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## Spatial and Temporal Variability in Growth of Hawaiian Spiny Lobsters in the Northwestern Hawaiian Islands

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**Abstract.**—Growth rates were estimated for recaptured Hawaiian spiny lobsters *Panulirus marginatus* tagged between 2002 and 2007 at Necker Island (23°30'N; 164°35'W), Gardner Pinnacles (25°00'N; 168°50'W), and Maro Reef (25°30'N; 170°45'W) in the Northwestern Hawaiian Islands (NWHI). The location and year-specific nature of the tag–recapture cruises enabled investigation of spatial and temporal variability in growth. Mean growth rates, estimated by using the von Bertalanffy and Schnute growth models and fitted via a maximum likelihood technique, differed between sexes and particularly among locations and years. Male lobster growth rates at Necker Island were, in general, one-third those at Gardner Pinnacles and one-half those at Maro Reef. Female lobsters exhibited the same pattern with less-pronounced differences. Maro Reef lobsters exhibited several abrupt, significant growth reductions among years, while Necker Island lobster growth rates increased significantly from 2004 to 2005. Model results also indicated substantial individual variability in growth. Neither density nor temperature could account for the differences in growth rates. No palinurid species studied to date has shown such spatial variability in growth, and only one panilurid species has exhibited similar variability on a temporal scale. Recognition of and accounting for the mean and individual variability in a life history trait of this endemic species will result in more accurate stock assessments and, ultimately, a better understanding of lobster dynamics and the NWHI coral reef ecosystem.

Sustainable exploitation of commercially valuable species requires accurate estimates of somatic growth and an in-depth understanding of the causes of any spatiotemporal variability associated with those estimates. Growth rates are important components of yield equations (Beverton and Holt 1957) and are also used in other methods to assess stocks, including mortality (Griffiths and Harrod 2007), maturity schedules (Clark et al. 1999), recruitment or settlement (Wahle et al. 2004), spawner–recruit relationships (Sheehy 2001), and fishing selectivity (Jeffery and Taggart 2000). Failing to account for growth variability in population models introduces biases to biomass estimates, which in turn can lead to inappropriate management schemes and exploitation levels. This may be especially true of palinurid lobsters because many species exhibit spatial and temporal growth variability associated with density (Pollock 1991; McGarvey et al. 1999), environmental conditions (Chittleborough 1975; Pollock and Shannon 1987; Pollock et al. 1997), and food availability (Newman and Pollock 1974; Edgar 1990).

The Hawaiian Archipelago consists of the main Hawaiian Islands (Hawaii to Niihau) and the Northwestern Hawaiian Islands (NWHI), a series of islands,

reefs, seamounts, and atolls (hereafter referred to as banks) extending approximately 2,000 km across the subtropical Pacific (Figure 1). The Hawaiian spiny lobster *Panulirus marginatus* is endemic to the Hawaiian Archipelago and Johnston Atoll. Pre-exploitation surveys (1976–1978) indicated that the average Hawaiian spiny lobster size increased in a northwesterly direction along the NWHI (Uchida et al. 1980). In particular, Necker Island, located near the southeastern end of the NWHI, had the smallest Hawaiian spiny lobsters but also the greatest abundance (Uchida et al. 1980).

A commercial fishery targeting Hawaiian spiny lobsters in the NWHI began in the mid-1970s. Fishing effort remained relatively low until 1984, when the introduction of new traps resulted in a threefold increase in effort in just 2 years. Despite near-annual stock assessments from 1985 to 2000, the NWHI lobster fishery suffered severe declines in catch per unit effort (CPUE; Figure 2) and was closed in 2000 because of increasing uncertainty in the population models used to assess stock status. The uncertainty stems from the assumption of synchronous dynamics, regardless of location and time period, among bank-specific lobster populations (Botsford et al. 2002). For example, previous Hawaiian spiny lobster tagging studies at the NWHI (Kure Atoll in 1979–1983 and French Frigate Shoals in 1981–1983) used Fabens' (1965) method to estimate the conventional von Bertalanffy growth parameters: asymptotic maximum length ( $L_{\infty}$ ) and growth constant ( $K$ ; MacDonald 1984).

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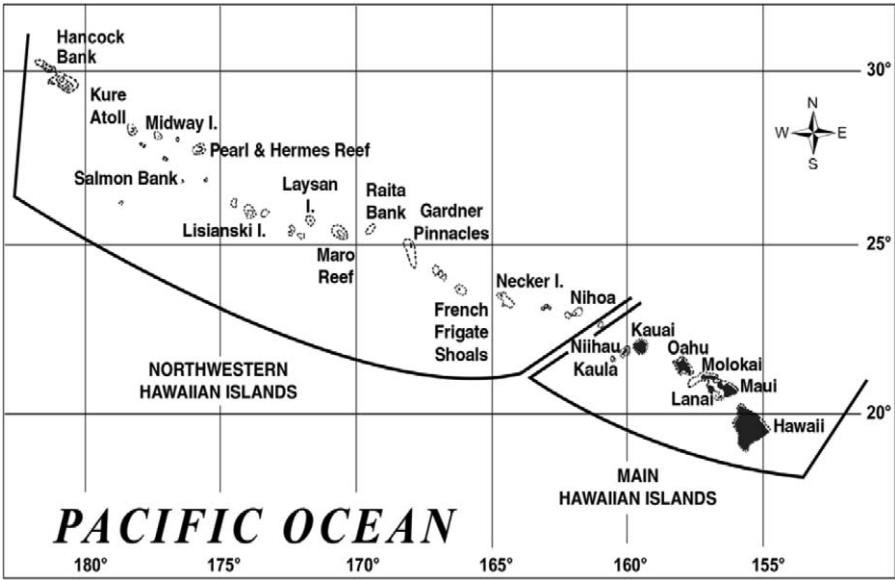


FIGURE 1.—Map of the Hawaiian Archipelago, including the Northwestern Hawaiian Islands.

The growth estimates (male  $K = 0.27$ , female  $K = 0.35$ ) were used in a Beverton–Holt yield equation to estimate Hawaiian spiny lobster abundance at all banks in the NWHI (Polovina 1985). Growth estimates were also derived from modal progression analysis of size frequencies of Hawaiian spiny lobsters captured during research cruises at Necker Island in 1986–1988

(growth rate values were unreported; Polovina and Moffitt 1989) and were used to determine size at recruitment to the fishery and age-specific CPUE in the 1989–1993 assessments. The use of these growth rates in NWHI-wide assessments ignored any spatial and temporal variability that might be associated with those values.

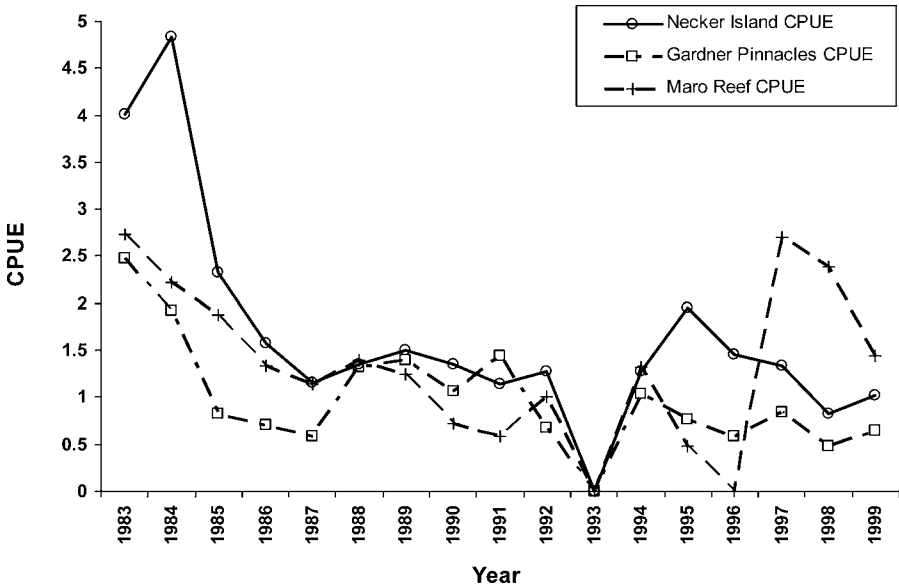


FIGURE 2.—Commercial catch per unit effort (CPUE; number of lobsters captured per trap haul) for Hawaiian spiny lobsters at Necker Island, Gardner Pinnacles, and Maro Reef, Northwestern Hawaiian Islands, from 1983 to 1999. The fishery was temporarily closed in 1993.

Acknowledging and discounting the assumption of synchronous dynamics represented a change in NWHI Hawaiian spiny lobster population dynamics theory from a single archipelago-wide population to a series of discrete metapopulations separated by the deep water between banks and connected only by larval dispersal. To examine bank-specific lobster dynamics, an extensive tagging program was conducted from 2002 to 2008. A preliminary examination of the 2002–2003 Necker Island tagging data indicated that a metapopulation approach is appropriate because the estimated Hawaiian spiny lobster growth differed significantly from that observed at Kure Atoll and French Frigate Shoals from 1979 to 1983 (O'Malley and MacDonald 2009).

The objective of this study is to investigate NWHI Hawaiian spiny lobster metapopulation dynamics by describing location-specific (Necker Island, Gardner Pinnacles, and Maro Reef) and year-specific (2002–2008) growth rates. Also examined are the effects of density and temperature, two common drivers of growth variability.

### Methods

**Tag-recapture data.**—Hawaiian spiny lobster tagging was conducted aboard chartered commercial fishing vessels at the banks where the NWHI commercial fishery was primarily concentrated: Necker Island (23°30'N; 164°35'W); Gardner Pinnacles (25°00'N; 168°50'W); and Maro Reef (25°30'N; 170°45'W; Figure 1). Trapping took place within the entirety of each bank, although effort was concentrated in areas with greater densities of Hawaiian spiny lobsters, as indicated by habitat (depth and sand versus hard bottom) and the captain's previous lobster fishing experiences in the NWHI. Tagging began at Necker Island in 2002, expanded to Maro Reef in 2004, and then extended to Gardner Pinnacles in 2006; tagging took place at all three banks in 2006–2008. The annual tag-recapture cruises occurred between June and September of each year.

