range (*Betancur-R et al.*, 2011), with populations in the invaded range reaching greater densities than those of their native range (*Kulbicki et al.*, 2012). These population booms have resulted in reduced abundance of native species (*Morris & Akins*, 2009; *Barbour et al.*, 2010; *Côté & Maljkovic*, 2010), decreases in recruitment (*Albins & Hixon*, 2008), and possible shifts in benthic community composition (*Lesser & Slattery*, 2011; *Albins & Hixon*, 2013; *Layman*, *Jud & Nichols*, 2014). Human intervention, in the form of culling, has become common to mitigate lionfish's negative effects (*Morris*, 2012).

Total eradication of lionfish across the invaded range is unlikely due to their high abundance, wide-spread distribution and the high resilience of the species (*Morris, Shertzer*

Submitted 26 April 2016 Accepted 28 January 2017 Published 23 February 2017

Corresponding author Jason D. Selwyn, jason.selwyn@tamucc.edu, jdselwyn@gmail.com

Academic editor Donald Kramer

Additional Information and Declarations can be found on page 12

DOI 10.7717/peerj.3043

© Copyright 2017 Usseglio et al.

Distributed under Creative Commons CC-BY 4.0

OPEN ACCESS

& Rice, 2010; Barbour et al., 2011; Arias-González et al., 2011). However, there is evidence that local-scale removal efforts can reduce lionfish abundance and have benefits for the local native reef community (Morris & Whitfield, 2009; Barbour et al., 2011; Frazer et al., 2012). In natural systems, the severity of the deleterious lionfish effects appear to be system or location-specific (Elise et al., 2015; Albins, 2015), suggesting that reef susceptibility to the negative lionfish effects may be driven by a complex number of locally determined factors, including community structure, complexity, and overall ecosystem health. Overall reef health and initial lionfish density both affect native reef resiliency and thus the necessary removal effort required to mitigate predation-induced declines of native species (Green et al., 2014). Importantly, Green et al. (2014) found that complete eradication of lionfish was not necessary for native species recovery, suggesting that culling of lionfish may be a practical solution to mitigate their impacts.

While spatially restricted culling can effectively reduce lionfish density, there is a lack of key metrics, such as the relationship between effort and percent of the population removed to assess the efficiency of this approach (*Frazer et al.*, 2012). Marine resource managers in the Caribbean, working under budget limitations, would greatly benefit from these metrics as it would be possible to know the effort needed to achieve specific removal targets. A framework to determine the fishing effort required to achieve reductions in lionfish densities, consistent with threshold levels estimated by the model proposed by *Green et al.* (2014), would be a useful management tool. Here we aim to develop such a framework to aid in the effective and efficient culling of lionfish.

To estimate the effort required to reduce lionfish populations by a given percentage, as specified by the difference between the initial and target densities, we followed a three-step approach. First, we culled lionfish populations at multiple sites by conducting removal dives over consecutive days and monitored changes in catch per unit effort at each site over time. Second, we used a depletion model and the aforementioned data to estimate initial lionfish population sizes and lionfish catchability at each site. Finally, we integrated the depletion model results from all sites into a simple exponential model to determine the percentage of the initial population removed for a given amount of effort. Our proposed framework can be easily implemented using data already collected in common culling efforts and as such could be incorporated into existing removal strategies.

METHODS

Study site

We sampled seven sites on Turneffe atoll (17.3638°N, 87.8581°W), Belize, Central America, (Fig. 1). Turneffe atoll is located 9–23 km offshore of the main Belizean Barrier Reef, the largest barrier reef in the Caribbean and second largest globally (*Gibson & Carter*, 2003). The atoll consists of a number of mangrove islands which provide important nursery habitat for a variety of fish species, including lionfish (*Gibson & Carter*, 2003; *Mumby*, 2006; *Claydon*, *Calosso & Traiger*, 2012). The perimeter of the atoll consists of a barrier reef that transitions into a drop-off (*Chittaro et al.*, 2006). Numerous reef patches are found within the reef lagoon. The northern portion of the windward side consists of spur

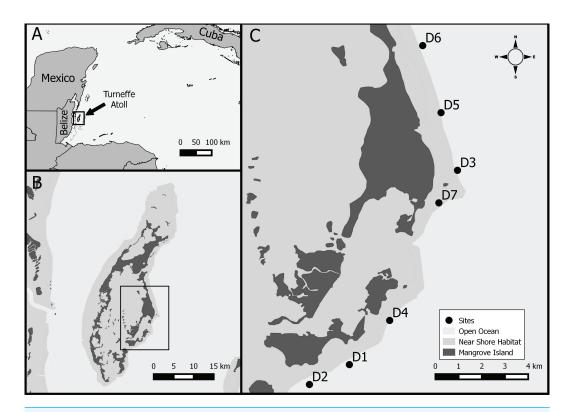


Figure 1 Study location. Study area, (A) Western Caribbean with location of Belize and Turneffe atoll, (B) Turneffe atoll, (C) study sites in southeast windward location on Turneffe Atoll.

and groove habitat while the southern portion is composed of drop-off and wall habitats starting at a depth of \sim 18 m. A more detailed description of the area can be found in (*Garcia & Holtermann*, 1998).

