























LETTER

Collaborative fisheries research reveals reserve size and age determine efficacy across a network of marine protected areas

Shelby L. Ziegler^{1,2}  | Rachel O. Brooks¹  | Lyall F. Bellquist^{3,4}  |
 Jennifer E. Caselle⁵  | Steven G. Morgan^{6,7}  | Timothy J. Mulligan⁸ |
 Benjamin I. Ruttenberg⁹  | Brice X. Semmens⁴  | Richard M. Starr¹  |
 Joe Tyburczy^{8,10}  | Dean E. Wendt⁹  | Andre Buchheister⁸ |
 Jose R. Marin Jarrin^{8,11}  | Christina Pasparakis⁶  | Salvador J. Jorgensen¹²  |
 Jennifer A. Chiu¹  | Jordan Colby⁶ | Connor L. Coscino⁴ | Leon Davis⁸ |
 Francine de Castro⁶ | Jack T. Elstner⁴  | Christopher Honeyman⁵  |
 Erica T. Jarvis Mason⁴  | Erin M. Johnston⁹  | Sadie L. Small⁶  | Jay Staton⁸ |
 Grant T. Waltz⁹  | Bonnie Basnett¹  | Erin V. Satterthwaite^{6,10}  |
 Helen Killeen⁶  | Connor D. Dibble⁶ | Scott L. Hamilton¹ 

¹Moss Landing Marine Laboratories, San Jose State University, Moss Landing, California, USA

²Odum School of Ecology, University of Georgia, Athens, Georgia, USA

³California Oceans Program, The Nature Conservancy, San Diego, California, USA

⁴Scripps Institution of Oceanography, University of California, San Diego, La Jolla, California, USA

⁵Marine Science Institute, University of California Santa Barbara, Santa Barbara, California, USA

⁶Bodega Marine Laboratory, University of California, Davis, Bodega Bay, California, USA

⁷Coastal Marine Sciences Institute, Department of Environmental Science and Policy, University of California Davis, Davis, California, USA

⁸Department of Fisheries Biology, Telonicher Marine Lab, California Polytechnic State University, Arcata, California, USA

⁹Bailey College of Science and Mathematics, California Polytechnic State University, San Luis Obispo, San Luis Obispo, California, USA

¹⁰California Sea Grant, Scripps Institution of Oceanography, University of California, San Diego, La Jolla, California, USA

¹¹Charles Darwin Research Station, Charles Darwin Foundation, Puerto Ayora, Galápagos, Ecuador

¹²Department of Marine Science, California State University, Monterey Bay, Seaside, California, USA

Correspondence

Shelby L. Ziegler, Odum School of Ecology, University of Georgia, 140 E. Green St, Athens, GA 30602 USA. Email: shelbyziegler@gmail.com

Funding information

California Ocean Protection Council, Grant/Award Number: C0752003;

Abstract

A variety of criteria may influence the efficacy of networks of marine protected areas (MPA) designed to enhance biodiversity conservation and provide fisheries benefits. Meta-analyses have evaluated the influence of MPA attributes on abundance, biomass, and size structure of harvested species, reporting that MPA size, age, depth, and connectivity influence the strength of MPA responses.

This is an open access article under the terms of the [Creative Commons Attribution](https://creativecommons.org/licenses/by/4.0/) License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

© 2023 The Authors. *Conservation Letters* published by Wiley Periodicals LLC.

California Department of Fish and Wildlife, Grant/Award Number: P1970018; California Sea Grant, University of California, San Diego, Grant/Award Number: R/MPA-45

However, few empirical MPA evaluation studies have used consistent sampling methodology across multiple MPAs and years. Our collaborative fisheries research program systematically sampled 12 no-take or highly protective limited-take MPAs and paired fished reference areas across a network spanning 1100 km of coastline to evaluate the factors driving MPA efficacy across a large geographic region. We found that increased size and age consistently contributed to increased fish catch, biomass, and positive species responses inside MPAs, while accounting for factors such as latitude, primary productivity, and distance to the nearest MPA. Our study provides a model framework to collaboratively engage diverse stakeholders in fisheries research and provide high-quality data to assess the success of conservation strategies.

KEYWORDS

biomass, California, community-based science, empirical data, fish communities, marine protected areas, productivity, reserve network

1 | INTRODUCTION

Conservation strategies such as marine protected areas (MPAs), including no-take marine reserves, are popular spatial management tools used to protect marine biodiversity, promote resilient ecosystems, and deliver benefits to society (Mouillot et al., 2020). Due to the success of many MPAs that limit fishing activities in supporting species abundances (Cinner et al., 2018) and human well-being (Ban et al., 2019), policymakers have also leaned on these strategies to help mitigate the effects of global climate change (Naeem et al., 2022). However, the measured outcomes of MPAs can vary greatly across space and time based on protection level, enforcement, MPA age, and various ecological conditions (Gill et al., 2017; Grorud-Colvert et al., 2021; Meehan et al., 2020).

Global meta-analyses suggest that the cessation of fishing in MPAs generally increases the abundance of targeted fish species (Edgar et al., 2014; Giakoumi et al., 2017), with more delayed indirect effects on nontargeted species (Babcock et al., 2010). However, for any single MPA, the abundance of fished species may remain constant or even decrease through time, possibly in response to external factors, such as oceanographic conditions, recruitment variability, changes in food webs or habitats (Edgar & Barrett, 2012) and gravity of human impacts (Cinner et al., 2018). Such variable results have led researchers to investigate the specific factors that drive organismal responses in MPAs (Nickols et al., 2019) along with increased development of MPA networks, which are designed to buffer populations from those external drivers, rather than stand-alone MPAs (Grorud-Colvert et al., 2014).

Theoretical benefits of MPA networks include protecting diversity of habitats and species, reducing localized

impacts of climate change, maintaining genetic diversity and population persistence (White et al., 2011), and creating a more equitable distribution of fisheries costs and benefits (Leenhardt et al., 2015). Understanding how species respond to MPA protection is essential to set appropriate expectations for stakeholders, as well as to adaptively manage the network into the future. Theoretical studies and meta-analyses suggest that reserve characteristics such as size, age, connectivity, location, and adjacent fishing effort may influence the magnitude of positive effects from MPA networks (Claudet et al., 2008; Edgar et al., 2014; Lester & Halpern, 2008). While meta-analyses can provide a wealth of information on diverse ecosystems and species across space and time, the data quality can vary greatly (Halpern & Warner, 2003) and empirical data are often collected in using different methods (e.g., visual census, fishing, underwater video; Claudet et al., 2010), influencing the conclusions researchers draw.

From 2003 to 2012, the government of California, a state located on the Pacific coast of the USA, implemented a statewide network of MPAs to restrict harvest of marine species and to provide conservation and fisheries benefits using the best available science on habitat, species diversity, dispersal capability, and distribution of human uses. This process resulted in a scientifically designed MPA network that includes 124 MPAs, spanning 1100 km of coastline and 2200 km² (16%) of state waters (Botsford et al., 2014). Baseline monitoring of the newly established network was conducted across diverse habitats, including kelp forests and rocky reefs. Data from these monitoring programs indicated that both MPAs and areas open to fishing within the same geographic area had similar abundance, biomass, and species composition when MPAs were first enacted (e.g., North coast: Mulligan et al., 2017;

Central coast: Starr et al., 2015, South coast: Caselle et al., 2015).

We evaluated the effects of no-take or limited take reserves at 12 MPAs within the California MPA network using 6 years (2017–2022) of standardized hook and line surveys conducted by the California Collaborative Fisheries Research Program (CCFRP). CCFRP is a unique partnership among scientists, managers, and anglers that combines the expertise and ideas of diverse stakeholders to gather information on fish stocks for management and conservation (Wendt & Starr, 2009) while improving buy-in for fisheries management (Mason et al., 2020). By using catch data collected by CCFRP inside and outside MPAs, we are able to assess how a variety of metrics including MPA age, size, adjacent fishing pressure, geographic location, network connectivity, and environmental factors affect total catch rates, catch biomass, and the number of species exhibiting positive responses to MPA protection (i.e., higher biomass inside MPAs compared with reference sites). We expected catch rates and catch biomass to be higher in larger and older MPAs (McClanahan & Graham, 2015). Further, we expected more rapid population responses and stronger MPA effects at lower latitudes in southern California where human population density (Free et al., 2023) and fishing pressure are greater and warmer water temperatures are associated with faster growth rates and more consistent recruitment of the dominant taxa (Caselle et al., 2023).