Hawaiian spiny lobsters were captured by using standard commercial molded black polyethylene Fathoms Plus (San Diego, California) traps, which are dome-shaped and single-chambered, with two entrance cones (dimensions = 980 × 770 × 295 mm, inside mesh dimensions = 45 × 45 mm). Mesh paneling was placed over the escape vents to prevent the release of small lobsters, thereby maximizing the size range of captured lobsters. Fifteen strings consisting of 20 traps/string were soaked overnight; each trap was baited with 1 kg of Pacific chub mackerel *Scomber japonicus*. Initially (i.e., at Necker Island in 2002), Hawaiian spiny lobsters were tagged by using polyethylene

streamer tags (Hallprint, Inc., Victor Harbor, South Australia) inserted through the abdominal flexor muscle at the juncture of the cephalothorax and abdomen. However, research indicated that these tags were not suitable for Hawaiian spiny lobsters because of excessive tag loss (O'Malley 2008). All subsequent tagging was done by using passive integrated transponder (PIT) tags (Destron Technologies, South St. Paul, Minnesota) injected into the distal tail segment (O'Malley 2008). The PIT tags were detected with a Destron Technologies Model 2001F-ISO portable transceiver (Digital Angel Corporation, St. Paul, Minnesota). For each tagged and recaptured lobster, the PIT tag number, carapace length (CL; nearest 0.01 mm), sex, date of capture, and position of release (latitude and longitude determined using a Global Positioning System unit) were recorded. All lobsters were released on the seafloor in the immediate area of recapture via a release cage (O'Malley 2008). Each tag-recapture cruise had at least one tagger from previous cruises, which provided continuity in the tagging process. An important detail of this work is that because the commercial fishery was closed and because poaching and recreational fishing are highly unlikely given the remoteness and regular monitoring of the NWHI, tagged lobsters were recaptured by researchers during the annual tag-recapture cruises only.

**Growth analysis.**—Hawaiian spiny lobster recapture data were first aggregated on a bank-specific level to examine spatial (among-bank) differences in growth rates (Necker Island, 2002–2008; Gardner Pinnacles, 2006–2008; and Maro Reef, 2004–2008). In these data sets, if individuals were recaptured more than once, only the initial and final capture information was used in the analysis and no immediate recaptures (i.e., lobsters captured during the same tagging cruise) were included. This was done to ensure equal weight in the analysis of any individual's specific growth rate and to maximize the time at liberty. To estimate annual growth and to examine temporal variability, recapture data were partitioned into year-specific data sets within each bank.

Bank-, year-, and sex-specific data were fitted to the von Bertalanffy and Schnute growth models with Francis' (1988a, 1995) maximum likelihood method by using the GROTAG program designed by Simpfendorfer (2000) for the Microsoft Excel solver function (Microsoft Excel version 2002; Microsoft Corp., Redmond, Washington). Both models estimate growth by using the parameters  $g_{\alpha}$  and  $g_{\beta}$ , which represent mean annual growth increments (mm/year) of chosen reference lengths  $\alpha$  and  $\beta$  (Francis 1988a, 1995). Growth rates at these specific sizes are directly

observable and are therefore biologically meaningful. In the case of the von Bertalanffy growth model, these parameters have better statistical properties than  $K$  and  $L_\infty$ , particularly when the entire size range of the species is not represented in the data (Sainsbury 1980; Francis 1988a, 1988b; Haddon 2001). Hawaiian spiny lobster reference lengths  $\alpha$  and  $\beta$  for both models were chosen so that within each data set, the values were well represented while maintaining the majority of individuals between the two values (Francis 1988a). This allowed direct spatial and temporal comparisons within the sexes. The male  $\alpha$  and  $\beta$  were set at 75 and 110 mm CL, respectively, whereas the female  $\alpha$  and  $\beta$  were set at 70 and 100 mm CL, respectively.

The von Bertalanffy growth model was applied by using Francis' (1988a) reparameterization of Fabens' (1965) growth model for tagging data. Following Francis (1988a), the expected length increment ( $\Delta L$ ) for a Hawaiian spiny lobster tagged at length  $L_1$  and at liberty for time  $\Delta T$  is given by:

$$\Delta L = \left[ \frac{\beta g_\alpha - \alpha g_\beta}{g_\alpha - g_\beta} - L_1 \right] \left[ 1 - \left( 1 + \frac{g_\alpha - g_\beta}{\alpha - \beta} \right)^{\Delta T} \right]. \quad (1)$$

To compare Hawaiian spiny lobster growth rates with other palinurid growth estimates, model outputs were converted to the von Bertalanffy growth parameters  $K$  and  $L_\infty$  following Francis (1988a):

$$K = \log_e [1 + (g_\alpha - g_\beta) / (\alpha - \beta)] \quad (2)$$

and

$$L_\infty = (\beta g_\alpha - \alpha g_\beta) / (g_\alpha - g_\beta). \quad (3)$$

Differences in the set reference values of  $\alpha$  and  $\beta$  for males and females prevented direct comparison of growth between the sexes; therefore, female lobster  $g_{75}$  and  $g_{110}$  were estimated by using (Francis 1988a):

$$g_\gamma = [(\gamma - \alpha)g_\beta + (\beta - \gamma)g_\alpha] / (\beta - \alpha). \quad (4)$$

Francis (1995) also created a mark-recapture "equivalent" of Schnute's (1981) size-at-age model by using the same parameters ( $\alpha$ ,  $\beta$ ,  $g_\alpha$ , and  $g_\beta$ ) to describe growth. Following Francis (1995: equation 5.1), the  $\Delta L$  for a Hawaiian spiny lobster tagged at length  $L_1$  and at liberty for  $\Delta T$  is given by:

$$\Delta L = -L_1 + [L_1^b e^{-a\Delta T} + c(1 - e^{-a\Delta T})]^{1/b}, \quad (5)$$

where

$$a = \log_e \left( \frac{\beta^b - \alpha^b}{\lambda_\beta^b - \lambda_\alpha^b} \right) \quad \text{if } b \neq 0, \quad (6)$$

$$c = \frac{\beta^b \lambda_\alpha^b - \alpha^b \lambda_\beta^b}{\lambda_\alpha^b - \alpha^b + \beta^b - \lambda_\beta^b} \quad \text{if } b \neq 0, \quad (7)$$

$$\lambda_\alpha = \alpha + g_\alpha, \quad (8)$$

and

$$\lambda_\beta = \beta + g_\beta. \quad (9)$$

Parameter  $b$ , which is analogous to the parameter  $b$  in Schnute's (1981) model, has no biological meaning and is generally thought of as describing curvature in the model.

Both growth models were fit by using the negative log-likelihood function (Francis 1988a):

$$\lambda = \sum_i \log[(1 - p)\lambda_i + p/R], \quad (10)$$

where

$$\lambda_i = \exp \frac{-1/2(\Delta L_i - \mu_i - m)^2 / (\sigma_i^2 + s^2)}{[2\pi(\sigma_i^2 + s^2)]^{1/2}}, \quad (11)$$

$R$  = the range of observed growth increments,  $\mu$  = the expected growth increment,  $m$  = the mean measurement error,  $s$  = standard deviation of measurement error, and  $\sigma$  = the standard deviation of the growth variability. Also estimated were the coefficient of variation of growth variability ( $v$ ) and outlier contamination ( $p$ ). To describe growth variability,  $\sigma$  was related to  $\mu$  ( $\sigma_i = v\mu_i$ ) by assuming an increase in growth variability as the size of the growth increment increases (Francis 1988a).

The GROTAG methodology cannot accurately determine  $m$  and  $s$  and cannot distinguish between variability in growth and measurement error because individuals with very short times at liberty were removed from the analysis. Therefore,  $m$  and  $s$  were fixed based on independent values estimated from differences in length at release and length at recapture for Hawaiian spiny lobsters that were recaptured within 2 d of tagging ( $n = 206$ ). The mean and standard deviation of these differences represent an exact measure of measurement error, assuming that no growth occurred between tagging and recapture. It is likely that researchers would detect a Hawaiian spiny lobster that molted within this time period (clean carapace, soft shell). The estimated mean and standard deviation of the differences in size were 0.13 and 0.12 mm, respectively, for males and were 0.16 and 0.17 mm, respectively, for females.

For each data set (bank, years at liberty, sex), the first model was the simplest, fitted with only  $g_\alpha$ ,  $g_\beta$ ,  $s$ , and in the case of Schnute model,  $b$ . Each subsequent model introduced an additional parameter. Likelihood

TABLE 1.—Tag–recapture data for Hawaiian spiny lobsters used in growth analysis by bank and years at liberty (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $n$  = sample size; DAL = days at liberty).

Bank	Year tagged– year recaptured	$n$ tagged	$n$ recaptured	DAL	
				Mean	SD
NI	2002–2003	13,944	472	360	11
	2003–2004	11,917	901	364	8
	2004–2005	5,990	629	318	1
	2005–2006	5,177	104	334	8
	2006–2007	1,605	165	373	1
	2007–2008	1,361	85	426	1
GP	2002–2008	39,994	3,158	609	359
	2006–2007	673	43	373	1
	2007–2008	1,146	37	429	4
	2006–2008	1,819	90	501	178
MR	2004–2005	1,911	80	316	4
	2005–2006	2,792	39	374	6
	2006–2007	1,663	105	370	2
	2007–2008	2,648	123	377	4
	2004–2008	9,014	415	503	243

ratio tests were used to determine the final model, where for a significant ( $P < 0.05$ ) improvement in fit, the likelihood value must increase by at least 1.92 with the introduction of one parameter and by at least 3.0 with the introduction of two parameters (Francis 1988a). The 95% confidence intervals (CIs) were estimated by using a bootstrapping method as implemented in GROTAG (Simpfendorfer 2000). Significant differences between data sets were identified by comparing 95% CIs and the extent of bootstrap overlap (Welsford and Lyle 2005). Comparison of CIs has been shown to be a conservative yet effective measure of significance (Schenker and Gentleman 2001).