Sampling sites were located along the outer edge of the windward forereef in the central portion of the atoll, ranging in depth from 15–30 m and area from 1,148 to 1,800 m² (1,643 (±238 s.d.) m², Fig. 1, Table 1). Similar to much of the Caribbean, sampling sites generally have low coral cover (4–19%), with some evidence of a coral to algal dominated community shift (*McClanahan & Muthiga, 1998*; *Wild, Jantzen & Kremb, 2014*). Between the reef crest and wall habitats, along the windward side, the reef is dominated by numerous *Orbicella* colonies.

Sampling sites were haphazardly located using natural breaks, such as sand channels, as site boundaries (Fig. 1). Additional marker buoys were placed at the corners of each site to act as guides and help minimize the accidental inclusion of offsite lionfish. Site area was measured using fiberglass tape measures.

Depletion fishing

Divers on SCUBA removed lionfish from each site using pole-spears. For each dive, the number of divers, the time spent hunting for lionfish, the total number of lionfish, and the total length (TL, cm) and mass (g) of each lionfish were recorded. Fishing effort was calculated as diver-hours per dive, the sum of time spent by all divers hunting lionfish at

Table 1 Site descriptions. Descriptions of seven experimental reef sites at Turneffe atoll, Belize, including size (m²), depth range (m), the mean and total number of diver-hours, the number of depletion 2 dives, the total number of lionfish caught, and whether or not we achieved our depletion criterion of two consecutive dives with zero lionfish caught (only the first of which counted towards the number of depletion dives).

Site	Area (m²)	Depth range (m)	Average diver hours per dive (±s.d.)	Total diver hours	Number of depletion dives	Number of lionfish caught	Depletion criterion achieved
D1	1,184	20–30	$1.02~(\pm 0.17)$	5.10	5	35	Yes
D2	1,800	15–30	$1.00~(\pm 0.28)$	7.03	7	43	Yes
D3	1,470	20–30	$1.07~(\pm 0.05)$	4.27	4	17	No
D4	1,800	15–30	$1.05\ (\pm0.40)$	5.23	5	27	Yes
D5	1,800	15–30	$0.98~(\pm 0.38)$	5.85	6	32	No
D6	1,800	19–35	$0.94~(\pm 0.29)$	5.67	6	32	Yes
D7	1,650	20–35	$1.44\ (\pm0.20)$	8.65	6	23	No

each site. For example, three divers hunting lionfish during a 20-min dive would equal one diver-hour. Each site was depleted by repetitively diving the area, over successive days, until no lionfish were observed or caught on consecutive dives. As the areas of the sample sites varied substantially, all dive times and catches were standardized to diver-hours per 1,000 m² and catch per 1,000 m², respectively. The effect of depletion fishing on abundance was estimated by calculating catch per unit effort (CPUE) as:

$$CPUE = \frac{C}{h}; (1)$$

where C is the catch per dive per 1,000 m² and h is the fishing effort expressed in diver-hours per 1,000 m². The use of diver-hours per 1,000 m² as the metric of effort assumes no change in catch efficiency with increased numbers of divers, for example if divers communicate to help find fish.

Depletion model

We used a Leslie depletion model (*Leslie & Davis*, 1939) to estimate the initial population size per 1,000 m² on all sites. A depletion model observes how the removal of fish affects the relative abundance of fish remaining in the population (*Hilborn & Walters*, 1992), assuming that repeated fishing over a small area will reduce local population size (*Ogle*, 2016a).

Initial population size is calculated as:

$$N_0 = \sum_{t=1}^{T} n_t; (2)$$

where N_0 is the initial population size per 1,000 m², T is the number of removal dives and is indexed by t, and n_t is the catch per 1,000 m² from the t-th removal.

The model assumes that catch per unit effort in the t-th removal event is proportional to the extant population at the time of the t-th removal event:

$$CPUE_t = qN_t; (3)$$

where $CPUE_t$ is the catch per unit effort, N_t is the population size per 1,000 m² for the t-th removal, and q is the catchability coefficient representing the fraction of the population that is removed by one unit of fishing effort.

The Leslie depletion model is in the form of a linear equation:

$$CPUE_t = qN_0 - qK_{t-1}; (4)$$

where $CPUE_t$ is the response variable, and K_{t-1} is the cumulative catch up to time t-1, q is the slope and qN_0 is the intercept (*Leslie & Davis*, 1939; *Ogle*, 2016a). The initial number of lionfish is thus estimated by dividing the intercept by the catchability coefficient q. Confidence intervals for estimates of qN_0 and q were derived from the regression results following the methods outlined in (*Ogle*, 2016a).