2 | METHODS

2.1 | CCFRP surveys

CCFRP collaborates with volunteer anglers and the fishing industry to conduct surveys of rocky reef-associated fish communities inside and outside of MPAs using standardized, repeatable hook-and-line fishing protocols (Wendt & Starr, 2009). Annually, between 2017 and 2022, CCFRP sampled 12 MPAs (10 no-take State Marine Reserves [SMRs], one highly protected limited-take State Marine Conservation Areas [SMCAs], and one paired SMR/SMCA) and 12 paired reference sites open to fishing (REFs) from Humboldt County in Northern California to San Diego County in Southern California (Figures S1–S3). Each of the 24 total sites was sampled with standardized hook-and-line protocols for 3 days (i.e., visits) per year, with four randomly selected grid cells (each 500 m × 500 m in area) sampled per day (Starr et al., 2015). Within each grid cell, we completed three 15-min fishing periods, where volunteer anglers actively fished over rocky reef habitat. All fish captured were identified to the lowest taxonomic level, measured to total length

(cm), and released. For additional details on CCFRP, reference site selection, and fishing protocols please refer to the Supporting Information Text 1.

2.2 | MPA characteristics

We calculated the age of each MPA by subtracting the year of implementation from the sampling year. MPAs were implemented in various years: Channel Islands in 2003, central coast in 2007 and northern and southern (non-island) coasts from 2010 to 2012. Total protected area (km²) was defined as the MPA area for solitary MPAs and the sum of protected areas for paired MPAs (locations where no-take SMRs and highly protective limited-take SMCAs were adjacent to each other). We extracted the latitude for the centroid of each MPA for spatial analyses. We calculated the distance from the nearest MPA (excluding SMR/SCMA pairs) within the network to assess the influence of connectivity and nearest port as a proxy for fishing pressure (Harborne et al., 2018). These distances were measured as the shortest linear distance over water from the boundary of the MPA to the nearest MPA or port using ArcGIS PRO (version 3.0.3). Across our study system, latitude and human population density are highly correlated and previous work along this coast suggests that human population density may not be the best indicator of human engagement with MPAs (Free et al., 2023), so we did not consider that metric in our MPA characteristics. Fishing pressure directly adjacent to an MPA has been shown to modulate the magnitude of positive effects on fish populations (biomass and catch) across a portion of this network (Ziegler et al., 2022), but recreational fishing data at the statewide-scale are limited and less reliable in some regions. Therefore, we calculated an index of recreational fishing pressure outside each MPA as the total number of groundfish landed between 2017 and 2019 for the nearest port complex (from the CalFISH database; Free et al., 2022) divided by the distance from the nearest port (under the assumption that realized fishing pressure declines with distance from port). Due to lack of high-quality habitat mapping across all of our sampling sites, we used the habitat rugosity metric from each CCFRP fishing drift (Supporting Information Text 1) to calculate the mean difference in vertical relief among the MPAs and REFs. The difference in relief was the average of all rugosity scores from drifts inside each MPA within a year divided by the average rugosity scores for drifts in the associated REF. Scores of 1 indicate the relief in the sampled areas of the MPAs and REFs were equal; while scores greater than 1 indicate higher relief in the sampled areas of the MPA compared with the REF. We also extracted environmental variables of sea surface temperature (°C; SST) and net pri-

mary production ($\text{mg C m}^{-2} \text{ day}^{-1}$; NPP) from the Central and Northern California Ocean Observing System Repository. SST was collected by Advanced Very High-Resolution Radiometer instrument aboard NOAA's Polar Operational Environmental Satellites from 2004 to 2022 at a 1.47 km resolution and NPP was collected by the California Current Merged Satellite daily from 1996 to 2022 at a 4-km spatial resolution. The spatial average of both SST and NPP within the boundaries of each MPA were extracted annually for each year of CCFRP sampling. See Table 1 for a summary of MPA characteristics for all sampling sites.

2.3 | Analysis

We calculated both catch per unit effort (CPUE) and biomass per unit effort (BPUE) as relative metrics of abundance and biomass in each MPA and REF site. CPUE was calculated by dividing the total number of fishes caught by total angler hours of fishing in all grid cells in a day ($\text{catch angler h}^{-1}$). For each individual fish, we converted length to biomass using published length-weight relationships (Love et al., 1990; Froese & Pauly, 2023). We then calculated BPUE as the total weight of fish caught in kilograms divided by the number of angler hours fished (kg angler h^{-1}). CPUE and BPUE for the grid cells sampled on a given day was then averaged to estimate CPUE or BPUE for the total fish community inside and outside each MPA for a given sampling year. Data were tested for normality with a Shapiro–Wilk test ($W = 0.94$, $p < 0.001$).

To assess the drivers of spatial variation in the strength of MPA responses across the network, we calculated a yearly response ratio for each MPA using both CPUE and BPUE values (Stewart, 2010). Response ratios were calculated as the log of the value (X) inside the MPA divided by the value (X) at the REF for year i :

$$\text{Log}_{10} \left(\frac{X_{\text{MPA}, i}}{X_{\text{REF}, i}} \right)$$

A ratio above 0 indicates higher CPUE or BPUE inside the MPA relative to REF sites, while the magnitude of the ratio reflects the relative difference in CPUE or BPUE between each MPA and its associated REF (i.e., the strength of MPA response) on the log_{10} scale. Using a BPUE response ratio calculated independently for each species, we determined the number of positive species responses for each MPA and year.

To determine the relative influence of MPA characteristics on the efficacy of MPAs to enhance fish catches (i.e., response ratios), we fit generalized linear mixed effects models with normally distributed errors using the lme4 package in R (Bates et al., 2015; Table 1). Prior to anal-

yses, we tested for collinearity among fixed effects using a Pearson correlation matrix (Figure S4). Both SST and the difference in relief between MPAs and REFs were highly correlated with latitude ($r > 0.7$) and therefore were removed from subsequent models. Our final models included MPA age, latitude, total protected area (km^2), relative fishing pressure (landings [n] per distance from nearest port [km]), mean NPP ($\text{mg C m}^{-2} \text{ day}^{-1}$), and distance to the nearest MPA (km) within the MPA network as fixed effects with a random effect of sampling site. We also considered interactions that were deemed the most ecologically relevant or plausible: MPA age and latitude, fishing pressure and latitude, and total protected area and fishing pressure. For the metric of the number of positive species responses, including the random effect overfit the models and therefore we used negative binomial models with no random effects. Akaike information criterion corrected for small sample sizes (AICc) was used to compare models with all possible variable combinations. Models were ranked using their AICc difference from the model with the minimum AICc (ΔAICc), and we considered all models with $\Delta\text{AICc} < 2$ for interpretation. Beta coefficients (β) were standardized by standard deviation for direct comparisons across models. We then conducted model averaging in the MuMIn package (Bartoń, 2023) to assess which variables best predicted the response variables across the top models and assessed the partial effects of each factor on MPA responses (Burnham & Anderson, 2002). All analyses were conducted in R statistical software version 4.3.0 (R Development Core Team 2023).

3 | RESULTS

We observed greater total catch rates (CPUE) and catch biomass (BPUE) inside MPAs compared with REFs at all locations surveyed across the network of MPAs from Northern to Southern California (Figures 1 and S5, S6).

Both CPUE and BPUE response ratios varied across the MPA network through time (Figure 2a). Of the 12 MPAs sampled, 7 had positive CPUE response ratios and 10 had positive BPUE response ratios in all sampling years (Figure 2b), indicating consistently higher total catch rates and greater biomass inside MPAs. The largest effect sizes for both CPUE and BPUE (100-fold higher BPUE in MPA relative to the REF) occurred at Swami's SMCA (where absolute values were low in both MPA and REF) and at Anacapa Island SMR/SMCA (where values for both CPUE and BPUE in REF were extremely low) (Figure 2). Overall, 19 of 72 possible year-site combinations for CPUE and 29 of the combinations for BPUE showed response ratios of $>=1$, respectively (10-fold higher values of catch and biomass inside MPAs relative to REFs).

TABLE 1 MPA characteristics for each of the 12 MPAs sampled by CCFRP. MPA characteristics include year of MPA establishment, MPA name, whether the MPA is solitary (no) or adjacent to another marine managed area with less restrictive no-take regulations (yes), the amount of total protected area (km²), the latitude of the centroid for the MPA, the nearest port name, distance to the nearest port in kilometers, groundfish landings (no. groundfish y⁻¹) extracted from the CalFish database (Free et al., 2022), fishing pressure (landings/distance to nearest port), distance to the nearest MPA in kilometers and the difference in relief calculated by dividing the mean rugosity score (1–3) from each fishing drift in a MPA by the rugosity score (1–3) from each drift in the REF. Mean net primary production (mg C m⁻² day⁻¹) and sea surface temperature (°C) are not shown as they were examined on an annual basis.