The von Bertalanffy and Schnute growth models were evaluated by using the second-order bias-corrected Akaike's information criterion ( $AIC_c$ ). For each data set, the model with the lowest  $AIC_c$  value was considered the model that provided the best fit to the data.

To examine the effects of density on Hawaiian spiny lobster growth, individual annual growth rates were regressed on CPUE (i.e., lobsters captured per hauled

trap) at Necker Island and Maro Reef, the banks with the longest time series of tag–recapture data. Trap CPUE has been shown to be an adequate proxy for density (MacDonald and Stimson 1980; Polovina 1989); therefore, string-specific trap CPUE (lobsters caught in the same string per hauled trap) was calculated for each recaptured Hawaiian spiny lobster at Necker Island (2002–2003 to 2007–2008) and Maro Reef (2004–2005 to 2007–2008). The regression included release size to account for the effect of size on growth:

$$\Delta CL / \Delta t = B_0 + B_1(\text{release size}) + B_2(\text{string-specific trap CPUE}) + [B_1(\text{release size}) \times B_2(\text{string-specific trap CPUE})] + \varepsilon, \quad (12)$$

where  $\varepsilon$  is the error term.

To test the influence of temperature on Hawaiian spiny lobster growth, the 2004–2005 to 2007–2008 Necker Island and Maro Reef estimated growth rates ( $g_x$ ,  $g_y$ ) were regressed on sea surface temperature (SST) data. The SST data, collected from the Geostationary Operational Environmental Satellite, were acquired from the National Oceanic and Atmospheric Administration's Oceanwatch Central Pacific website (<http://oceanwatch.pifsc.noaa.gov/index.html>). The annual mean SST at a given bank and period was calculated by using the start date of the tagging cruise year and the end date of the following year's cruise. The regression included the additional sources of variation to account for their effects on growth:

$$\text{estimated growth rate} = B_0 + B_1(\text{year}) + B_2(\text{bank}) + B_3(\text{sex}) + B_4(\text{SST}) + \varepsilon. \quad (13)$$

## Results

### Tag–Recapture Data

In total, 50,827 Hawaiian spiny lobsters were tagged at Necker Island, Gardner Pinnacles, and Maro Reef

TABLE 2.—Hawaiian spiny lobster catch, effort (number of traps hauled), and catch per unit effort (CPUE; lobsters/trap) during tagging cruises by year and location in the Northwestern Hawaiian Islands.

Year	Necker Island			Gardner Pinnacles			Maro Reef		
	Catch	Effort	CPUE	Catch	Effort	CPUE	Catch	Effort	CPUE
2002	13,950	15,000	0.93	—	—	—	—	—	—
2003	11,917	9,400	1.27	—	—	—	—	—	—
2004	7,019	6,900	1.02	—	—	—	1,911	6,000	0.32
2005	6,619	7,660	0.86	—	—	—	2,888	6,300	0.46
2006	2,049	3,400	0.60	703	3,400	0.21	1,764	2,900	0.61
2007	1,845	3,300	0.56	1,222	3,300	0.37	2,834	3,900	0.73
2008	1,441	1,800	0.80	514	1,800	0.29	1,381	1,800	0.77

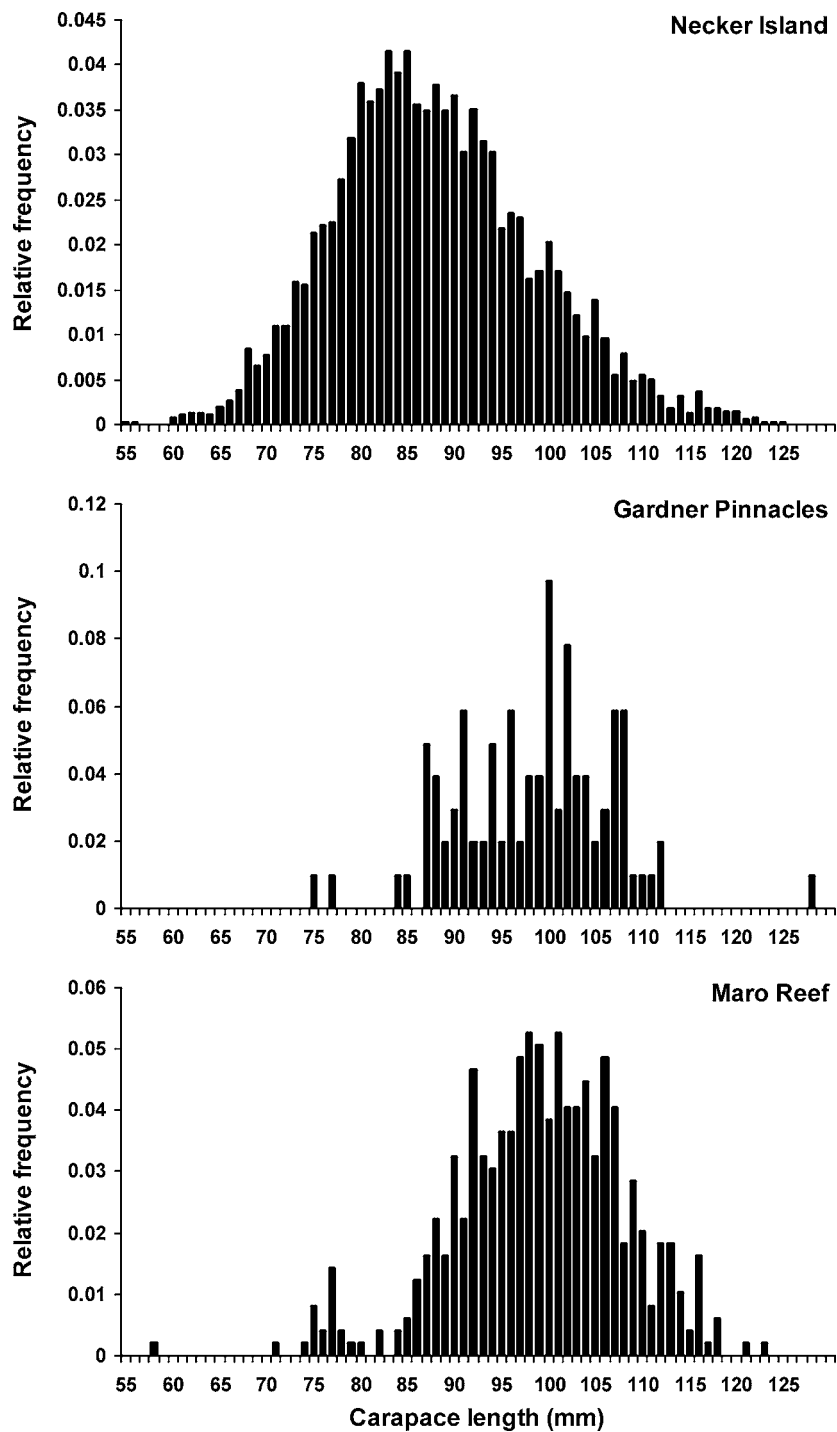


FIGURE 3.—Size-frequency distribution (carapace length, mm) of recaptured Hawaiian spiny lobsters at Necker Island (2003–2008), Gardner Pinnacles (2007 and 2008), and Maro Reef (2005–2008), Northwestern Hawaiian Islands.

TABLE 3.—Negative log-likelihood values for differently parameterized von Bertalanffy growth models (Francis 1988a method) used in selection of the optimal model of male and female Hawaiian spiny lobster growth by bank and year (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $g_\alpha$  and  $g_\beta$  = mean annual growth increments [mm/year] of chosen reference lengths  $\alpha$  and  $\beta$ ;  $s$  = SD of measurement error;  $v$  = coefficient of variation of growth variability;  $m$  = mean measurement error; and  $p$  = outlier contamination). Values with an asterisk indicate final model selected. Male  $s$  and  $m$  were fixed at 0.12 and 0.13, respectively; female  $s$  and  $m$  were fixed at 0.17 and 0.16, respectively.

Bank	Year tagged– year recaptured	Model							
		$g_\alpha, g_\beta, s$		$g_\alpha, g_\beta, s, v$		$g_\alpha, g_\beta, s, v, m$		$g_\alpha, g_\beta, s, v, m, p$	
		Male	Female	Male	Female	Male	Female	Male	Female
NI	2002–2003	2,155.2	1,580.1	399.2*	310.2*	399.2	309.1	399.2	309.1
	2003–2004	4,211.2	6,645.0	930.2*	516.6*	929.8	516.6	929.8	516.6
	2004–2005	2,444.9	3,042.9	504.2*	574.7*	503.9	574.5	503.9	574.5
	2005–2006	324.2	403.7	77.0*	89.2*	77.1	89.5	77.0	93.7
	2006–2007	469.5	412.3	131.5*	107.0*	131.5	107.1	131.1	107.1
GP	2007–2008	248.8	781.9	81.8*	61.9*	81.8	61.8	78.9	61.8
	2002–2008	14,009.1	12,871.2	3,457.5*	2,838.2*	3,461.0	2,840.0	3,461.0	2,840.0
	2006–2007	129.2	289.7	42.8*	66.3*	42.7	66.1	42.7	66.1
	2007–2008	52.7	310.2	32.6*	52.0*	32.6	52.2	32.6	52.2
	2006–2008	225.2	754.5	93.1*	141.9*	93.1	141.7	93.1	141.7
MR	2004–2005	271.4	177.9	105.9*	80.5*	106.1	80.5	106.1	80.5
	2005–2006	123.9	185.4	50.7*	46.7*	50.7	46.6	50.7	46.6
	2006–2007	340.1	607.5	101.2*	134.5*	101.2	134.6	101.2	133.8
	2007–2008	203.4	665.9	116.2*	136.8*	116.3	136.4	116.2	137.5
	2004–2008	1,777.2	2,699.9	519.6*	586.7*	520.0	587.4	520.0	619.1

between 2002 and 2007 (Table 1). From 2003 to 2008, 5,052 Hawaiian spiny lobsters were recaptured across all banks; of these recaptures, 2,315 occurred during the next tagging cruise (i.e., lobsters were at liberty for approximately 1 year; Table 1). The variability in the numbers tagged reflects changes in trapping effort when tagging was expanded to different banks rather

than large changes in abundance, as evident by CPUE (Table 2). The variability in days at liberty was a result of the research cruise scheduling.