The model assumes populations are closed to migration over the sampling period, catchability is constant through time and among individuals, enough fish are removed to substantially reduce CPUE, and the catch removes more than 2% of the population (Ogle, 2016a). We attempted to meet these assumptions by conducting depletion dives on consecutive days, with the same observers, and whenever possible, continuing depletion until zero fish were caught or seen on two consecutive dives. While some fish can exhibit fleeing behavior in response to depletion fishing activities (Giddens et al., 2014), and this has been documented for lionfish (Côté et al., 2014), we saw no such change in lionfish behavior during the course of our dives. This is likely a reflection of high capture efficiency, preventing wounded individuals from learning to avoid diver activity. Additionally, the depletion models for all sites met the standard assumptions of normality and homoscedasticity for linear regression (Figs. S1 and S2). This approach does not assume similarity in catchability among sites, or any other site characteristic, as the initial number of lionfish at each site was estimated using only measurements from that site. Despite this, there may be a relationship between the initial density of lionfish at a site and the catchability coefficient at that site due to, for example, handling time being of relatively greater importance at higher densities. To determine if the estimated initial lionfish density affected the catchability coefficient we performed a linear regression.

Lionfish caught versus dive time

Using the estimated initial population size per 1,000 m² we modeled the percentage of lionfish caught as a function of cumulative dive time per 1,000 m² using an exponential asymptotic growth model following the formula:

$$C = 100\left(1 - e^{-\alpha qt}\right) \tag{5}$$

where α is the marginal reduction in CPUE, t is the cumulative number of diver-hours per 1,000 m², q is the site-specific catchability coefficient, and C is cumulative percentage of lionfish caught at time t. Models were built both including and not including the catchability coefficient to test the hypothesis that catchability interacts with dive-time to affect the cumulative percentage of lionfish caught. Including catchability coefficient as a multiplier of time acts as a site-specific scalar of effort in order to standardize effort across sites with disparate degrees of removal efficacy. To account for potential variation among

sites (beyond what is captured by the catchability coefficient) in coral cover or habitat complexity, for example, this function was modeled to include site as a random factor influencing α . The best model was chosen based on minimization of the bias corrected Akaike Information Criteria (AICc) (*Akaike*, 1973; *Arnold*, 2010), which maximizes model fit while avoiding overfitting by incorporating the log-likelihood of the model, while penalizing for the number of parameters and the small sample size. The best-fit model was used to predict the time required to remove 50% and 90% of the estimated total population of lionfish per 1,000 m² based on the mean catchability coefficient.

All of the above analyses were performed with the statistical package R, version 3.0.2 (*R Development Core Team*, 2013), depletion models were calculated with the FSA package (*Ogle*, 2016b), exponential asymptotic growth models were calculated using the nlme package (*Pinheiro et al.*, 2013), and graphing was done using the package ggplot2 (*Wickham*, 2009). All relevant data, analysis, and code are provided in the supplemental material (Data S1 and Supplemental Information 1).

This research was conducted under permission 000033-14 from the Belize Fisheries Department. Animal use for this project received approval by the Animal Care & Use Committee, Texas A&M University—Corpus Christi, protocol number #05-14.

RESULTS

Depletion fishing

Across all seven sites, divers captured 209 lionfish, ranging in size from 12–41 cm TL (mean $= 25.9 \pm 0.9 \pm 0.9$ s.d.) cm). The total number of fish captured at each site ranged from 17 to 43 lionfish (Table 1). In four of seven sites, we were unable to achieve our depletion criterion (i.e., no lionfish observed on two consecutive dives; Table 1).

Depletion model

Leslie depletion models independently estimated the initial number of lionfish present as 11 to 29 individuals per 1,000 m² per site (mean = 19.5 (± 6.5 s.d.) per 1,000 m⁻², Table 2, Fig. 2). The catchability coefficient was found to range from 0.43 to 0.91 (mean = 0.694 (± 0.170 s.d.), Table 2, Fig. 2) with higher values indicating lionfish were easier to capture at a particular site. The models' r^2 values ranged from 0.68–0.98 and were all significant, suggesting a good fit of the data to the model (Table 2). There was no effect of the initial estimated density of lionfish on the catchability coefficient ($r^2 = 0.13$, p = 0.45; Fig. S3A).

Lionfish caught versus dive time

The best-fit model incorporated among-site variability as a random factor affecting α and the interaction between the catchability coefficient and dive-time (Eq. 6; Table 3; Fig. 3). The model predicted, with a 95% CI, that one hour of fishing effort per 1,000 m², scaled using the mean catchability coefficient (0.69), results in a 58%–66% reduction in the lionfish population per 1,000 m². Four diver-hours per 1,000 m², scaled by the mean catchability coefficient, achieves 97%–99% reductions of the initial population per 1,000 m² (Fig. 3A). The percent reduction of lionfish for a specified amount of effort is inversely related to the catchability coefficient (Fig. 3B). As such when one hour of fishing effort

Table 2 Depletion model results. Summary of results of the depletion diving and estimating initial lionfish abundance based on the Leslie depletion model (Eq. 4; Fig. 2). The initial number of lionfish (N_0), catchability coefficient (q), coefficient of determination (r^2), and p-value (p) are shown for each site (D1–D7). Numbers in parentheses represent 95% confidence intervals of estimates.

