Short-term management of kelp forests for marine heatwaves requires pre-emptive action

Jess K. Hopf [a]*, Anita Giraldo-Ospina [b], Jennifer E. Caselle [b], Kristy J. Kroeker [c], Mark H. Carr [c], Louis W. Botsford [d], Alan Hastings [e], J. Wilson White [a,f]

[a] Coastal Oregon Marine Experiment Station, Oregon State University, Newport, Oregon USA

[b] Marine Science Institute, University of California, Santa Barbara, California, USA

[c] Department of Ecology and Evolutionary Biology, University of California, Santa Cruz, California, USA.

[d] Department of Wildlife, Fish, and Conservation Biology, University of California, Davis, California, USA

[e] Department of Environmental Science and Policy, University of California, Davis, California, USA and Santa Fe Institute, Santa Fe, New Mexico, USA

[f] Department of Fisheries, Wildlife, and Conservation Sciences, Oregon State University, Newport, Oregon, USA

*Corresponding author:

Jess Hopf

jess.k.hopf@gmail.com

+61 473 350 870

2030 SE Marine Science Dr,

Newport OR 97365 USA

Data availability statement: All modelling code and relevant data is available here [link provided upon acceptance].

Conflict of interest disclosure: The authors have no conflicts of interest to declare.

Abstract

Heatwaves are now pervasive stressors to marine ecosystems globally and it is urgent to consider mitigation tools that support ecosystem resilience and persistence in the immediate future. We modeled a system of kelp, herbivorous urchin, and predatory fish to compare how potential management mitigation actions (kelp seeding, urchin removal, and fishery closures) could reduce the likelihood of a heatwave shifting a kelp forest into a degraded urchin barren state. We found that those interventions must commence at or before the start of a heatwave to be effective. Closing the predatory fish fishery was more effective when done earlier and for longer while urchin removal and kelp seeding needed to be undertaken immediately before, and continue throughout the heatwave. Our results suggest the need for improved heatwave forecasting and nimble management protocols to enact mitigation actions quickly if a heatwave is forecasted or occurs.

Keywords: Marine heatwaves; mitigation actions; resilience; ecosystem persistence; modelling; marine protected areas; fishery management; kelp restoration; urchin removals;

Introduction

The acceleration of climate change has come with an urgent need to understand what local actions can mitigate impacts. In particular, the increased frequency, intensity and duration of marine heatwaves (hereafter 'heatwave(s)') are a threat to marine systems and the ecosystem services they provide (Frölicher et al. 2018, Smith et al. 2021, 2023). Heatwaves can be defined as discrete pulses of extreme ocean temperatures occurring against the background of chronic warming (Hobday et al. 2016, Amaya et al. 2023). While long-term management responses are required to address long-term temperature trends, short-term, temporary interventions (our focus here) are critical to support systems through discrete heatwave events. The key questions are what management actions will be most effective, and when to implement them (Ainsworth et al. 2020, Pinsky et al. 2021, Smith et al. 2021, 2023).

Heatwaves can stress certain species while benefiting others, resulting in complex ecosystem responses. In northeastern Pacific kelp forests, warm, low-nutrient conditions during heatwaves reduce growth and recruitment of the dominant species, giant kelp (*Macrocystis pyrifera*) (Zimmerman and Kremer 1986, Hollarsmith et al. 2020, Michaud et al. 2022). Conversely, sea urchins (key herbivores; *Mesocentrotus franciscanus* and *Strongylocentrotus purpuratus*) experience increased grazing rates (Spindel 2023). The decrease in standing kelp also reduces production of detached kelp fronds ('drift kelp'), a preferred food source for urchins (Kriegisch et al. 2019, Randell 2022). The loss of drift kelp combined with increased grazing rates can trigger urchins to switch behavior, from cryptic feeding on drift and detritus in rocky crevices to exposed roaming and targeting standing kelp stipes (Kriegisch et al. 2019, Rennick et al. 2022; Figure 1). As a result, heatwaves can sometimes tip the ecosystem from a kelp-dominated forest to an urchin barren (McPherson et al. 2021). Recovery from the degraded barren state can be challenging even after returning to kelp-beneficial conditions (i.e., the system exhibits hysteresis; Filbee-Dexter and Scheibling 2014, Ling et al. 2015) largely because the barren-state urchins have high survival, despite low food availability and exposure to predators. In fact, some predators actively avoid starved, barren-state urchins (Eurich et al. 2014, Liebergesell 2022). Thus, kelp resilience to heatwaves is a fine balance of consumer-resource dynamics.