Region	Year established	MPA name	Paired MPA	Total protected area (km ²)	Latitude (°N)	Nearest port	Landings			Distance to nearest MPA (km)	Difference in relief
							Distance to nearest port (km)	no. groundfish y ⁻¹	Fishing pressure		
North	2012	Cape Mendocino SMR	No	23.52	40.425	Eureka	54.49	61,597	1130	9.11	0.82
	2012	Ten Mile SMR	Yes	40.12	39.585	Fort Bragg (Noyo River)	20.12	61,597	3062	5.94	1.00
Central	2010	Stewarts Point SMR	No	62.24	38.626	Bodega Bay	59.61	136,415	2289	7.26	1.02
	2010	Bodega Head SMR	Yes	56.07	38.279	Bodega Bay	10.5	136,415	12,992	11.48	0.97
South	2007	Año Nuevo SMR	Yes	28.88	37.078	Santa Cruz	39.86	151,247	3795	31.85	0.92
	2007	Point Lobos SMR	Yes	36.18	36.505	Monterey	26.03	151,247	5811	2.58	0.99
	2007	Piedras Blancas SMR	Yes	49.94	35.683	Morro Bay	60.71	254,141	4186	6.98	1.01
	2007	Point Buchon SMR	Yes	48.87	35.219	Morro Bay	20.13	254,141	12,625	12.22	1.11
	2003	Anacapa Island SMR/SMCA	Yes	48.82	34.045	Ventura	27.94	562,145	20,120	7.63	1.11
	2003	Carrington Point SMR	No	33.10	34.038	Santa Barbara	52.48	562,145	10,712	4.27	0.98
	2012	Swamis SMCA	No	58.33	33.025	Oceanside	22.33	94,255	4221	12.96	1.18
	2012	South La Jolla SMR	Yes	39.47	32.813	Quivira Basin (San Diego)	10.27	117,194	11,411	3.55	1.15

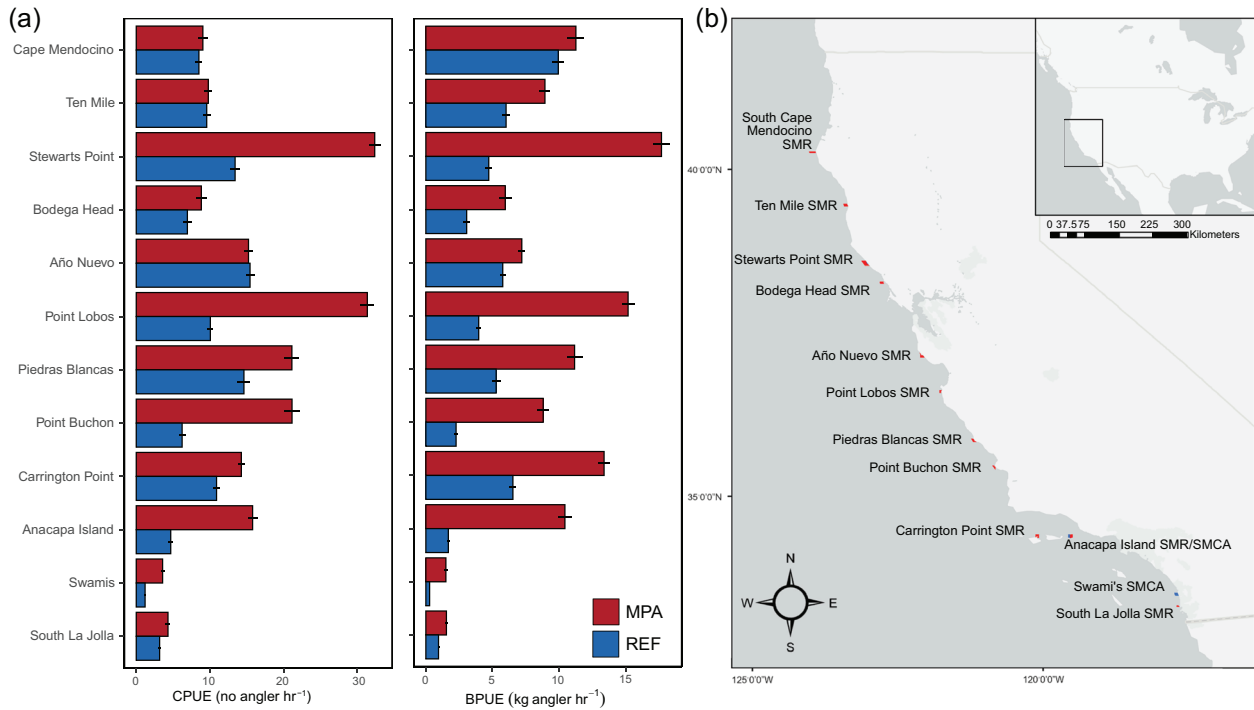


FIGURE 1 (a) The average catch per unit effort (CPUE; no angler h⁻¹) and biomass per unit effort (BPUE; kg angler h⁻¹) of fishes at 12 marine protected areas (MPAs, red bars) and reference sites (REFs, blue bars) along the coast of California across all 6 years sampled statewide (2017–2022). Values are mean \pm 95% CI. (b) Map of 12 MPAs (red polygons) sampled along the California coast.

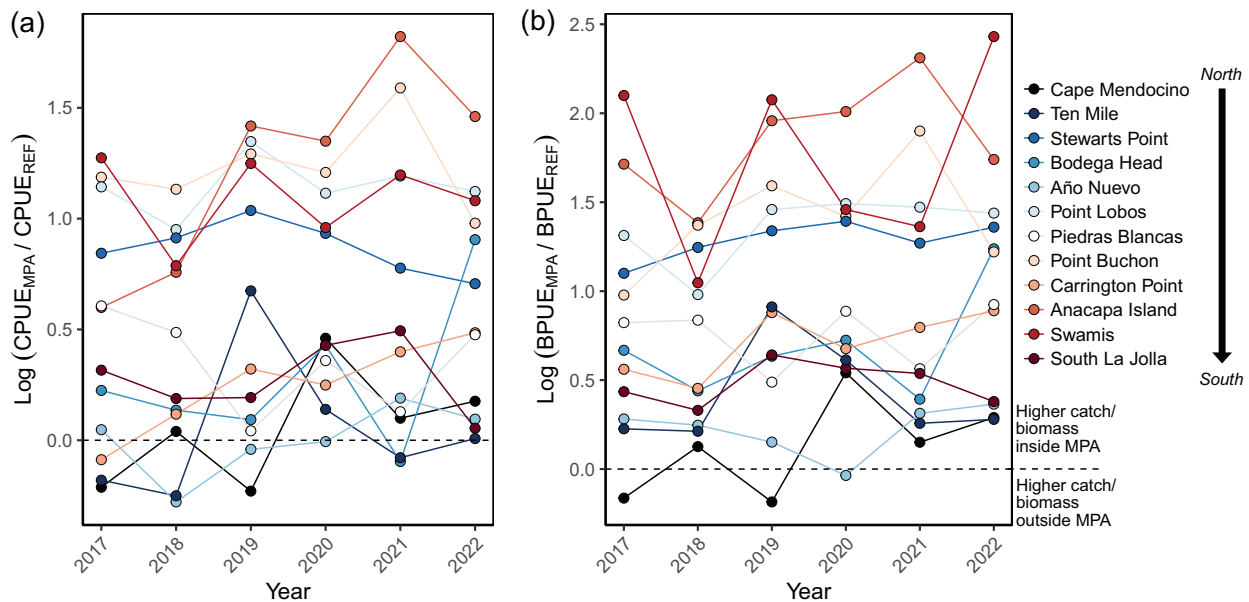


FIGURE 2 Catch per unit effort (CPUE; no angler h⁻¹) and Biomass per unit effort (BPUE; kg angler h⁻¹) response ratios for the 12 marine protected areas (MPAs) spanning 10 degrees of latitude along the coast California from 2017 to 2022. Paired MPA/REF sites in the legend are ordered and colored by latitude and relative differences in sea surface temperature (blue = northern and cool, red = southern and warm). Response ratio values (y-axis) greater than zero indicate higher total fish biomass inside the MPAs relative to the reference sites. Dashed line indicates no difference in CPUE or BPUE between MPA and REF sites.