Site	N_0	q	r^2	p
D1	29 (26–33)	0.70 (0.53-0.87)	0.98	0.0009
D2	25 (16–34)	0.66 (0.20–1.11)	0.68	0.0137
D3	11 (7–16)	0.91 (0.26–1.56)	0.92	0.0262
D4	15 (9–22)	0.76 (0.15–1.37)	0.78	0.0291
D5	23 (17–29)	0.43 (0.24–0.62)	0.88	0.0035
D6	19 (15–22)	0.86 (0.58–1.15)	0.93	0.0011
D7	14 (13–15)	0.54 (0.43–0.65)	0.97	0.0001

Table 3 Lionfish catch as a function of cumulative dive time. Comparison of the non-linear mixed effects models of cumulative lionfish catch as a function of cumulative dive time. Scaling indicates whether the cumulative dive time was scaled by the catchability coefficient (indicated as q-scaled). Random indicates which models included site as a random factor influencing the multiplication coefficient α . Parameter estimates (\pm SE) are shown based on each model. α is the coefficient in the exponential models multiplied by cumulative dive time, or catchability coefficient (q) and cumulative dive time. K is the number of parameters estimated in the model. logLik is the log likelihood of the model and Δ AICc is the difference in AICc from the best model

Model	Scaling	Random	α	Intercept	K	logLik	AICc	Δ AIC	AICc Weight
1	q-scaled	Site	1.39 ± 0.08		3	-116.56	239.81	0.00	0.99
2	q-scaled		1.35 ± 0.05		2	-122.13	248.59	8.78	0.01
3		Site	0.99 ± 0.12		3	-122.51	251.70	11.89	0.00
4			0.92 ± 0.06		2	-147.47	299.27	59.46	0.00
Null Model				78.09 ± 3.51	2	-173.08	350.50	110.69	0.00

per 1,000 m² is scaled based on the minimum observed catchability coefficient (0.43) there is a 41%–49% reduction in lionfish; when scaled with the maximum catchability coefficient (0.91) there is a 68%–76% reduction in lionfish (Fig. 3B). Similarly, a 50% decrease of the initial population size was achieved after between 1.16–1.17 (minimum observed catchability) to 0.54–0.55 (maximum observed catchability) diver-hours per 1,000 m² depending on the catchability coefficient (0.71–0.72 diver-hours per 1,000 m² based on the mean catchability to 1.80–1.83 (maximum observed catchability) diver-hours per 1,000 m² depending on the catchability coefficient (2.37–2.41 diver-hours per 1,000 m² based on the mean catchability coefficient; Fig. 3B).

$$C = 100(1 - e^{-1.39qt}). (6)$$

DISCUSSION

Here we created a simple, yet powerful, framework for predicting the removal effort required to achieve specific lionfish removal targets (represented as the percent of lionfish remaining on the reef). We demonstrated an important trade-off between time spent removing lionfish and the density reduction. We suggest that—when coupled with a model

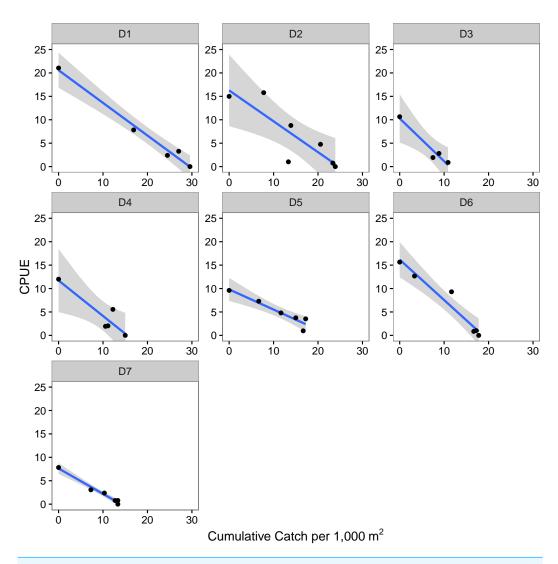


Figure 2 Depletion models. Leslie depletion models relating catch per unit effort (CPUE) to cumulative catch per 1,000 m² that were used to estimate the initial number of lionfish present per 1,000 m² (N_0) and catchability coefficient (q) at each site (D1–D7). The line and shaded area are the regression line and 95% confidence intervals.

explicitly used to determine critical lionfish densities required for native reef recovery (see the model by *Green et al.*, 2014)—our model allows managers to create strategies that not only consider local ecosystem demographics and lionfish densities, but also the resources available (i.e., the number of available divers). This would enable managers to tailor their culling strategy to maximize efficiency and rate of success.