Plots of bank-specific size frequency distributions of recaptured Hawaiian spiny lobsters displayed differences in the population size structure among banks (Figure 3). Mean ( $\pm$ SD) carapace sizes of recaptured

TABLE 4.—Negative log-likelihood values for differently parameterized Schnute growth models (Francis 1995 method) used in selection of the optimal model of male and female Hawaiian spiny lobster growth by bank and year (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $g_\alpha$  and  $g_\beta$  = mean annual growth increments [mm/year] of chosen reference lengths  $\alpha$  and  $\beta$ ;  $s$  = SD of measurement error;  $v$  = coefficient of variation of growth variability;  $m$  = mean measurement error;  $p$  = outlier contamination; and  $b$  = curvature). Values with an asterisk indicate final model selected. Male  $s$  and  $m$  were fixed at 0.12 and 0.13, respectively; female  $s$  and  $m$  were fixed at 0.17 and 0.16, respectively.

Bank	Year tagged– year recaptured	Model							
		$g_\alpha, g_\beta, s, b$		$g_\alpha, g_\beta, s, v, b$		$g_\alpha, g_\beta, s, v, m, b$		$g_\alpha, g_\beta, s, v, m, p, b$	
		Male	Female	Male	Female	Male	Female	Male	Female
NI	2002–2003	2,151.0	1,578.0	399.2*	300.4*	398.9	299.3	398.7	446.8
	2003–2004	4,996.0	6,354.0	927.8*	515.7*	927.7	515.7	927.7	515.7
	2004–2005	7,025.0	716.4	493.6*	570.1*	493.4	570.2	493.4	570.2
	2005–2006	999.5	1,001.0	76.1*	87.6*	76.1	89.8	76.1	118.4
	2006–2007	1,424.0	981.0	130.4*	106.7*	130.5	106.8	129.8	106.8
GP	2007–2008	770.1	759.3	80.3*	61.7*	80.2	61.7	78.1	62.9
	2002–2008	44,314.0	12,682.0	3,440.0*	2,827.0*	3,444.0	2,830.0	3,444.0	2,830.0
	2006–2007	317.4	857.1	41.4*	65.3*	41.3	65.2	41.3	65.2
	2007–2008	134.2	827.7	31.6*	51.2*	31.6	51.5	31.6	51.5
	2006–2008	533.5	2,003.0	86.3*	141.0*	86.3	141.2	86.5	141.1
MR	2004–2005	703.4	485.9	102.5*	79.9*	102.8	79.9	102.8	79.9
	2005–2006	398.9	529.5	50.6*	46.5*	50.6	46.5	50.6	46.5
	2006–2007	1,019.0	604.0	98.6*	133.8*	98.6	134.2	98.6	133.2
	2007–2008	967.9	1,744.8	109.5*	136.3*	109.5	136.0	109.5	169.8
	2004–2008	1,513.0	5,138.0	495.2*	509.9*	497.2	512.8	497.6	512.3



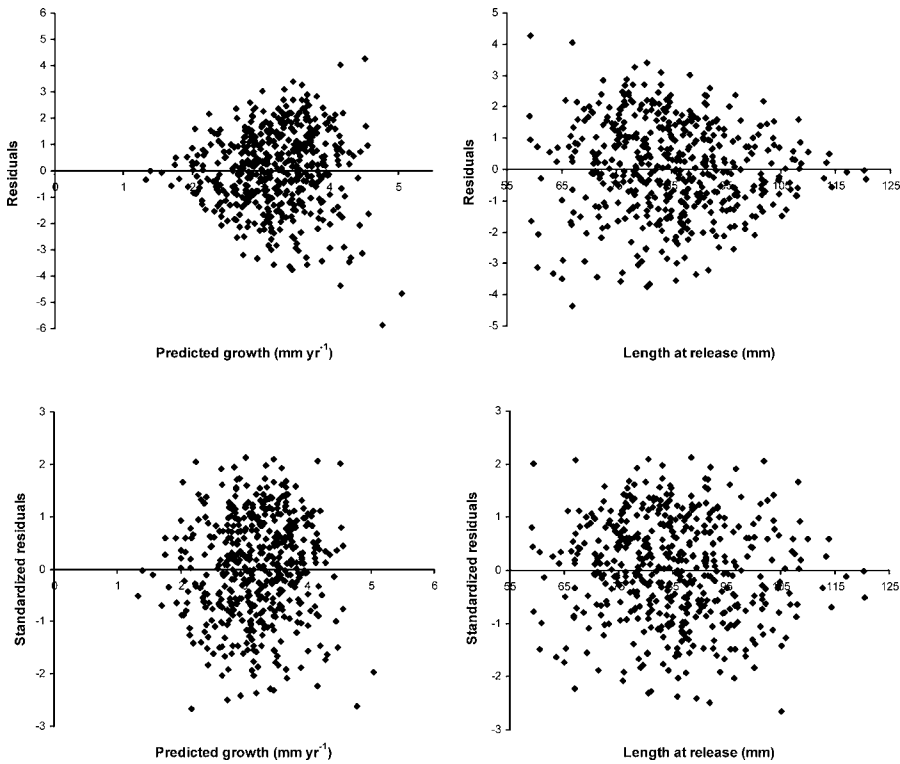


FIGURE 4.—Plots of von Bertalanffy growth model residuals (top panels) and standardized residuals (bottom panels) against predicted growth (left panels; mm carapace length [CL]/year) and length at release (right panels; CL, mm) for male Hawaiian spiny lobsters at Necker Island, Northwestern Hawaiian Islands, 2003–2004.

lobsters were  $87.37 \pm 10.56$  mm CL at Necker Island,  $98.93 \pm 8.16$  mm CL at Gardner Pinnacles, and  $99.52 \pm 8.98$  mm CL at Maro Reef. Using a Kruskal–Wallis test, a significant difference in size distributions was found among banks ( $\chi^2 = 617.13$ ,  $df = 2$ ,  $P < 0.0001$ ). Specifically, Kolmogorov–Smirnov tests indicated significant differences in size distributions between Necker Island and Gardner Pinnacles (Kolmogorov–Smirnov  $D$ -statistic = 0.49,  $P < 0.0001$ ) and between Necker Island and Maro Reef ( $D = 0.53$ ,  $P < 0.0001$ ) but not between Gardner Pinnacles and Maro Reef ( $D = 0.08$ ,  $P = 0.69$ ). The size structure differences between Necker Island and the other banks were probably not due to sampling issues because the same trapping gear and sampling protocol were used during each tagging cruise.

Growth Model Results

The models containing  $g_\alpha$ ,  $g_\beta$ ,  $s$ , and  $v$  resulted in the best fit to the von Bertalanffy growth model for all data sets (Table 3), and  $g_\alpha$ ,  $g_\beta$ ,  $s$ ,  $v$ , and  $b$  resulted in the best fit to the Schnute growth model (Table 4); introducing additional parameters did not result in a significant

improvement in fit, as evident in the likelihood ratio tests.

To assess model fits, residuals and standardized residuals were plotted against length at release and predicted growth; a representative data set of the von Bertalanffy model (Necker Island 2003–2004 males) is shown in Figure 4. Both growth models displayed declining residuals with increasing length at release because mean growth declines with length (McGarvey et al. 1999), while residuals versus predicted growth showed the opposite trend, as would be expected. Standardized residuals (residuals divided by  $\sigma_i$ , which, in the selected model, equals  $s$ ) showed no pattern, indicating that the model assumption that growth variability is dependent on mean growth was not violated (Francis 1988b). Residuals of all bank-, year-, and sex-specific data sets were generally symmetric around a mean of zero; the von Bertalanffy model residuals are shown in Figure 5. Overall, residual plots indicated satisfactory fits such that both models were suitable for the NWHI Hawaiian spiny lobster tagging data.

Both growth models generated similar estimates of

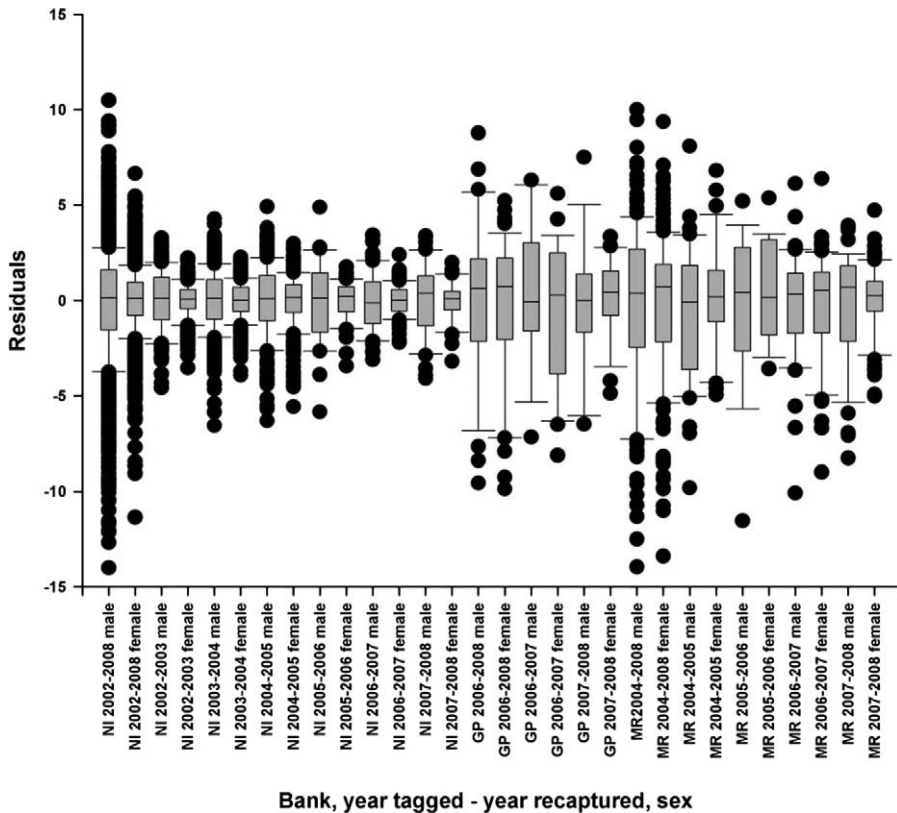


FIGURE 5.—Box-and-whisker plots of von Bertalanffy growth model residuals for Hawaiian spiny lobsters by bank (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef), study period (year tagged–year recaptured), and sex.