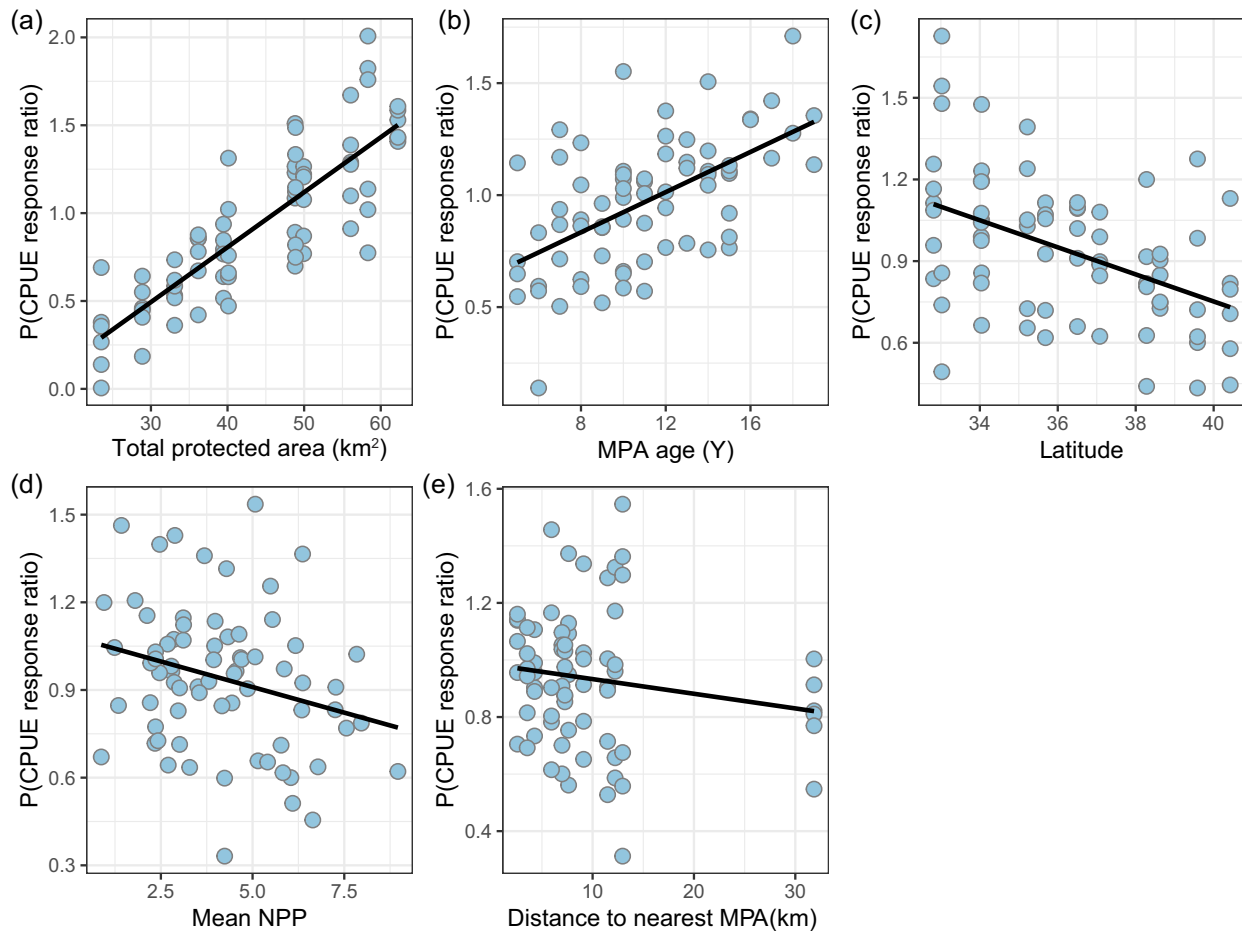


FIGURE 3 Partial effects plot of the full model for CPUE response ratios with all variables included in model averaging. Partial residuals of response ratios as a function of (a) total amount of protected area (km^2), (b) MPA age, (c) latitude, (d) mean NPP, and (e) distance to nearest MPA while other fixed effects were held constant at median values. Trend lines indicate conditional fit of the models without the random effect of site.

Four competing models best fit the CPUE data, including variables of MPA age, area, latitude, mean NPP, and distance to the nearest MPA. After model averaging the predictor variables that best explained CPUE response ratios were MPA age (with coefficient, $\beta = 0.29$, $p = 0.006$) and total protected area ($\beta = 0.50$, $p = 0.01$; Figure 3). For BPUE response ratios, six competing models best fit the data, including variables of MPA age, area, latitude, mean NPP, distance to the nearest MPA, and fishing pressure. However, similar to CPUE, MPA age ($\beta = 0.29$, $p = 0.002$) and total protected area ($\beta = 0.29$, $p = 0.002$) were the top predictors of BPUE response ratios across all MPAs. MPA age and total protected area were positively associated with CPUE and BPUE responses ratios observed (Table 2), such that MPAs that were both larger and older tended to have higher total catch rates and fish biomass inside MPAs relative to the reference sites (Figure 4).

The number of positive species responses varied across MPAs and as a function of sampling year, with the percentage of positive species responses ranging from 40 to 100%.

Of the 72 possible year-site combinations, 65 had more positive species responses than negative species responses (>50%; Figure 5). Three models best explained, the total number of positive species responses within MPAs over time. After model averaging, the top predictor variables were total protected area ($\beta = 0.12$, $p < 0.001$), fishing pressure ($\beta = 0.45$, $p = 0.03$), as well as interactive effects of area and fishing pressure ($\beta = -0.50$, $p < 0.001$; Table 2). The number of species exhibiting positive responses to MPAs was higher in larger MPAs and in locations where fishing pressure was higher outside the MPA; however, the fishing pressure effect was dampened in locations with large MPAs.

4 | DISCUSSION

Size and age of no-take or highly protective MPAs positively affected fish catch rates and catch biomass. For many of the MPAs sampled, response ratios were greater in 2022

TABLE 2 Comparison of top eight models predicting the influence of MPA characteristics (MPA age, total protected area, relative fishing pressure, latitude, mean NPP and distance to the nearest MPA) on CPUE and BPUE response ratios, and number of positive species responses across 12 MPAs in a temperate marine MPA network. For each model, marginal R^2 (R^2_m), conditional R^2 (R^2_c), degrees of freedom (df), log likelihood (LL), AICc, and Δ AICc, are shown. Shaded models had a Δ AICc < 2.

Response variable	Model rank	MPA age	Total protected area	Latitude	Fishing pressure	NPP	nearest MPA	Age × latitude	Area × fishing pressure	Fishing pressure × latitude	R^2_m	R^2_c	df	LL	AICc	Δ AICc
CPUE response ratios	1	X	X								0.33	0.79	5	-15.51	41.9	0
	2	X	X	X							0.39	0.80	6	-15.03	43.3	1.42
	3	X	X		X						0.35	0.78	6	-15.23	43.8	1.83
	4	X	X			X					0.35	0.79	6	-15.28	43.9	1.93
	5	X	X	X	X						0.36	0.80	6	-15.37	44	2.11
	6	X	X								0.08	0.79	4	-18.11	44.8	2.9
	7	X	X	X	X		X				0.40	0.80	7	-14.84	45.4	3.51
	8	X	X	X	X	X					0.40	0.79	7	-14.85	45.4	3.51
BPUE response ratios	1	X	X								0.47	0.82	5	-19.10	49.1	0
	2	X	X	X							0.55	0.83	6	-18.13	49.6	0.44
	3	X	X			X					0.52	0.82	6	-18.19	49.7	0.55
	4	X	X	X	X	X					0.57	0.83	7	-17.46	50.7	1.55
	5	X	X	X			X				0.49	0.82	6	-18.86	51	1.89
	6	X	X			X					0.50	0.83	6	-18.89	51.1	1.95
	7	X	X	X	X		X				0.56	0.83	7	-17.93	51.6	2.5
	8	X	X	X	X	X					0.53	0.82	7	-18.04	51.8	2.72
Positive species responses	1		X	X	X				X		0.19	-	6	-162.6	338.4	0
	2	X	X	X	X		X		X		0.23	-	8	-160.5	339.2	0.79
	3		X	X	X	X			X	X	0.20	-	7	-162.3	340.3	1.86
	4		X	X	X	X			X		0.19	-	7	-162.5	340.7	2.24
	5	X	X	X	X	X			X		0.19	-	7	-162.5	340.8	2.31
	6	X	X	X	X	X	X		X		0.19	-	7	-162.5	340.8	2.32
	7	X	X	X	X	X		X	X	X	0.23	-	9	-160.4	341.8	3.36
	8	X	X	X	X	X	X	X	X	X	0.23	-	9	-160.5	341.8	3.37