Generally, our model estimated a 50% reduction of lionfish in less than one diver-hour per 1,000 m², and up to 90% reduction in fewer than 2.5 diver-hours per 1,000 m². Given the importance of among-site variation in our model, these values should be applied cautiously, but may be applicable to similar reef systems in the western Caribbean, or used as reference values when other data is unavailable. Importantly, the results demonstrate a well-fit model that requires relatively little data to parameterize. Thus, the model provides

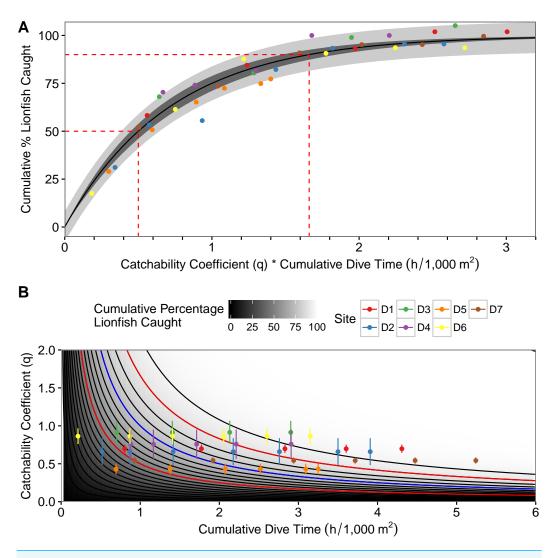


Figure 3 Asymptotic exponential growth model. (A) Plot of asymptotic exponential growth model with percent removal of lionfish per 1,000 m² vs. the product of dive-time per 1,000 m² and the catchability coefficient. Dashed red lines represent time needed to remove 50% and 90% of the population, the darkly shaded area represents the 95% confidence interval, the lighter shaded area represents the 95% prediction interval, and colored points represent different depletion sites (D1–D7). (B) Contour plot of the results of the best-fit model of percent removal of lionfish per 1,000 m² based on dive-time per 1,000 m² and catchability coefficient (Eq. 4). Points represent observed values with standard deviations, and colored points represent different depletion sites (D1–D7). Contours represent the cumulative percent of lionfish caught and increment by 5% with darker colors being closer to 0% and lighter colors closer to 100%. Red contours are the 50% and 90% contour lines and the blue contour is at 75% removal, as used in the theoretical example.

a valuable framework for creating local or system-specific estimates for removal effort per unit area.

Variability among sites was partially accounted for within the model by including a site-specific catchability coefficient. This value reflects unquantified variables that cause catch rate to vary among sites, such as subtle changes in reef structure, rugosity, lionfish behavior, shifts in lionfish size structure, and changes in diver hunting efficiency.

Furthermore, habitat structure and an adaptive behavioral response to culling (i.e., lionfish wariness) have been shown to negatively affect hunting efficiency (*Green et al., 2014*; *Côté et al., 2014*).

The initial lionfish density could affect removal time due to relatively more time spent on handling than on search at high density. Interestingly, we found no effect of initial lionfish density on the catchability coefficient. This result should be interpreted with caution as our analysis was performed over a relatively small range of densities compared to what has been observed by others (110–290 ha⁻¹ this study; \sim 20 ha⁻¹, (*Whitfield et al.*, 2007); 390 ha⁻¹, (*Green & Côté*, 2009); \sim 1,450 ha⁻¹, (*Dahl & Patterson*, 2014).

The efficacy of any removal strategy is expected to be driven partially by the rate of immigration and recruitment into the managed area after a removal. In species with complex, life-stage dependent habitat utilization patterns, the size and abundance of immigrating individuals may be habitat-specific. In lionfish, several studies have found evidence of an ontogenetic shift, with new recruits and juveniles utilizing protected, shallow water habitat prior to migrating out to deeper forereefs (Barbour et al., 2010; Claydon, Calosso & Traiger, 2012; A Downey-Wall & JD Hogan, 2015, unpublished data). Consequently, management strategies that focus on removing lionfish from forereefs (i.e., this study) will likely experience the highest immigration pressure from juvenile and adult life-stage fish. Lionfish at these life stages have demonstrated an ability to move thousands of meters, but tend to exhibit comparatively small home range tendencies (Bacheler et al., 2015; Tamburello & Côté, 2015). Furthermore, lionfish immigration on patch reefs has been shown to be density independent (Benkwitt, 2013), suggesting immigration pressure will remain similar regardless of the removal area density. Overall, creating estimates for immigration rate will be essential for long-term management success, as it can drive both the frequency and intensity of removals. We suggest resolving recolonization rate, particularly at different removal areas, habitat types, and depths, should be an important priority for future research.

The framework presented here for predicting lionfish removal effort provides a starting point for management. However, there are a number of factors that can affect long-term sustainability of culling programs, the effect of which are not yet understood. For example, the size of the area to be managed, the number and experience of team members and the speed at which recolonization happens post-culling can affect costs and efficiency of removals and therefore impact sustained management. This study only investigated relatively small sites (~1,600 m²) using a consistent and minimal team of experienced divers (2–3) and did not repetitively cull a single site to maintain lionfish densities as one might as part of a management strategy. It is unknown how the effort required to remove lionfish to achieve or maintain target densities will vary in response to changes in these factors. Some factors may cause non-linear changes in efficiency or costs; for example, increasing the number of divers may lead to greater improvement than expected due to diver communication, but will also cost more than increasing dive times or number of dives per personnel member.