$g_{\alpha}$ ,  $g_{\beta}$ , and  $v$  (Tables 5, 6). However, the von Bertalanffy growth model had lower  $AIC_c$  values relative to the Schnute growth model in 67% of the male data sets (Necker Island: 2002–2003, 2003–2004, 2005–2006, 2006–2007, 2007–2008; Gardner Pinnacles: 2006–2007, 2007–2008; Maro Reef: 2005–2006, 2006–2007) and 73% of the female data sets (Necker Island: 2003–2004, 2005–2006, 2006–2007, 2007–2008; Gardner Pinnacles: 2006–2007, 2007–2008, 2006–2008; Maro Reef: 2004–2005, 2005–2006, 2006–2007, 2007–2008) and is therefore considered the better model. All reported growth estimates are from the von Bertalanffy model.

#### *Individual Variability and Mean Growth Rates*

Hawaiian spiny lobsters exhibited substantial individual variability in growth, with the estimated  $v$  for all data sets ranging from 0.24 to 0.71 (Table 5). Estimates of  $g_{75}$  and  $g_{110}$  indicated that Hawaiian spiny lobsters displayed sexual dimorphism in growth, with males growing faster than females (Table 7). The estimated Hawaiian spiny lobster growth rates also indicated that

male and female growth was consistently slowest at Necker Island (Table 7). In years when tagging took place at all three banks (2006–2007 and 2007–2008), Necker Island 75- and 110-mm CL males grew three times slower than those at Gardner Pinnacles and two times slower than those at Maro Reef. Female Hawaiian spiny lobsters exhibited the same pattern, but the differences between banks were slightly less pronounced. During the same time period, male and female Hawaiian spiny lobsters grew slightly faster at Gardner Pinnacles than at Maro Reef except for female  $g_{110}$  in 2007–2008. The slower growth trend of Necker Island lobsters was further confirmed by comparing growth rates between Necker Island and Maro Reef lobsters during the extended time series of 2004–2005 to 2007–2008. Plots of bootstrap growth estimates (Figure 6) and the nonoverlapping 95% CIs (Table 5) indicate that the differences between Necker Island and the other banks were significant. Although smaller, the difference in growth between Gardner Pinnacles and Maro Reef lobsters was also significant.

Temporal variability in Hawaiian spiny lobster

TABLE 5.—Final von Bertalanffy model (Francis 1988a method) estimated growth parameters (95% confidence intervals in parentheses) for male and female Hawaiian spiny lobsters (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $g_{\alpha}$  = mean annual growth of  $\alpha$ , where  $\alpha = 75$  and 70 mm carapace length [CL] for males and females, respectively;  $g_{\beta}$  = mean annual growth of  $\beta$ , where  $\beta = 110$  and 100 mm CL for males and females, respectively;  $v$  = coefficient of variation of growth variability;  $K$  = von Bertalanffy growth constant; and  $L_{\infty}$  = asymptotic maximum length). The SD of measurement error ( $s$ ) included in the final model was fixed at 0.17 and 0.12 for males and females, respectively.

Bank	Year tagged– year recaptured	$g_{\alpha}$ , mm/year		$g_{\beta}$ , mm/year	
		Male	Female	Male	Female
NI	2002–2003	3.70 (3.36–4.08)	2.28 (2.07–2.50)	1.78 (1.51–2.02)	0.56 (0.45–0.66)
	2003–2004	3.71 (3.53–3.89)	2.29 (2.15–2.44)	1.93 (1.72–2.11)	0.62 (0.50–0.73)
	2004–2005	4.75 (4.42–5.08)	3.18 (2.96–3.41)	1.57 (1.32–1.82)	0.91 (0.75–1.08)
	2005–2006	5.69 (4.70–6.71)	2.75 (2.41–3.16)	2.37 (1.51–3.17)	0.74 (0.36–0.91)
	2006–2007	4.60 (4.00–5.23)	2.48 (2.20–2.78)	2.31 (1.79–2.93)	0.51 (0.32–0.70)
	2007–2008	5.35 (4.61–6.05)	2.68 (2.25–3.12)	2.07 (1.25–2.88)	0.51 (0.25–0.78)
GP	2002–2008	4.42 (4.31–4.54)	2.51 (2.43–2.59)	1.81 (1.71–1.90)	0.71 (0.66–0.76)
	2006–2007	12.96 (10.83–15.58)	10.46 (8.38–12.79)	5.66 (4.48–6.85)	3.39 (2.71–4.02)
	2007–2008	14.86 (12.46–17.69)	8.65 (6.06–12.14)	1.71 (0.24–3.36)	2.21 (1.68–2.90)
	2006–2008	12.46 (10.88–14.31)	9.01 (7.39–10.95)	5.29 (4.24–6.33)	2.75 (2.29–3.31)
MR	2004–2005	16.22 (14.38–18.09)	15.06 (13.56–16.74)	4.02 (3.37–4.65)	1.9 (1.56–2.26)
	2005–2006	16.53 (13.38–20.17)	12.16 (9.00–15.45)	2.93 (1.44–4.07)	3.07 (2.40–3.76)
	2006–2007	10.45 (8.62–12.40)	10.48 (9.03–12.44)	3.38 (2.72–3.94)	2.4 (2.08–2.76)
	2007–2008	10.65 (8.95–12.32)	7.17 (6.13–8.36)	3.14 (2.54–3.75)	2.22 (1.96–2.49)
	2004–2008	12.36 (11.28–13.50)	8.43 (7.69–9.19)	3.53 (3.2–3.88)	3.05 (2.78–3.28)

growth was also evident within banks (Table 5). In particular, females at Maro Reef experienced a 19% decline in growth between lobsters at liberty from 2004–2005 to 2005–2006, a 14% decline in growth from 2005–2006 to 2006–2007, and a 35% decline in growth from 2006–2007 to 2007–2008 (Table 5). Plots of bootstrap growth estimates (Figure 7) and the nonoverlapping 95% CIs indicated that differences in the first and last intervals were significant (Table 5). Males at Maro Reef experienced a

significant 37% decline in growth from 2005–2006 to 2006–2007 (Table 5; Figure 7). Hawaiian spiny lobsters at Necker Island did not experience a significant change in growth during the same time periods, but males displayed a 28% increase in growth and females displayed a 39% increase in growth from 2003–2004 to 2004–2005 (Table 5). The only significant change in Gardner Pinnacle lobster growth was a decrease in male  $g_{110}$  from 2006–2007 to 2007–2008 (Table 5).

TABLE 6.—Final Schnute model (Francis 1995 method) estimated growth parameters (95% confidence intervals in parentheses) for male and female Hawaiian spiny lobsters (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $g_{\alpha}$  = mean annual growth of  $\alpha$ , where  $\alpha = 75$  and 70 mm carapace length [CL] for males and females, respectively;  $g_{\beta}$  = mean annual growth of  $\beta$ , where  $\beta = 110$  and 100 mm CL for males and females, respectively;  $v$  = coefficient of variation of growth variability; and  $b$  = curvature). The SD of measurement error ( $s$ ) included in the final model was fixed at 0.17 and 0.12 for males and females, respectively.

Bank	Year tagged– year recaptured	$g_{\alpha}$ , mm/year		$g_{\beta}$ , mm/year	
		Male	Female	Male	Female
NI	2002–2003	3.60 (3.12–4.05)	3.04 (2.59–3.53)	1.77 (1.48–2.04)	0.61 (0.48–0.73)
	2003–2004	3.68 (3.52–3.85)	2.34 (2.16–2.51)	2.14 (1.90–2.38)	0.66 (0.52–0.78)
	2004–2005	4.72 (4.37–5.05)	3.24 (3.00–3.51)	2.0 (1.63–2.35)	1.15 (0.93–1.38)
	2005–2006	5.21 (4.15–6.28)	2.99 (2.47–3.44)	3.40 (2.00–4.63)	0.71 (0.48–0.94)
	2006–2007	4.67 (3.98–5.29)	2.36 (1.97–2.73)	2.60 (1.67–3.25)	0.49 (0.28–0.68)
	2007–2008	4.99 (4.35–5.79)	2.77 (2.20–3.27)	1.91 (1.20–2.52)	0.54 (0.26–0.87)
GP	2002–2008	3.60 (3.12–4.05)	3.04 (2.59–3.53)	1.77 (1.48–2.04)	0.61 (0.48–0.73)
	2006–2007	14.46 (10.56–18.43)	11.02 (6.96–15.29)	4.80 (3.26–6.53)	2.95 (2.14–3.92)
	2007–2008	15.55 (12.70–19.71)	9.92 (3.25–20.42)	2.95 (–0.27–5.06)	2.06 (1.48–2.95)
	2006–2008	14.05 (11.89–16.59)	9.19 (6.37–12.54)	4.30 (3.43–5.37)	2.65 (2.07–3.22)
MR	2004–2005	17.61 (15.36–19.90)	14.44 (12.64–16.42)	3.42 (2.83–3.92)	1.89 (1.56–2.25)
	2005–2006	16.81 (12.87–21.26)	13.12 (6.90–20.61)	3.19 (1.57–4.65)	2.98 (2.18–3.91)
	2006–2007	13.38 (10.21–16.83)	11.04 (8.32–14.64)	3.10 (2.47–3.71)	2.21 (1.82–2.67)
	2007–2008	14.19 (10.76–17.46)	7.94 (5.38–10.70)	3.13 (2.56–3.73)	2.16 (1.87–2.48)
	2004–2008	14.58 (13.43–16.39)	15.12 (13.09–17.12)	3.17 (2.85–3.48)	5.96 (5.53–6.49)

TABLE 5.—Extended.