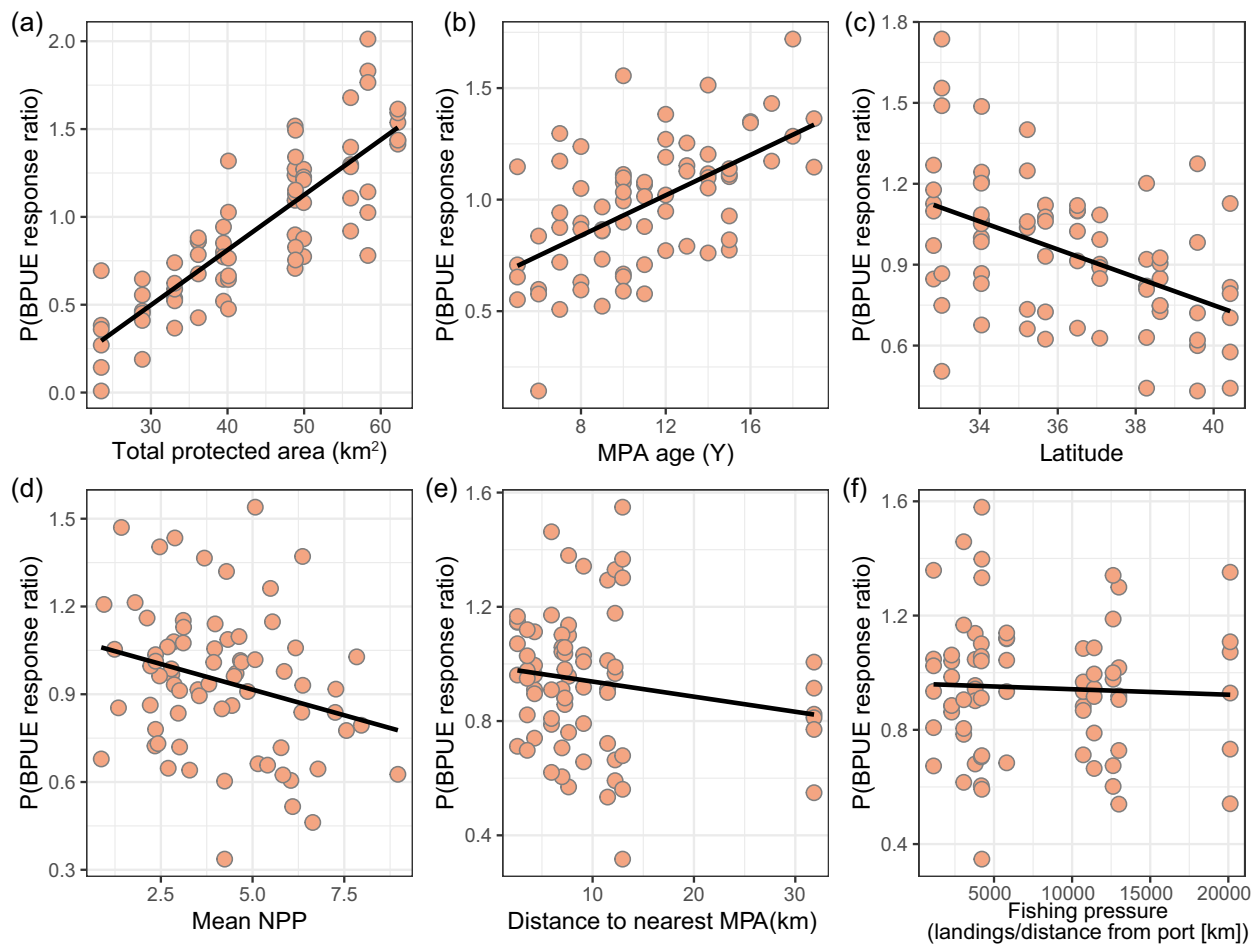


FIGURE 4 Partial effects plot of the full model for BPUE response ratios with all variables included in model averaging. Partial residuals of response ratios as a function of (a) total amount of protected area (km^2), (b) MPA age, (c) latitude, (d) mean NPP, (e) distance to nearest MPA, and (f) fishing pressure, while other fixed effects were held constant at median values. Trend lines indicate conditional fit of the model without the random effect of site.

compared with 2017 when CCFRP expanded sampling across the state (the program started in central California in 2007; Starr et al., 2015), even though the MPAs were implemented at different times in different regions between 2003 and 2012. Extensive efforts (such as consulting multibeam habitat maps and members of the charter fishing community) were made at the beginning of the CCFRP program to identify reference areas that were equivalent in habitat to the MPAs sampled (Yochum et al., 2011). Similarly, baseline sampling and times series analysis in select regions indicated that fish density and biomass values were similar among MPAs and REFs at the time of MPA implementation across the California coast (Caselle et al., 2015; Hamilton et al., 2010; Starr et al., 2015; Mulligan et al., 2017; Ziegler et al., 2022). As a result, we interpret that the differences we observed in catch rates and biomass are due to the removal of fishing pressure inside the MPAs and not pre-existing differences.

In general, we observed stronger responses to MPA protection in older MPAs (Anacapa, Carrington Pt., Pt. Lobos) compared with younger MPAs implemented in 2010–2012, closer to the expansion of CCFRP to a statewide MPA monitoring program. Synthetic analyses of European MPAs similarly show that fish density increases with the age of the MPA (Claudet et al., 2008) while global analyses report that older MPAs (>10 years since implementation) host increased fish biomass (Edgar et al., 2014). However, we did observe comparably high responses in some newer MPAs in each region (e.g., Swami's, Stewarts Pt.), although these MPAs tended to have higher total amounts of protected area and Swami's had much lower BPUE overall. Many of the response ratios are relatively flat over the time series, possibly indicating that the effects of the MPAs have begun to saturate (McClanahan et al., 2007; Russ & Alcala, 2010), that extensive spillover may be occurring (Russ & Alcala, 2011) or that recovery is occurring very

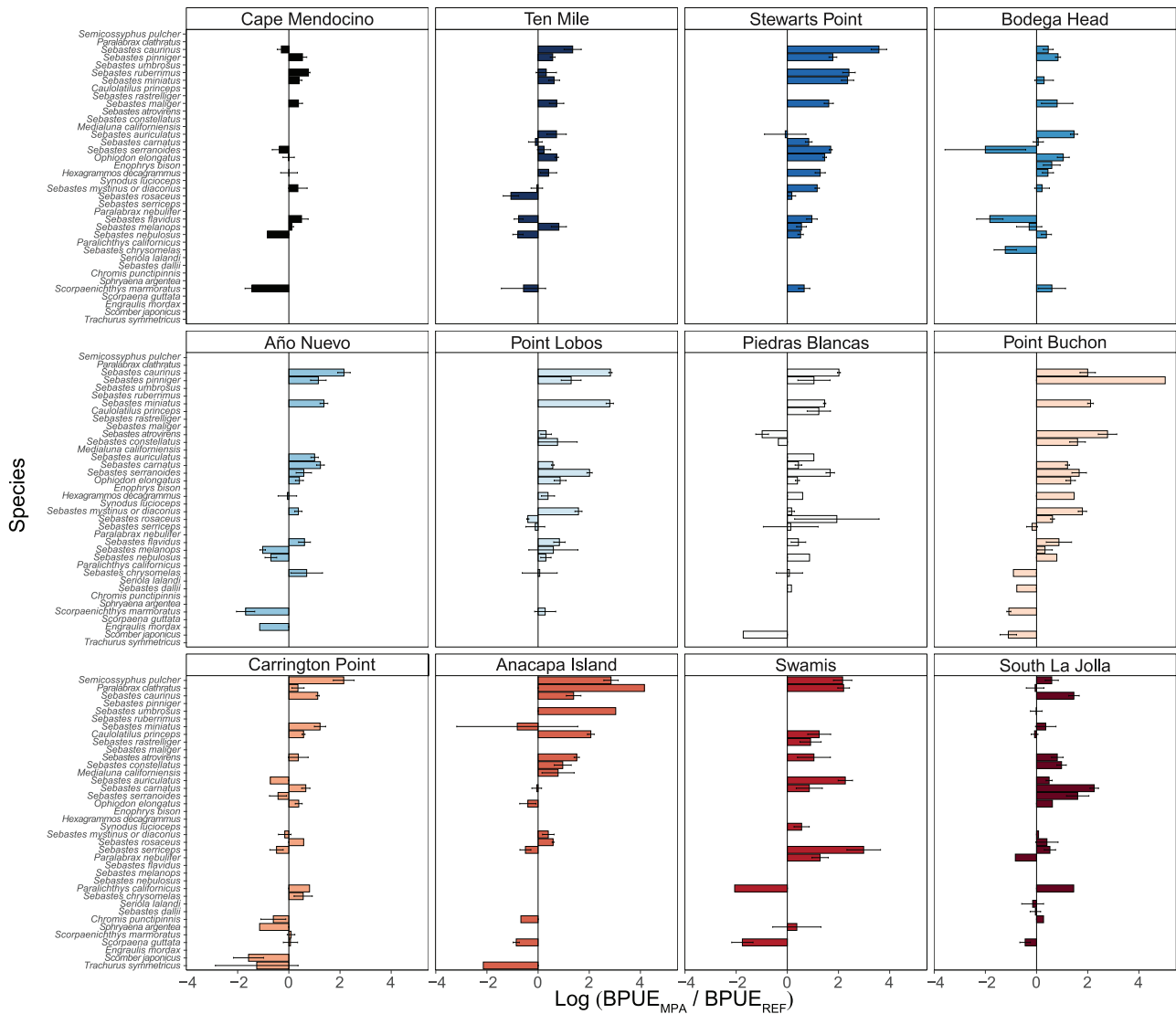


FIGURE 5 Tornado plots of species-specific average BPUE response ratios for all MPAs surveyed between 2017 and 2022. Values > 0 indicate a positive species response (i.e., higher biomass inside MPA relative to REF), while values < 0 indicate a negative species response. Paired MPA/REF sites are ordered and colored by latitude (and relative differences in sea surface temperature) (blue = northern and cool, red = southern and warm). Bars are the mean response ratio per species and errors are $\pm 95\%$ CI.

slowly. McClanahan et al. (2007) reported that densities of coral reef fish in Kenya may reach an asymptote anywhere from 5 to 10 years or to up to 37 years after MPA protection, while Russ and Alcala (2010) demonstrated that predatory fish biomass saturated after 20–40 years of protection in the Philippines. While it is possible that total fish abundance and biomass in MPAs across this network may saturate relatively quickly, the studies that found rapid recovery examined tropical species that are likely to have much higher growth rates than those in temperate waters in California.