Despite these remaining challenges, it is clear that a simple framework for lionfish culling programs can aid in management decision-making. In order to maximize the outcome of

lionfish fishing activities, we suggest a framework with a series of steps that managers could follow. The first step is to survey the target reefs to assess densities of both lionfish and native species. Following *Green et al.* (2014), managers can use survey data as inputs into their model in order to calculate reef specific lionfish density thresholds. To determine the appropriate amount of effort required to meet those threshold densities, lionfish depletion dives can then be performed at all or some of the sensitive sites following the methods presented in this paper. Subsequent or additional removal efforts can then be planned using the relationship between diver effort and percent removal of lionfish.

As a hypothetical example, upon surveying fish populations at a site and using the model in *Green et al.* (2014) we found a target density of 7 lionfish/1,000 m² with an initial density of 29 lionfish/1,000 m² (a reduction of 75%). We would next use the model derived from the depletion dives shown here (Eq. 6) to calculate that the product of the cumulative dive time and catchability required to achieve this goal is ~1 scaled diver-hour. Then, we divide this estimate by various catchability values to see a range of estimates based on different scenarios (e.g., inexperienced divers, particularly high complexity reef; Fig. 3B blue contour). Using the range of catchability estimates from our depletion dives (0.43–0.91, Table 2) we would then estimate that it would take between 1 and 2.3 diver hours to achieve the target density at the hypothetical site. Having this information, we can determine the optimal number of divers needed and identify specific divers with known catch rates to determine our personnel composition. This allows us to estimate the amount of fuel and boat time required to accomplish the task. These estimates are vital to agencies with limited resources as they allow for more accurate budgeting of time and money.

Controlling invasive lionfish populations has become a major objective of many agencies involved in the management of coral reefs in the tropical and sub-tropical Atlantic. Given that resources such as money, time and personnel are limited, managers need tools to effectively plan and allocate resources to achieve lionfish control objectives. Current best practices suggest targeting removal sites of particular interest: ecologically sensitive sites (such as marine protected areas) or tourism areas (*Morris*, 2012). Generally, it is advisable for these areas to be small to allow for tractable removal efforts with limited resources. The methods applied here are broadly applicable as a straightforward way for managers to estimate the effort needed to achieve target levels of lionfish in an area of interest. As this framework incorporates only data commonly collected during removal efforts (catch per dive and dive time), it could be implemented as part of already existing management strategies, including using previously culled sites to calculate the continued effort needed into the future. By following these steps, managers can balance their field needs with their assets in order to choose the best course of action based on the specific circumstances they face.

ACKNOWLEDGEMENTS

We thank E Anderson for help with removal dives, and staff at the University of Belize Institute for Marine Science, Calabash Caye field station for field and logistical support. We are also grateful to C van Slyck, D Kramer, H Vallès, and one anonymous reviewer whose comments and critique greatly improved this manuscript.

ADDITIONAL INFORMATION AND DECLARATIONS

Funding

This work was funded by Texas A&M University and Corpus Christi start-up funds (JDH). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Grant Disclosures

The following grant information was disclosed by the authors: Texas A&M University and Corpus Christi start-up funds (JDH).

Competing Interests

The authors declare there are no competing interests.

Author Contributions

- Paolo Usseglio and Jason D. Selwyn conceived and designed the experiments, performed
 the experiments, analyzed the data, contributed reagents/materials/analysis tools,
 wrote the paper, prepared figures and/or tables, reviewed drafts of the paper, authors
 contributed equally.
- Alan M. Downey-Wall and J. Derek Hogan conceived and designed the experiments, performed the experiments, contributed reagents/materials/analysis tools, wrote the paper, reviewed drafts of the paper.

Animal Ethics

The following information was supplied relating to ethical approvals (i.e., approving body and any reference numbers):

Animal use for this project received approval by the Animal Care & Use Committee, Texas A&M University—Corpus Christi protocol number #05-14.

Field Study Permissions

The following information was supplied relating to field study approvals (i.e., approving body and any reference numbers):

This research was conducted under permission 000033-14 from the Belize Fisheries Department.

Data Availability

The following information was supplied regarding data availability:

The raw data has been supplied as a Supplemental Dataset.

Supplemental Information

Supplemental information for this article can be found online at http://dx.doi.org/10.7717/peerj.3043#supplemental-information.