Bank	Year tagged– year recaptured	$v$		$K$		$L_{\infty}$ , mm	
		Male	Female	Male	Female	Male	Female
NI	2002–2003	0.54 (0.48–0.60)	0.57 (0.50–0.64)	0.06	0.06	142.42	109.70
	2003–2004	0.47 (0.43–0.50)	0.54 (0.49–0.59)	0.05	0.06	147.75	111.09
	2004–2005	0.50 (0.45–0.56)	0.57 (0.51–0.63)	0.10	0.08	127.33	112.00
	2005–2006	0.48 (0.34–0.62)	0.51 (0.39–0.62)	0.10	0.07	135.04	110.98
	2006–2007	0.40 (0.32–0.47)	0.45 (0.37–0.54)	0.07	0.07	145.40	107.79
	2007–2008	0.37 (0.28–0.47)	0.45 (0.32–0.56)	0.10	0.07	132.06	107.10
	2002–2008	0.48 (0.46–0.49)	0.53 (0.51–0.56)	0.08	0.06	134.22	111.89
GP	2006–2007	0.37 (0.22–0.49)	0.54 (0.36–0.69)	0.23	0.27	137.16	114.35
	2007–2008	0.24 (0.13–0.29)	0.71 (0.45–0.93)	0.49	0.24	113.41	110.33
	2006–2008	0.36 (0.26–0.46)	0.67 (0.52–0.85)	0.23	0.23	135.85	113.20
MR	2004–2005	0.37 (0.28–0.46)	0.32 (0.23–0.39)	0.43	0.58	121.52	104.33
	2005–2006	0.36 (0.21–0.47)	0.49 (0.28–0.65)	0.49	0.36	117.55	110.13
	2006–2007	0.42 (0.32–0.52)	0.56 (0.43–0.68)	0.23	0.31	126.76	108.94
	2007–2008	0.41 (0.31–0.51)	0.54 (0.43–0.63)	0.24	0.18	124.67	113.45
	2004–2008	0.46 (0.40–0.52)	0.60 (0.53–0.67)	0.29	0.20	124.00	117.03

*Density and Sea Surface Temperature Effects*

Examination of density effects on Hawaiian spiny lobster growth by using individual growth rates was valid because of the near-equal time at liberty for all individuals at a given bank in a specific year; this removes the effect of time at liberty when comparing growth rates relative to density. String-specific trap CPUE ranged from 0.05 to 3.55 lobsters/trap at Necker Island and from 0.05 to 2.80 lobsters/trap at Maro Reef. At both banks and in all years except for

2 years at Necker Island (2005–2006, 2006–2007), there was a significant effect of release size on Hawaiian spiny lobster growth (Table 8). After correcting for release size, density was suggested to have a negative effect on growth at Necker Island in only one of the 6 years examined and no effect in any year at Maro Reef. The regression analysis indicated no effect of SST on growth of 75-mm ( $P = 0.30$ ) and 110-mm CL ( $P = 0.17$ ) Hawaiian spiny lobsters at Necker Island and Maro Reef from 2005

TABLE 6.—Extended.

Bank	Year tagged– year recaptured	$v$		$b$	
		Male	Female	Male	Female
NI	2002–2003	0.53 (0.47–0.60)	0.56 (0.49–0.62)	0.26 (–2.12–2.24)	5.73 (3.44–7.77)
	2003–2004	0.47 (0.43–0.50)	0.54 (0.50–0.59)	2.16 (0.97–3.12)	1.83 (0.30–3.06)
	2004–2005	0.49 (0.44–0.54)	0.56 (0.51–0.61)	4.11 (2.69–5.21)	3.28 (1.65–4.74)
	2005–2006	0.47 (0.33–0.59)	0.50 (0.39–0.59)	5.11 (0.78–9.64)	2.93 (0.28–5.09)
	2006–2007	0.40 (0.31–0.46)	0.45 (0.37–0.52)	3.08 (–0.99–5.64)	–0.05 (–3.22–2.38)
	2007–2008	0.36 (0.26–0.43)	0.44 (0.33–0.55)	–2.34 (–7.70–0.96)	1.89 (–1.73–4.75)
	2002–2008	0.54 (0.47–0.60)	0.56 (0.49–0.62)	0.26 (–2.12–2.24)	5.73 (3.44–7.77)
GP	2006–2007	0.34 (3.26–6.53)	0.56 (0.36–0.72)	3.88 (–0.72–8.66)	2.66 (–2.24–7.45)
	2007–2008	0.23 (0.11–0.30)	0.72 (0.42–1.05)	5.12 (–1.92–17.36)	2.93 (–8.75–12.75)
	2006–2008	0.30 (0.21–0.36)	0.72 (0.53–0.91)	4.67 (1.51–7.11)	1.50 (–1.71–4.09)
MR	2004–2005	0.34 (0.25–0.42)	0.31 (0.23–0.38)	5.35 (2.70–9.17)	–0.75 (–3.97–1.63)
	2005–2006	0.35 (0.22–0.48)	0.48 (0.28–0.64)	2.46 (–3.65–9.52)	3.47 (–5.09–11.68)
	2006–2007	0.40 (0.29–0.48)	0.58 (0.45–0.70)	4.72 (0.74–8.82)	2.30 (–1.25–5.78)
	2007–2008	0.36 (0.28–0.44)	0.53 (0.41–0.64)	7.17 (2.91–12.26)	2.12 (–1.62–4.83)
	2004–2008	0.41 (0.36–0.46)	0.45 (0.39–0.50)	5.37 (3.70–7.29)	2.07 (0.38–3.44)

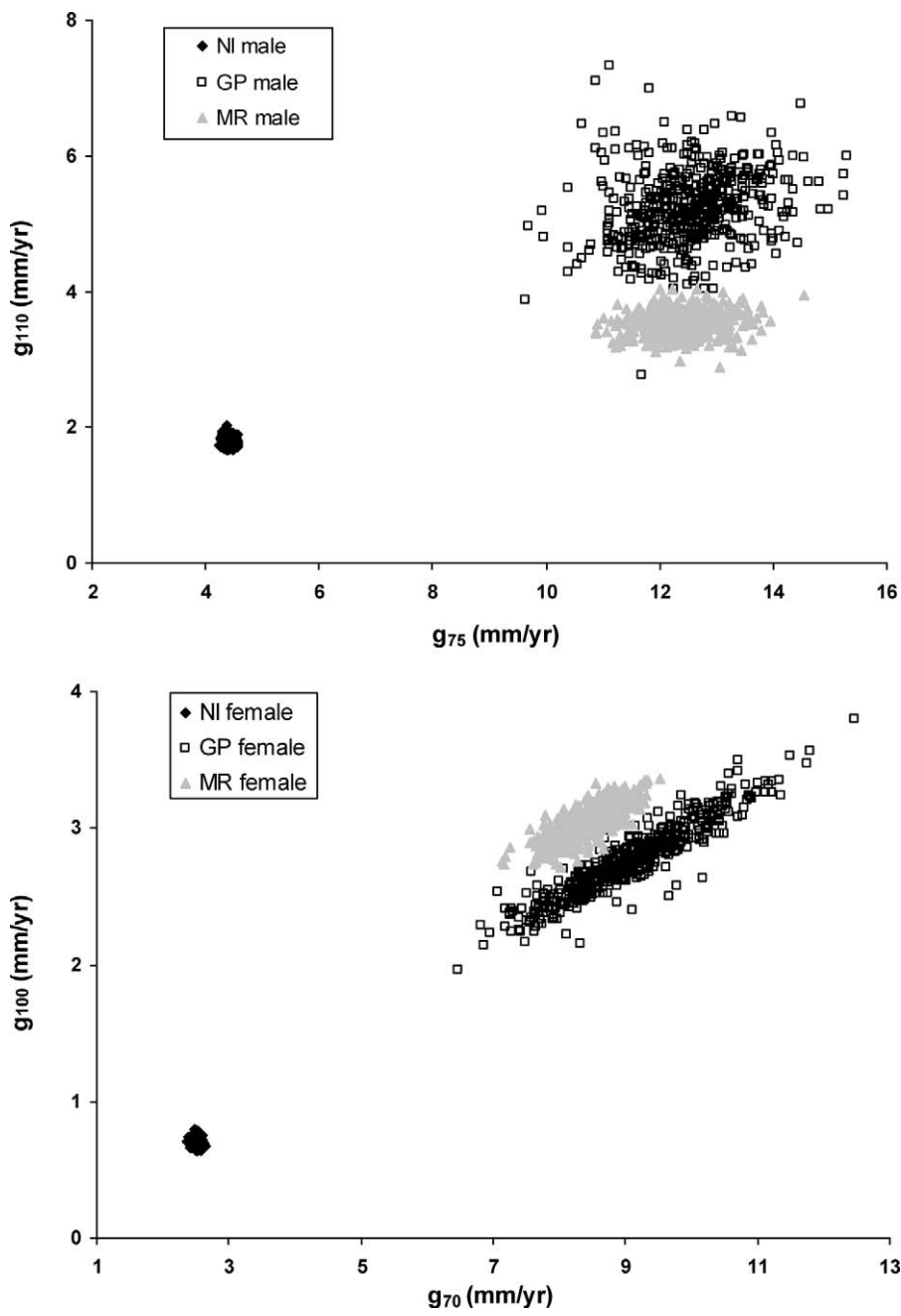


FIGURE 6.—von Bertalanffy growth model bootstrap estimates of growth ( $g$ ; mm carapace length [CL]/year) for male (top; reference lengths = 75 and 110 mm CL) and female (bottom; reference lengths = 70 and 100 mm CL) Hawaiian spiny lobsters by bank for all years (NI = Necker Island, 2002–2008; GP = Gardner Pinnacles, 2006–2008; MR = Maro Reef, 2004–2008).

to 2008. The analysis also indicated significant effects of bank and sex on the growth rates of 75-mm (bank:  $P < 0.001$ ; sex:  $P = 0.004$ ) and 110-mm CL (bank:  $P = 0.04$ ; sex:  $P < 0.01$ ) lobsters, providing additional evidence of spatial and sexual

variability in Hawaiian spiny lobster growth. Mean monthly averages of SST at Necker Island (mean  $\pm$  SD =  $25.6 \pm 1.40^{\circ}\text{C}$ ) and Maro Reef ( $25.4 \pm 1.83^{\circ}\text{C}$ ) were not significantly different from 2005 to 2008 (paired  $t$ -test:  $P = 0.21$ ).