We also found that the differences in fish CPUE and BPUE between MPAs and REFs were greater in locations where the MPAs were larger. Theoretical studies have hypothesized that larger MPAs would be more effective at

increasing biodiversity (Krueck et al., 2018) and density of economically important species (Krueck et al., 2017). Past empirical studies have provided evidence that larger MPAs are characterized by greater increases in fish density and biomass (Claudet et al., 2008; Edgar et al., 2014). Larger MPAs may allow a higher proportion of fishes with larger home ranges to remain protected within the MPA, compared with smaller MPAs, where more mobile species may be likely to spillover into unprotected areas and be caught by anglers (Di Lorenzo et al., 2020). In contrast, other large-scale global syntheses of MPAs are more equivocal on the importance of MPA size (Guidetti & Sala, 2007) or report greater effects in smaller MPAs (Giakoumi et al., 2017).

A recent meta-analysis reported that edge effects in MPAs can effectively reduce the MPA area (Ohayon

et al., 2021). The authors concluded that fishing-the-line behavior—where fishers disproportionately fish the edge of a protected area—may be largely responsible. These MPA edge effects may be particularly pernicious for small MPAs (<10 km²), since the total area to perimeter ratio is small, providing more opportunities for fishing-the-line relative to the size of the MPA. Larger MPAs may also increase larval retention, promoting self-recruitment and the build-up of biomass inside the MPA boundary, increasing overall MPA efficacy (Krueck et al., 2017). Our results indicated that catch biomass was higher inside every MPA compared with its associated REF, with MPA effectiveness increasing further with increased with MPA size. Thus, while small MPAs are still effective at protecting fish populations, our statewide data revealed that larger ones within the network were even more effective.

Overall, we expected more rapid population responses and stronger MPA effects with decreasing latitude in the warmer waters of Southern California (Caselle et al., 2023) due to faster growth rates and higher recruitment of the dominant southern taxa (kelp bass, California sheephead, ocean whitefish) compared with northern taxa that generally have slower growth and more sporadic recruitment (rockfishes and lingcod) (Caselle et al., 2010). For example, Hamilton et al. (2010) observed rapid increases in biomass of southern taxa following the establishment of the Channel Islands MPAs, while Starr et al. (2015) and Ziegler et al. (2022) reported much slower responses of rockfishes in the initial years following MPA establishment in Central California. Consistent with our hypotheses, MPA responses tended to decline with increasing latitude ($\Delta\text{AICc} < 2$). Latitude is also negatively correlated with SST and our metric of fishing pressure, but positively correlated with NPP (Figure S4), making it challenging to disentangle which factor(s) are ultimately driving the latitudinal response to MPAs.

We also observed a weak positive effect of adjacent fishing pressure on the strength of MPA responses, especially the number of positive species responses; shown to be important in other California studies (Jaco & Steele, 2020; Ziegler et al., 2022) and globally (Griffiths et al., 2022). Fishing pressure is much more intense in southern California (~1 million recreational anglers), and much of the coast is easily accessible. In contrast, human population density and recreational fishing pressure are lower in central and northern California and there are smaller, more disparate ports in these regions, creating more remote sections of the coastline that are much less accessible to boat-based anglers (Free et al., 2023). The limited strength of the relationship between fishing pressure and MPA response ratios in this study may be due to a variety of factors. Most importantly, the fishing pressure metric (e.g., landings / distance from port) was likely too coarse to provide

strong predictive power at the individual MPA scale. Still, we observed a positive association between the strength of the MPA response and fishing effort, similar to Ziegler et al. (2022) on the central California coast, indicating that fishing pressure is positively correlated with MPA responses across the larger scale of the statewide network. Higher fishing pressure in reference sites equates with bigger differences in catches and biomass inside MPAs compared with those fished areas. These results emphasize the need for high quality, spatially explicit fishing pressure data to better assess the effectiveness of MPAs across large spatial scales.

Overall, our results—using standardized data collected through a collaborative fisheries research program, CCFRP—support and expand much of the previous literature on the attributes that result in positive effects of MPAs to enhance abundance and biomass of targeted species. Our results show that targeted fish abundance increased in MPAs throughout the network, and that MPA size, age, and likely some combination of latitude, fishing pressure, and temperature had significant effects on the strength of the MPA response. Critically, this partnership of academic and agency scientists with the sportfishing industry and local anglers is able to generate high quality, reliable, and transparent information on the status of species, communities, and ecosystems protected by MPAs essential for adaptive management. While meta-analytical frameworks and studies can provide insight into global patterns of MPA efficacy, standardized sampling programs that engage stakeholders and span MPA networks can provide essential data for local and regional marine resource management. Our approach combining community engagement and a standardized and statistically rigorous sampling design not only provides valuable insight for the design and adaptive management of MPA networks spanning large geographic scales, it provides a model framework for the engagement of local stakeholders, often wary of conservation strategies, into MPA monitoring and fisheries management.

AUTHOR CONTRIBUTION

Shelby L. Ziegler: Conceptualization; formal analysis; investigation; visualization; writing—original draft. **Rachel O. Brooks:** Investigation; data curation; visualization; writing—review and editing. **Lyall F. Bellquist:** Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Jennifer E. Caselle:** Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Steven G. Morgan:** Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Timothy J. Mulligan:** Conceptualization;

methodology; project administration; funding acquisition; writing—review and editing. **Benjamin I. Ruttenberg**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Brice X. Semmens**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Richard M. Starr**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Joe Tyburczy**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Dean E. Wendt**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Andre Buchheister**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Jose R. Marin Jarrin**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Christina Pasparakis**: Conceptualization; methodology; project administration; funding acquisition; writing—review and editing. **Salvador J. Jorgensen**: Investigation; data curation; writing—review and editing. **Jennifer A. Chiu**: Investigation; data curation; writing—review and editing. **Jordan Colby**: Investigation; data curation; writing—review and editing. **Connor L. Coscino**: Investigation; data curation; writing—review and editing. **Leon Davis**: Investigation; data curation; writing—review and editing. **Francine de Castro**: Investigation; data curation; writing—review and editing. **Jack T. Elstner**: Investigation; data curation; writing—review and editing. **Christopher Honeyman**: Investigation; data curation; writing—review and editing. **Erica T. Jarvis Mason**: Investigation; data curation; writing—review and editing. **Erin M. Johnston**: Investigation; data curation; writing—review and editing. **Sadie L. Small**: Investigation; data curation; writing—review and editing. **Jay Staton**: Investigation; data curation; writing—review and editing. **Grant T. Waltz**: Investigation; data curation; writing—review and editing. **Bonnie Basnett**: Investigation; data curation; writing—review and editing. **Erin V. Satterthwaite**: Investigation; data curation; writing—review and editing. **H. Killeen**: Investigation; data curation; writing—review and editing. **Connor D. Dibble**: Investigation; data curation; writing—review and editing. **Scott L. Hamilton**: Conceptualization; methodology; project administration; funding acquisition; writing—original draft.

ACKNOWLEDGMENTS

This research was possible by the investment and dedication of collaborating sportfishing vessel staff, captains, and crew, CCFRP science crew, and >1800 volunteer



anglers. This work was supported by the California Ocean Protection Council (C0752003), California Department of Fish and Wildlife (P1970018), and California Sea Grant (R/MPA-45). We would also like to thank historic funding sources and program partners (Supporting Information Text 2). Data are openly available in the California OPC DataOne Repository: <https://opc.dataone.org/view/urn%3Auuid%3Af843f110-e691-4d26-bf12-3854a4b641cd>. Data and code used for analyses are publicly available: <https://github.com/slziegler/CCFRP2017-2022Statewide>.

DATA AVAILABILITY STATEMENT

Data are openly available in the California OPC DataOne Repository: <https://opc.dataone.org/view/urn%3Auuid%3Af843f110-e691-4d26-bf12-3854a4b641cd>. Data and code used for analyses are publicly available: <https://github.com/slziegler/CCFRP2017-2022Statewide>.