REFERENCES

- **Akaike H. 1973.** Information theory as an extension of the maximum likelihood principle. In: Petrov BN, Csaki F, eds. *Second international symposium on information theory*. Budapest: Akademiai Kiado, 267–281.
- **Albins M. 2015.** Invasive Pacific lionfish *Pterois volitans* reduce abundance and species richness of native Bahamian coral-reef fishes. *Marine Ecology Progress Series* **522**:231–243 DOI 10.3354/meps11159.
- Albins M, Hixon M. 2008. Invasive Indo-Pacific lionfish *Pterois volitans* reduce recruitment of Atlantic coral-reef fishes. *Marine Ecology Progress Series* 367:233–238 DOI 10.3354/meps07620.
- **Albins MA, Hixon MA. 2013.** Worst case scenario: potential long-term effects of invasive predatory lionfish (*Pterois volitans*) on Atlantic and Caribbean coral-reef communities. *Environmental Biology of Fishes* **96(10)**:1151–1157 DOI 10.1007/s10641-011-9795-1.
- Arias-González JE, González-Gándara C, Luis Cabrera J, Christensen V. 2011. Predicted impact of the invasive lionfish *Pterois volitans* on the food web of a Caribbean coral reef. *Environmental Research* 111(7):917–925 DOI 10.1016/j.envres.2011.07.008.
- **Arnold TW. 2010.** Uninformative parameters and model selection using Akaike's Information Criterion. *The Journal of Wildlife Management* **74(6)**:1175–1178 DOI 10.1111/j.1937-2817.2010.tb01236.x.
- Bacheler N, Whitfield P, Muñoz R, Harrison B, Harms C, Buckel C. 2015. Movement of invasive adult lionfish *Pterois volitans* using telemetry: importance of controls to estimate and explain variable detection probabilities. *Marine Ecology Progress Series* 527:205–220 DOI 10.3354/meps11241.
- Barbour AB, Allen MS, Frazer TK, Sherman KD. 2011. Evaluating the potential efficacy of invasive lionfish (*Pterois volitans*) removals. *PLOS ONE* **6**(5):e19666 DOI 10.1371/journal.pone.0019666.
- Barbour A, Montgomery M, Adamson A, Díaz-Ferguson E, Silliman B. 2010. Mangrove use by the invasive lionfish *Pterois volitans*. *Marine Ecology Progress Series* 401:291–294 DOI 10.3354/meps08373.
- **Benkwitt CE. 2013.** Density-dependent growth in invasive lionfish (*Pterois volitans*). *PLOS ONE* **8(6)**:e66995 DOI 10.1371/journal.pone.0066995.
- Betancur-R R, Hines A, Acero PA, Ortí G, Wilbur AE, Freshwater DW. 2011. Reconstructing the lionfish invasion: insights into Greater Caribbean biogeography. *Journal of Biogeography* 38(7):1281–1293 DOI 10.1111/j.1365-2699.2011.02496.x.
- **Chittaro PM, Usseglio P, Fryer BJ, Sale PF. 2006.** Spatial variation in otolith chemistry of *Lutjanus apodus* at Turneffe Atoll, Belize. *Estuarine Coastal and Shelf Science* **67(4)**:673–680 DOI 10.1016/j.ecss.2005.12.014.
- Claydon J, Calosso M, Traiger S. 2012. Progression of invasive lionfish in seagrass, mangrove and reef habitats. *Marine Ecology Progress Series* 448:119–129 DOI 10.3354/meps09534.

- Côté IM, Darling ES, Malpica-Cruz L, Smith NS, Green SJ, Curtis-Quick J, Layman C. 2014. What doesn't kill you makes you wary? Effect of repeated culling on the behaviour of an invasive predator. *PLOS ONE* **9**(4):e94248

 DOI 10.1371/journal.pone.0094248.
- **Côté I, Maljkovic A. 2010.** Predation rates of Indo-Pacific lionfish on Bahamian coral reefs. *Marine Ecology Progress Series* **404**:219–225 DOI 10.3354/meps08458.
- **Dahl KA, Patterson WF. 2014.** Habitat-specific density and diet of rapidly expanding invasive red lionfish, *Pterois volitans*, populations in the northern Gulf of Mexico. *PLOS ONE* **9(8)**:e105852 DOI 10.1371/journal.pone.0105852.
- Elise S, Urbina-Barreto I, Boadas-Gil H, Galindo-Vivas M, Kulbicki M. 2015. No detectable effect of lionfish (*Pterois volitans* and *P. miles*) invasion on a healthy reef fish assemblage in Archipelago Los Roques National Park, Venezuela. *Marine Biology* **162(2)**:319–330 DOI 10.1007/s00227-014-2571-y.
- Frazer TK, Jacoby CA, Edwards MA, Barry SC, Manfrino CM. 2012. Coping with the lionfish invasion: can targeted removals yield beneficial effects? *Reviews in Fisheries Science* 20(4):185–191 DOI 10.1080/10641262.2012.700655.
- Garcia E, Holtermann K. 1998. Calabash Caye, Turneffe Islands Atoll, Belize. In: Kjerfve B, ed. *Caricomp: Caribbean coral reef, seagrass and mangrove sites. Coastal region and small islands unit.* Paris: UNESCO, 347.
- **Gibson J, Carter J. 2003.** The reefs of Belize. In: *Latin American coral reefs*. Amsterdam: Elsevier Press.
- Giddens J, Friedlander A, Conklin E, Wiggins C, Stamoulis K, Donovan M. 2014.