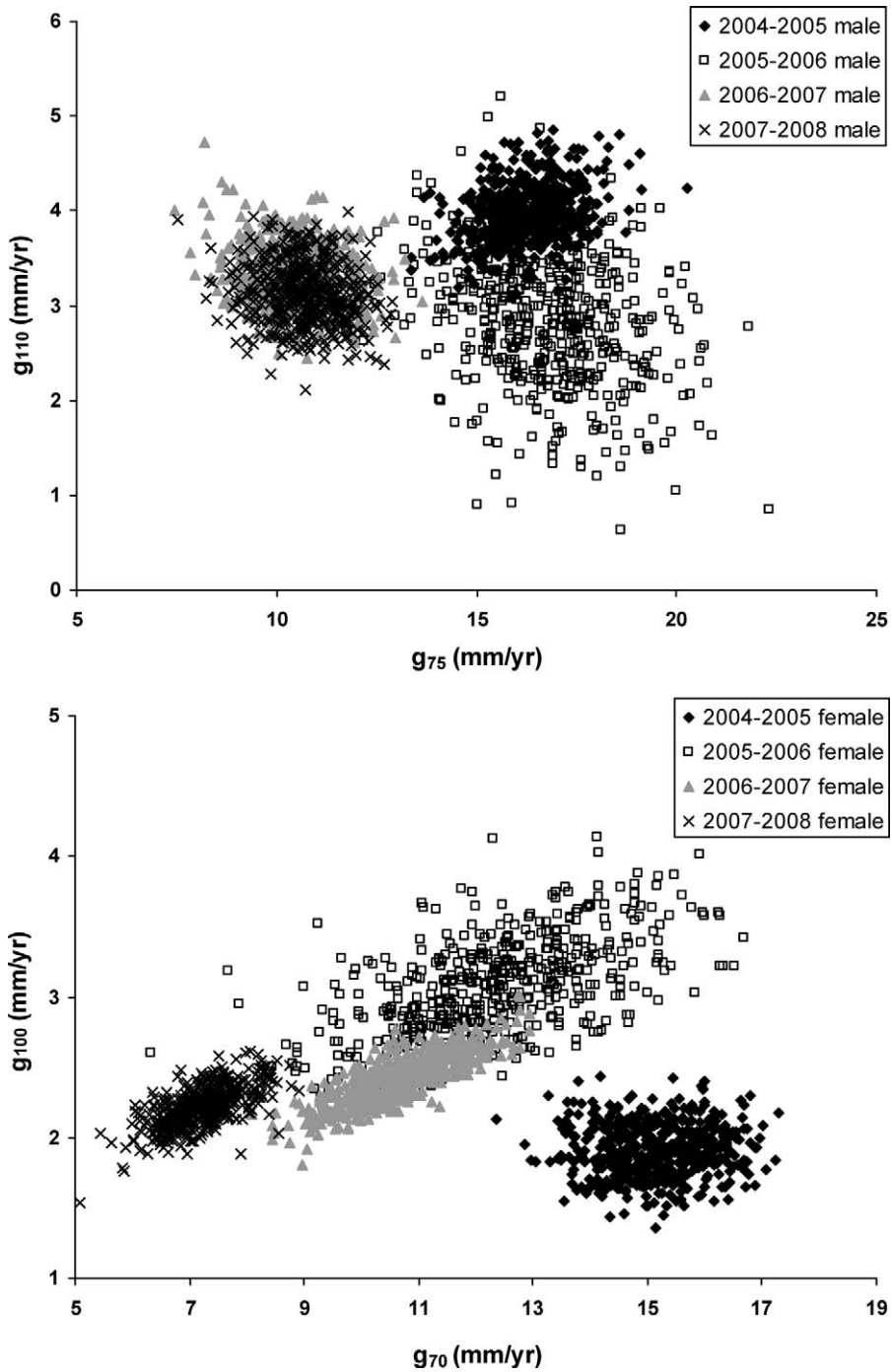


FIGURE 7.—von Bertalanffy growth model bootstrap estimates of growth ( $g$ ; mm carapace length [CL]/year) for male (top; reference lengths = 75 and 110 mm CL) and female (bottom; reference lengths = 70 and 100 mm CL) Hawaiian spiny lobsters at Maro Reef, Northwestern Hawaiian Islands, 2004–2008.

TABLE 7.—Estimated mean annual growth of 75- and 110-mm (carapace length) male and female Hawaiian spiny lobsters (NI = Necker Island; GP = Gardner Pinnacles; MR = Maro Reef;  $g_{75}$  = mean annual growth of 75-mm lobsters;  $g_{110}$  = mean annual growth of 110-mm lobsters).

Bank	Year tagged– year recaptured	$g_{75}$ , mm/year		$g_{110}$ , mm/year	
		Male	Female	Male	Female
NI	2002–2003	3.70	1.99	1.78	0.00
	2003–2004	3.71	1.78	1.93	0.07
	2004–2005	4.75	2.80	1.57	0.15
	2005–2006	5.69	2.42	2.37	0.07
	2006–2007	4.60	2.15	2.31	0.00
	2007–2008	5.35	2.34	2.07	0.00
	2002–2008	4.42	2.21	1.81	0.11
GP	2006–2007	12.96	9.28	5.66	1.03
	2007–2008	14.86	7.58	1.71	0.08
	2006–2008	12.46	7.97	5.29	0.66
MR	2004–2005	16.22	12.87	4.02	0.00
	2005–2006	16.53	10.65	2.93	0.04
	2006–2007	10.45	7.12	3.38	1.34
	2007–2008	10.65	9.17	3.14	0.74
	2004–2008	12.36	7.53	3.53	1.26

Discussion

Model and Tagging Assumptions

A reparameterization of the von Bertalanffy growth model (Francis 1988a) and a mark–recapture analog of the Schnute growth model (Francis 1995) were used to examine spatial and temporal variability in growth of NWHI Hawaiian spiny lobsters. Although both model estimates of growth were similar and residual plots indicated that both were suitable for the Hawaiian spiny lobster tag–recapture data, the von Bertalanffy model was considered the preferred model based on AIC<sub>c</sub> values. Palinurid growth has been adequately described by the von Bertalanffy model (Morgan 1980; Wahle and Fogarty 2006) and, more specifically, utilizing the GROTAG method (McGarvey et al.

1999; Montgomery et al. 2009). The Hawaiian spiny lobsters’ small growth increments relative to maximum size allow use of the von Bertalanffy model despite the assumption of continuous growth (Frisch 2007; Montgomery et al. 2009). Further, decreasing growth with increasing size and the lack of extreme outliers, particularly at small and large sizes, indicate that this growth model is suitable for the Hawaiian spiny lobster. Biases in growth rates can arise when individuals with a higher probability of zero growth (short time at liberty) are included in the analysis. All data sets only included lobsters that had been at liberty for longer time periods, hence avoiding this source of bias. Individuals recaptured after a short time at liberty usually provide information for GROTAG to estimate measurement error. Fixing the model’s measurement error parameters at values estimated independently still allowed the model to accurately estimate the growth parameters while excluding those individuals that were at liberty for shorter amounts of time.

An important assumption in tagging studies is that the tagging process does not significantly affect growth of tagged individuals relative to untagged individuals. Tag-induced effects were found in similar lobster species (Waugh 1981; Brown and Caputi 1985; Hunt and Lyons 1986). Despite taking precautions to minimize handling and exposure during the tagging process, it is impossible to disregard this assumption, although there is no evidence that the assumption was violated in this study. Even with the possibility of tag-induced effects occurring, it would not change the conclusion of growth variability because the exact tagging methodology replication spreads any bias evenly across banks and years, with the exception being Necker Island in 2002, when a different tag type was used.

TABLE 8.—Multiple regression analysis of the effects of Hawaiian spiny lobster density on individual-specific growth rates at Necker Island (NI; 2002–2008) and Maro Reef (MR; 2004–2008). Sources of variation are release size (carapace length at release, mm), catch per unit effort (CPUE; number of lobsters caught on the same string/trap; 20 traps/string) and the release size × CPUE interaction term (MS = mean square; SS = sum of squares; and df = degrees of freedom).