ORCID

Shelby L. Ziegler  <https://orcid.org/0000-0001-7218-0811>
Rachel O. Brooks  <https://orcid.org/0000-0002-8443-699X>
Lyall F. Bellquist  <https://orcid.org/0000-0003-1247-0483>
Jennifer E. Caselle  <https://orcid.org/0000-0002-1364-3123>
Steven G. Morgan  <https://orcid.org/0000-0002-9971-5414>
Benjamin I. Ruttenberg  <https://orcid.org/0000-0001-7874-1915>
Brice X. Semmens  <https://orcid.org/0000-0001-5663-9194>
Richard M. Starr  <https://orcid.org/0000-0002-4010-5152>
Dean E. Wendt  <https://orcid.org/0000-0002-1732-5616>
Jose R. Marin Jarrin  <https://orcid.org/0000-0002-4474-8323>
Christina Pasparakis  <https://orcid.org/0000-0002-7173-9801>
Salvador J. Jorgensen  <https://orcid.org/0000-0002-4331-1648>
Jennifer A. Chiu  <https://orcid.org/0000-0002-7800-594X>
Jack T. Elstner  <https://orcid.org/0009-0000-8610-449X>
Christopher Honeyman  <https://orcid.org/0000-0003-0152-1775>
Erica T. Jarvis Mason  <https://orcid.org/0000-0003-3366-157X>
Erin M. Johnston  <https://orcid.org/0000-0002-1094-5708>
Sadie L. Small  <https://orcid.org/0000-0003-3874-5924>
Grant T. Waltz  <https://orcid.org/0000-0001-7879-5390>
Bonnie Basnett  <https://orcid.org/0000-0001-9019-7270>
Erin V. Satterthwaite  <https://orcid.org/0000-0003-0177-7770>

Helen Killeen  <https://orcid.org/0000-0002-6942-7315>
 Scott L. Hamilton  <https://orcid.org/0000-0001-5034-4213>

REFERENCES

- Babcock, R. C., Shears, N. T., Alcalá, A. C., Barrett, N. S., Edgar, G. J., Lafferty, K. D., Mcclanahan, T. R., & Russ, G. R. (2010). Decadal trends in marine reserves reveal differential rates of change in direct and indirect effects. *PNAS*, *107*, 18256–18261.
- Ban, N. C., Gurney, G. G., Marshall, N. A., Whitney, C. K., Mills, M., Gelcich, S., Bennett, N. J., Meehan, M. C., Butler, C., Ban, S., Tran, T. C., Cox, M. E., & Breslow, S. J. (2019). Well-being outcomes of marine protected areas. *Nat Sustain*, *2*, 524–532.
- Bartoń, K. (2023). *MuMIn: Multi-Model Inference*. R package version 1.47.5, <https://CRAN.R-project.org/package=MuMIn>
- Bates, D., Mächler, M., Bolker, B., & Walker, S. (2015). Fitting linear mixed-effects models using lme4. *Journal of Statistical Software*, *67*, 1–48.
- Botsford, L. W., White, J. W., Carr, M. H., & Caselle, J. E. (2014). Chapter six—Marine protected area networks in California, USA. In *Advances in marine biology, marine managed areas and fisheries*. (pp. 205–251) (eds. M. L. Johnson, & J. Sandell). Academic Press.
- Burnham, K. P., & Anderson, D. R. (2002). *Model selection and multimodel inference: A practical information-theoretic approach*. Springer.
- Caselle, J., Nickols, K., Smith, J., Lopazanski, C., Brun, J., Anderson, C., Carr, M., Claudet, J., Dugan, J., Eurich, J., Francis, T., Free, C., Gill, D., Hamilton, S., Kaschner, K., Mouillot, D., Raimondi, P., Starr, R., & Ziegler, S. (2023). *A synthesis of ecological and social outcomes from the California marine protected area (MPA) network*. <https://doi.org/10.25494/P6430V>
- Caselle, J. E., Rassweiler, A., Hamilton, S. L., & Warner, R. R. (2015). Recovery trajectories of kelp forest animals are rapid yet spatially variable across a network of temperate marine protected areas. *Scientific Reports*, *5*, 14102.
- Caselle, J. E., Wilson, J. R., Carr, M. H., Malone, D. P., & Wendt, D. E. (2010). Can we predict interannual and regional variation in delivery of pelagic juveniles to nearshore populations of rockfishes (genus *Sebastes*) using simple proxies of ocean conditions? *Reports of California Cooperative Oceanic Fisheries Investigations*, *51*, 91–105.
- Cinner, J. E., Maire, E., Huchery, C., Macneil, M. A., Graham, N. A. J., Mora, C., Mcclanahan, T. R., Barnes, M. L., Kittinger, J. N., Hicks, C. C., D'agata, S., Hoey, A. S., Gurney, G. G., Feary, D. A., Williams, I. D., Kulbicki, M., Vigliola, L., Wantiez, L., Edgar, G. J., ... Mouillot, D. (2018). Gravity of human impacts mediates coral reef conservation gains. *Proceedings of the National Academy of Sciences of the United States of America*, *115*, E6116–E6125.
- Claudet, J., Osenberg, C. W., Benedetti-Cecchi, L., Domenici, P., García-Charton, J.-A., Pérez-Ruzafa, Á., Badalamenti, F., Bayle-Sempere, J., Brito, A., Bulleri, F., Culioli, J.-M., Dimech, M., Falcón, J. M., Guala, I., Milazzo, M., Sánchez-Meca, J., Somerfield, P. J., Stobart, B., Vandeperre, F., ... Planes, S. (2008). Marine reserves: Size and age do matter. *Ecology Letters*, *11*, 481–489.
- Claudet, J., Osenberg, C. W., Domenici, P., Badalamenti, F., Milazzo, M., Falcón, J. M., Bertocci, I., Benedetti-Cecchi, L., García-Charton, J.-A., Goñi, R., Borg, J. A., Forcada, A., De Lucia, G. A., Pérez-Ruzafa, Á., Afonso, P., Brito, A., Guala, I., Diréach, L. L., Sanchez-Jerez, P., ... & Planes, S. (2010). Marine reserves: Fish life history and ecological traits matter. *Ecological Applications*, *20*, 830–839.
- Development Core Team, R. (2023). *R: A language and environment for statistical computing*. R Foundation for Statistical Computing.
- Di Lorenzo, M., Guidetti, P., Di Franco, A., Calò, A., & Claudet, J. (2020). Assessing spillover from marine protected areas and its drivers: A meta-analytical approach. *Fish and Fisheries*, *21*, 906–915.
- Edgar, G. J., & Barrett, N. S. (2012). An assessment of population responses of common inshore fishes and invertebrates following declaration of five Australian marine protected areas. *Environmental Conservation*, *39*, 271–281.
- Edgar, G. J., Stuart-Smith, R. D., Willis, T. J., Kininmonth, S., Baker, S. C., Banks, S., Barrett, N. S., Becerro, M. A., Bernard, A. T. F., Berkhout, J., Buxton, C. D., Campbell, S. J., Cooper, A. T., Davey, M., Edgar, S. C., Försterra, G., Galván, D. E., Irigoyen, A. J., Kushner, D. J., ... Thomson, R. J. (2014). Global conservation outcomes depend on marine protected areas with five key features. *Nature*, *506*, 216–220.
- Free, C. M., Smith, J. G., Lopazanski, C. J., Brun, J., Francis, T. B., Eurich, J. G., Claudet, J., Dugan, J. E., Gill, D. A., Hamilton, S. L., Kaschner, K., Mouillot, D., Ziegler, S. L., Caselle, J. E., & Nickols, K. J. (2023). If you build it, they will come: Coastal amenities facilitate human engagement in marine protected areas. *People and Nature*, <https://doi.org/10.1002/pan3.10524>
- Free, C. M., Vargas Poulsen, C., Bellquist, L. F., Wassermann, S. N., & Oken, K. L. (2022). The CALFISH database: A century of California's non-confidential fisheries landings and participation data.
- Froese, R., & Pauly, D. (2023). *FishBase. Fisheries centre*. University of British Columbia Vancouver.
- Giakoumi, S., Scianna, C., Plass-Johnson, J., Micheli, F., Grorud-Colvert, K., Thiriet, P., Claudet, J., Di Carlo, G., Di Franco, A., Gaines, S. D., García-Charton, J. A., Lubchenco, J., Reimer, J., Sala, E., & Guidetti, P. (2017). Ecological effects of full and partial protection in the crowded Mediterranean Sea: A regional meta-analysis. *Scientific Reports*, *7*, 8940.
- Gill, D. A., Mascia, M. B., Ahmadi, G. N., Glew, L., Lester, S. E., Barnes, M., Craigie, I., Darling, E. S., Free, C. M., Geldmann, J., Holst, S., Jensen, O. P., White, A. T., Basurto, X., Coad, L., Gates, R. D., Guannel, G., Mumby, P. J., Thomas, H., ... Fox, H. E. (2017). Capacity shortfalls hinder the performance of marine protected areas globally. *Nature*, *543*, 665–669.
- Griffiths, L. L., Andradi-Brown, D. A., Ahmadi, G. N., Purwanto, Awaludinnoer, Bryan-Brown, D., & Brown, C. J. (2022). Linking historical fishing pressure to biodiversity outcomes to predict spatial variation in marine protected area performance. *Marine Policy*, *139*, 105024.
- Grorud-Colvert, K., Claudet, J., Tissot, B. N., Caselle, J. E., Carr, M. H., Day, J. C., Friedlander, A. M., Lester, S. E., De Loma, T. L., Malone, D., & Walsh, W. J. (2014). Marine protected area networks: Assessing whether the whole is greater than the sum of its parts. *PLoS ONE*, *9*, e102298.
- Grorud-Colvert, K., Sullivan-Stack, J., Roberts, C., Constant, V., Horta E Costa, B., Pike, E. P., Kingston, N., Laffoley, D., Sala, E., Claudet, J., Friedlander, A. M., Gill, D. A., Lester, S. E., Day, J. C., Gonçalves, E. J., Ahmadi, G. N., Rand, M., Villagomez, A., Ban, N. C., ... Lubchenco, J. (2021). The MPA guide: A framework to achieve global goals for the ocean. *Science*, *373*, eabf0861.