 Experimental removal of the invasive peacock hind (roi) *Cephalopholis argus*, in Puakō, Hawai'i: methods for assessing and managing marine invasive species. *Marine Ecology Progress Series* 511:209–221 DOI 10.3354/meps10919.
- **Green SJ, Côté IM. 2009.** Record densities of Indo-pacific lionfish on Bahamian coral reefs. *Coral Reefs* **28**(1):107–107 DOI 10.1007/s00338-008-0446-8.
- Green SJ, Dulvy NK, Brooks AM, Akins JL, Cooper AB, Miller S, Côté IM. 2014. Linking removal targets to the ecological effects of invaders: a predictive model and field test. *Ecological Applications* 24(6):1311–1322 DOI 10.1890/13-0979.1.
- **Hilborn R, Walters C. 1992.** *Quantitative fisheries stock assessment: choice, dynamics and uncertainty.* New York: Chapman and Hall.
- Kulbicki M, Beets J, Chabanet P, Cure K, Darling E, Floeter S, Galzin R, Green A, Harmelin-Vivien M, Hixon M, Letourneur Y, De Loma T, McClanahan T, McClanahan T, McClanahan G, Myers R, O'Leary J, Planes S, Vigliola L, Wantiez L. 2012. Distributions of Indo-Pacific lionfishes *Pterois spp* in their native ranges: implications for the Atlantic invasion. *Marine Ecology Progress Series* 446:189–205 DOI 10.3354/meps09442.
- **Layman CA, Jud ZR, Nichols P. 2014.** Lionfish alter benthic invertebrate assemblages in patch habitats of a subtropical estuary. *Marine Biology* **161**:2179–2182 DOI 10.1007/s00227-014-2491-x.
- **Leslie PH, Davis DHS. 1939.** An attempt to determine the absolute number of rats on a given area. *Journal of Animal Ecology* **8**:94–113 DOI 10.2307/1255.

- **Lesser MP, Slattery M. 2011.** Phase shift to algal dominated communities at mesophotic depths associated with lionfish (*Pterois volitans*) invasion on a Bahamian coral reef. *Biological Invasions* **13(8)**:1855–1868 DOI 10.1007/s10530-011-0005-z.
- McClanahan TR, Muthiga NA. 1998. An ecological shift in a remote coral atoll of Belize over 25 years. *Environmental Conservation* 25(2):122–130 DOI 10.1017/S0376892998000174.
- **Morris JA. 2012.** *Invasive lionfish: a guide to control and management.* Marathon: Gulf and Caribbean Fisheries Institute.
- **Morris JA, Akins JL. 2009.** Feeding ecology of invasive lionfish (*Pterois volitans*) in the Bahamian archipelago. *Environmental Biology of Fishes* **86**:389–398 DOI 10.1007/s10641-009-9538-8.
- Morris JA, Shertzer KW, Rice JA. 2010. A stage-based matrix population model of invasive lionfish with implications for control. *Biological Invasions* 13(1):7–12 DOI 10.1007/s10530-010-9786-8.
- Morris JA, Whitfield PE. 2009. Biology, ecology, control and management of the invasive Indo-Pacific lionfish: an updated integrated assessment. North Carolina: NOAA Technical Memorandum NOS NCCOS 99.
- **Mumby PJ. 2006.** Connectivity of reef fish between mangroves and coral reefs: algorithms for the design of marine reserves at seascape scales. *Biological Conservation* **128(2)**:215–222 DOI 10.1016/j.biocon.2005.09.042.
- **Ogle DH. 2016a.** *Introductory fisheries analyses with R.* Boca Raton: CRC Press.
- **Ogle DH. 2016b.** FSA: fisheries stock analysis. *R Package Version 0.8.5*.
- **Pinheiro J, Bates D, Debroy S, Sarkar D. R Core Team. 2013.** nlme: linear and nonlinear mixed effects models. *R Package Version 3.1-111*.
- **R Development Core Team. 2013.** R: a language and environment for statistical computing. Version 3.0.2. Vienna: Foundation for Statistical Computing.
- **Tamburello N, Côté IM. 2015.** Movement ecology of Indo-Pacific lionfish on Caribbean coral reefs and its implications for invasion dynamics. *Biological Invasions* **17(6)**:1639–1653 DOI 10.1007/s10530-014-0822-y.
- Whitfield PE, Hare JA, David AW, Harter SL, Muñoz RC, Addison CM. 2007. Abundance estimates of the Indo-Pacific lionfish *Pterois volitans/miles* complex in the Western North Atlantic. *Biological Invasions* **9**(1):53–64

 DOI 10.1007/s10530-006-9005-9.
- Wickham H. 2009. Ggplot2: elegant graphics for data analysis. New York: Springer.
- **Wild C, Jantzen C, Kremb SG. 2014.** Turf algae-mediated coral damage in coastal reefs of Belize, Central America. *PeerJ* **2**:e571 DOI 10.7717/peerj.571.