		Effect											
Bank	Year tagged– year recaptured	Release size			CPUE			Release size × CPUE			Error		
		MS	<i>F</i>	<i>P</i>	MS	<i>F</i>	<i>P</i>	MS	<i>F</i>	<i>P</i>	df	SS	MS
NI	2002–2003	3.02	10.39	0.001	0.78	2.67	0.10	0.68	2.33	0.13	460	133.63	0.29
	2003–2004	2.26	8.64	0.003	0.001	0.005	0.95	0.0007	0.003	0.96	896	234.45	0.26
	2004–2005	17.13	54.14	<0.001	2.74	8.67	0.003	2.31	7.30	0.007	625	197.80	0.32
	2005–2006	1.15	3.04	0.08	0.12	0.32	0.57	0.13	0.32	0.57	100	37.87	0.38
	2006–2007	0.22	0.81	0.37	<0.001	<0.001	0.991	0.0005	0.002	0.97	161	44.17	0.27
	2007–2008	2.30	7.44	0.008	1.28	4.14	0.05	1.33	4.29	0.04	80	24.71	0.31
MR	2004–2005	13.32	26.49	<0.001	0.36	0.72	0.40	0.13	0.26	0.61	76	38.21	0.50
	2005–2006	3.36	6.69	0.01	0.22	0.44	0.51	0.32	0.63	0.43	35	17.58	0.50
	2006–2007	7.09	15.24	<0.001	0.31	0.66	0.42	0.43	0.92	0.34	105	48.83	0.47
	2007–2008	2.60	6.79	0.01	1.30	3.39	0.07	1.13	2.95	0.09	119	45.52	0.39

*Hawaiian Spiny Lobster Growth Rates in the Northwestern Hawaiian Islands*

The estimated values of  $v$  indicate that the Hawaiian spiny lobster does exhibit substantial individual variability in growth, thus warranting the use of Francis' (1988a) method. Failure to explicitly account for individual variability in growth (i.e., assuming all individuals in a population grow according to fixed parameters) can result in biased mean growth estimates (Sainsbury 1980; Tyler and Rose 1994; Smith et al. 1997). The high values also indicate that such variability should be taken into account in future NWHI lobster assessments (Parma and Deriso 1990; Hampton 1991).

Significant differences in Hawaiian spiny lobster mean growth rates were found between sexes, among banks, and among years within the NWHI. Sexual dimorphism is common in palinurid growth (Aiken 1980), with males commonly growing faster (Hunt and Lyons 1986; McGarvey et al. 1999; Robertson and Butler 2003). The bank- and year-specific nature of the tag-recapture events in this study allowed for estimation of Hawaiian spiny lobster growth rates that included temporal and spatial variability. Spatial variability was particularly evident between Necker Island and the other banks. Spatial differences in growth rates have been observed in other palinurids (Newman and Pollock 1974; Joll and Phillips 1984; Pollock 1991; Skewes et al. 1997; McGarvey et al. 1999); however, at their extreme, the differences between the slowest- and fastest-growing individuals in those studies were less than the differences at the NWHI banks reported here. Temporal variability was evident to some extent within each NWHI bank, but several particularly abrupt increases and decreases in growth were found at Necker Island and Maro Reef. Temporal differences in growth similar to those observed in Hawaiian spiny lobsters were exhibited by South African Cape rock lobsters *Jasus lalandii* (Pollock et al. 1997).

The GROTAG method does allow for conversion of  $g_{\alpha}$  and  $g_{\beta}$  into  $K$  and  $L_{\infty}$  for the sake of comparison with similar species. The wide range of  $K$  estimates prevents grouping of the NWHI Hawaiian spiny lobster, as a species, with any specific group of palinurids. The Hawaiian spiny lobsters at Gardner Pinnacles and Maro Reef grew at a rate similar to that of tropical palinurids, yet Necker Island lobsters grew at a rate more typical of temperate palinurids (Wahle and Fogarty 2006; Frisch 2007).

Density-dependent growth has been exhibited in other palinurids, including the southern rock lobster *J. edwardsii* (McGarvey et al. 1999) and Tristan rock

lobster *J. tristani* (Pollock 1991). Hawaiian spiny lobster density-dependent growth was suggested by pre-exploitation surveys that documented smaller individuals but greater abundance at Necker Island relative to other banks in the NWHI (Uchida et al. 1980). Polovina (1989) concluded that density influenced NWHI Hawaiian spiny lobster population dynamics based on (1) a positive relationship between density (CPUE) and the ratio of total mortality ( $Z$ ) to growth ( $K$ ) among seven banks and (2) indications that changes in natural mortality and  $K$  occurred at Necker Island and Maro Reef after heavy exploitation. However, whether density dependence was operating on mortality, growth, or both was beyond the scope of that study. The current study did not find a significant influence of density on growth; density could not explain variability in growth at Necker Island, at Maro Reef, or between the two banks. Further, if density dependence was operating, the relatively slow growth of Hawaiian spiny lobsters at Necker Island would be associated with higher lobster densities, and a comparison of CPUEs between Necker Island and Maro Reef does not support this hypothesis. This theory also applies to density dependence within Necker Island. If density dependence was operating, fast growth would be expected in 2002–2008 because CPUE was lower relative to CPUE during the advent of the fishery (Figure 2). It now seems plausible that the smaller Hawaiian spiny lobsters observed at Necker Island in the pre-exploitation surveys (Uchida et al. 1980) were a result of the relatively slow growth rather than density dependence.

Direct temperature effects have been linked to growth variability in other palinurid species (Wahle and Fogarty 2006); however, growth variability of NWHI Hawaiian spiny lobsters was not associated with differences in temperature. Temperatures were very similar between Necker Island and Maro Reef during the span of this study, yet Hawaiian spiny lobster growth was not, further discounting temperature as the cause of growth differences between banks.

It also appears likely that the differences in Hawaiian spiny lobster growth between Necker Island and the other banks cannot be attributed to genetic differences. An analysis of allozyme frequencies prior to exploitation found a single panmictic stock throughout the Hawaiian Archipelago (Shaklee and Samollow 1984), and postexploitation analysis of allozyme frequencies in Necker Island and Maro Reef Hawaiian spiny lobsters found no differences in six out of seven loci (Seeb et al. 1990). It is also unlikely that the differences in growth among banks are a result of genetic selection for slower-growing individuals by the commercial fishery (Conover and Munch 2002) based on the



historically smaller Hawaiian spiny lobsters at Necker Island; the similar levels of exploitation at Necker Island, Gardner Pinnacle, and Maro Reef; and the lack of genetic structure in this species across the archipelago.

Because the slower growth of Hawaiian spiny lobsters at Necker Island cannot be explained by common drivers of variability (e.g., density, temperature, or genetics), the reason for the slower growth may be that a crucial dietary component of "normal" growth is missing at Necker Island. Temporal differences in Cape rock lobster growth were attributed to the collapse of their primary food source during the anomalous El Niño years of 1990–1993 (Pollock et al. 1997). Spatial variability in growth rates of juvenile western rock lobsters *Panulirus cygnus* was associated with local abundances of the top shell *Cantharidus lepidus*; western rock lobsters in the fast-growth areas ingested more top shells and less of the relatively nutritionally poor plant material than did lobsters in the slow-growth area (Edgar 1990).

The NWHI is a dynamic ecosystem that is poorly understood, and the factors that caused the growth differences between Necker Island and the other banks in the NWHI are, at this point, not readily identifiable. Future research should focus on aspects of the Hawaiian spiny lobster's diet, such as stable isotope analysis and nutritional indices (Parrish and Martinelli-Liedtke 1999), to help pinpoint the drivers of the growth variability. The variability in growth rates also suggests variability in other Hawaiian spiny lobster life history traits (e.g., mortality) as well as in life histories of other lobster species. The assumption that the scaly slipper lobster *Scyllarides squammosus*, an important component of the NWHI lobster fishery, also displays synchronous life history dynamics should now be examined. The results from such studies would not only provide species-specific information but would also provide insights into the dynamics of the NWHI ecosystem.

The results of this tagging study indicate that the assumption of homogenous Hawaiian spiny lobster population dynamics, a foundation of previous population assessments, is invalid. The application of growth rates estimated at one bank to other banks is likely inappropriate, as is the assumption of uniform annual growth within a given bank. A re-examination of the 1979–1983 Kure Atoll and French Frigate Shoals tagging data using the von Bertalanffy model fitted with the same maximum likelihood method employed here documented growth rates (Kure Atoll males:  $g_{75} = 15.85$  mm/year,  $g_{145} = 1.06$  mm/year; Kure Atoll females  $g_{70} = 15.81$  mm/year,  $g_{135} = 0.41$  mm/year; French Frigate Shoals males:  $g_{75} = 11.84$

mm/year,  $g_{110} = 5.01$  mm/year; French Frigate Shoals females:  $g_{70} = 10.99$  mm/year,  $g_{110} = 7.04$  mm/year; O'Malley and MacDonald 2009) that were similar to the contemporary Maro Reef and Gardner Pinnacles growth rates. The documented smallest average size of Hawaiian spiny lobsters at Necker Island from the pre-exploitation surveys (Uchida et al. 1980) also suggests that the slower growth rates exhibited by Necker Island lobsters in 2002–2008 are not a recent phenomenon. If, in the past, Hawaiian spiny lobsters at these banks grew at rates similar to those of contemporary populations, then the application of Necker Island growth rates to other banks was erroneous, as was applying Kure Atoll and French Frigate Shoals growth rates to Necker Island lobsters. Both were damaging in that (1) the Necker Island growth rates were significantly different from those at other banks and (2) because Necker Island was a focal point of the commercial fishery, the catch information contributed significantly to archipelago-wide assessments (Botsford et al. 2002).

Small differences in growth rates significantly affect outputs from population models (Jones 1979; Majowski et al. 1987; Lai and Gallucci 1988; Helser and Brodziak 1998). The identified spatial, temporal, and individual variability in growth of the Hawaiian spiny lobster lends credence to the possibility that ignoring variability in life history parameters introduced significant biases to previous NWHI lobster assessments. Recognition of and accounting for the mean and individual variability in a life history trait of this important endemic species will result in more accurate stock assessments and, ultimately, better understanding of lobster dynamics and the NWHI coral reef ecosystem.

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