- Guidetti, P., & Sala, E. (2007). Community-wide effects of marine reserves in the Mediterranean Sea. *Marine Ecology Progress Series*, 335, 43–56.
- Halpern, B. S., & Warner, R. R. (2003). Matching marine reserve design to reserve objectives. *Proceedings of the Royal Society of London. Series B: Biological Sciences*, 270, 1871–1878.
- Hamilton, S. L., Caselle, J. E., Malone, D. P., & Carr, M. H. (2010). *Incorporating biogeography into evaluations of the Channel Islands marine reserve network*. (Vol. 107, pp. 18272–18277) PNAS.
- Harborne, A. R., Green, A. L., Peterson, N. A., Beger, M., Golbuu, Y., Houk, P., Spalding, M. D., Taylor, B. M., Terk, E., Treml, E. A., Victor, S., Vigliola, L., Williams, I. D., Wolff, N. H., Zu Ermgassen, P. S. E., & Mumby, P. J. (2018). Modelling and mapping regional-scale patterns of fishing impact and fish stocks to support coral-reef management in Micronesia. *Diversity and Distributions*, 24, 1729–1743.
- Jaco, E. M., & Steele, M. A. (2020). Pre-closure fishing pressure predicts effects of marine protected areas. *Journal of Applied Ecology*, 57, 229–240.
- Krueck, N. C., Ahmadi, G. N., Possingham, H. P., Riginos, C., Treml, E. A., & Mumby, P. J. (2017). Marine reserve targets to sustain and rebuild unregulated fisheries. *PLoS Biology*, 15, e2000537.
- Krueck, N. C., Legrand, C., Ahmadi, G. N., Estradivari, Green, A., Jones, G. P., Riginos, C., Treml, E. A., & Mumby, P. J. (2018). Reserve sizes needed to protect coral reef fishes. *Conservation Letters*, 11, e12415.
- Leenhardt, P., Teneva, L., Kininmonth, S., Darling, E., Cooley, S., & Claudet, J. (2015). Challenges, insights and perspectives associated with using social-ecological science for marine conservation. *Ocean & Coastal Management*, 115, 49–60.
- Lester, S., & Halpern, B. (2008). Biological responses in marine no-take reserves versus partially protected areas. *Marine Ecology Progress Series*, 367, 49–56.
- Love, M. S., Morris, P., McCrae, M., & Collins, R. (1990). *Life history aspects of 19 rockfish species (Scorpaenidae: Sebastes) from the Southern California Bight*. NOAA Technical Report NMFS 87.
- Mason, E. T., Kellum, A. N., Chiu, J. A., Waltz, G. T., Murray, S., Wendt, D. E., Starr, R. M., & Semmens, B. X. (2020). Long-term participation in collaborative fisheries research improves angler opinions on marine protected areas. *PeerJ*, 8.
- McClanahan, T. R., & Graham, N. A. J. (2015). Marine reserve recovery rates towards a baseline are slower for reef fish community life histories than biomass. *Proceedings of the Royal Society B: Biological Sciences*, 282, 20151938.
- McClanahan, T. R., Graham, N. A. J., Calnan, J. M., & Macneil, M. A. (2007). Toward pristine biomass: Reef fish recovery in coral reef marine protected areas in Kenya. *Ecological Applications*, 17, 1055–1067.
- Meehan, M. C., Ban, N. C., Devillers, R., Singh, G. G., & Claudet, J. (2020). How far have we come? A review of MPA network performance indicators in reaching qualitative elements of Aichi Target 11. *Conservation Letters*, 13, e12746.
- Mouillot, D., Velez, L., Maire, E., Masson, A., Hicks, C. C., Moloney, J., & Troussellier, M. (2020). Global correlates of terrestrial and marine coverage by protected areas on islands. *Nature Communications*, 11, 4438.
- Mulligan, T., Tyburczy, J., Staton, J., Kelmartin, I., & Barrett, D. (2017). *Baseline characterization of fish communities associated with nearshore rocky reefs in the Northern California Marine Protected Area study regions (Final Report)*. <https://caseagrant.ucsd.edu/sites/default/files/37-Mulligan-Final.pdf>
- Naeem, S., Lu, Y., & Jackson, J. (2022). Curtailing the collapse of the living world. *Science Advances*, 8, ead9317.
- Nickols, K. J., White, J. W., Malone, D., Carr, M. H., Starr, R. M., Baskett, M. L., Hastings, A., & Botsford, L. W. (2019). Setting ecological expectations for adaptive management of marine protected areas. *Journal of Applied Ecology*, 56, 2376–2385.
- Ohayon, S., Granot, I., & Belmaker, J. (2021). A meta-analysis reveals edge effects within marine protected areas. *Nature Ecology & Evolution*, 5, 1301–1308.
- Russ, G. R., & Alcala, A. C. (2010). Decadal-scale rebuilding of predator biomass in Philippine marine reserves. *Oecologia*, 163, 1103–1106.
- Russ, G. R., & Alcala, A. C. (2011). Enhanced biodiversity beyond marine reserve boundaries: The cup spillith over. *Ecological Applications*, 21, 241–250.
- Starr, R. M., Wendt, D. E., Barnes, C. L., Marks, C. I., Malone, D., Waltz, G., Schmidt, K. T., Chiu, J., Launer, A. L., Hall, N. C., & Yochum, N. (2015). Variation in responses of fishes across multiple reserves within a network of marine protected areas in temperate waters. *PLoS ONE*, 10, e0118502.
- Stewart, G. (2010). Meta-analysis in applied ecology. *Biology Letters*, 6, 78–81.
- Wendt, D. E., & Starr, R. M. (2009). Collaborative research: An effective way to collect data for stock assessments and evaluate marine protected areas in California. *Marine and Coastal Fisheries*, 1, 315–324.
- White, J. W., Botsford, L. W., Baskett, M. L., Barnett, L. A., Barr, R. J., & Hastings, A. (2011). Linking models with monitoring data for assessing performance of no-take marine reserves. *Frontiers in Ecology and the Environment*, 9, 390–399.
- Yochum, N., Starr, R. M., & Wendt, D. E. (2011). Utilizing fishermen knowledge and expertise: Keys to success for collaborative fisheries research. *Fisheries*, 36, 593–605.
- Ziegler, S. L., Brooks, R. O., Hamilton, S. L., Ruttenberg, B. I., Chiu, J. A., Fields, R. T., Waltz, G. T., Shen, C., Wendt, D. E., & Starr, R. M. (2022). External fishing effort regulates positive effects of no-take marine protected areas. *Biological Conservation*, 269, 109546.

SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.

How to cite this article: Ziegler, S. L., Brooks, R. O., Bellquist, L. F., Caselle, J. E., Morgan, S. G., Mulligan, T. J., Ruttenberg, B. I., Semmens, B. X., Starr, R. M., Tyburczy, J., Wendt, D. E., Buchheister, A., Jarrin, J. R. M., Pasparakis, C., Jorgensen, S. J., Chiu, J. A., Colby, J., Coscino, C. L., Davis, L., . . . , Hamilton, S. L. (2024). Collaborative fisheries research reveals reserve size and age determine efficacy across a network of marine protected areas. *Conservation Letters*, 17, e13000. <https://doi.org/10.1111/conl.13000>