Kelp forests are ecologically and economically important, and we need to understand how to mitigate heatwave effects on them (McPherson et al. 2021, Hamilton et al. 2022, Eger et al. 2023, CDFW 2023). Management actions can be broadly grouped by which link in the kelp-urchin-predator trophic chain they address (Figure 1). First, direct kelp restoration aims to increase kelp biomass via seeding spores or transplanting sporophytes (Eger et al. 2022). Second, urchin removal can promote kelp recovery by removing the key herbivore via harvest, culling, or translocation (Eger et al. 2022). Third, because many urchin predators (e.g., fishes, lobsters, sea otters) have declined due to human harvest,

increasing predator densities through fishery closures can suppress herbivory and support kelp forest
persistence (Eisaguirre et al. 2020, Kumagai et al. 2024).

Here we focus on potential short-term management actions: temporary changes enacted in response to
an impending, current, or recently-ended heatwave that may prevent a shift from a kelp forest to an
urchin barren (i.e., ecosystem resistance). We help address the management question, 'what action(s)
could be taken in the face of a heatwave and when?' This reflects a proactive or reactive response to
sudden, acute changes in the system rather than permanent management actions that would better
orient the system towards longer-term goals (Pinsky et al. 2021, Arroyo-Esquivel et al. 2023). To do
this, we analyzed a three-species model that captures the interactions among giant kelp, purple urchins
(*Strongylocentrotus purpuratus*), and a fishery-targeted urchin predator (Californian Sheephead;
Bodianus pulcher, henceforth 'Sheephead') (Figure 1). We use Southern Californian (USA) kelp
forests as our study system, but our results are applicable to kelp systems globally where urchin
predators are targeted by fisheries. We consider the ability of three main management actions (kelp
seeding, urchin removal, and a temporary predator fishery closure) to reduce the likelihood of a kelp
forest shifting to an urchin barren because of a heatwave when the actions are implemented before,
during, or after a 2-year heatwave. This analysis complements studies that consider these management
actions only after forests have shifted to barrens states (e.g., Dunn et al. 2021, Arroyo-Esquivel et al.
2023).

Figure 1: Three-species model overview. Main processes included in our kelp-urchin-predator
model, including those affected by heatwaves (orange symbol). Purple and green text show processes
relevant to the kelp-urchin sub-model, and blue text show processes relevant to the predator sub-
model. Bold, black text indicates potential mitigation actions considered in this study. All artwork by

Jess K. Hopf, except giant kelp icons, which are modified from artwork by Jane Thomas sourced from

ian.umces.edu/media-library under the CC BY-SA 4.0 license.

Methods

Three-species population model

We developed a three-species, single patch (spatially-implicit) population model that captures the

interactions between giant kelp, herbivorous purple urchins, and a fishery-targeted predatory fish

(Californian Sheephead) (Figure 1). The model is comprised of two sub-models that run over time: 1)

a kelp-urchin stage-based sub-model and 2) a predatory fish integral-projection sub-model. To

account for seasonal variations in ecological process (recruitment, kelp growth, urchin grazing, and

heatwave effects), the model uses seasonal (3-month) time steps. Details are provided in the

Supplemental Information, and all code is publicly available at repository [xxx].

Our model captured the characteristic kelp forest dynamic in which overgrazing by urchins shifted the

system from a kelp forest to an urchin barren with local kelp extinction. Underpinning this was an

urchin behavioral feeding switch: when drift kelp was abundant, urchins fed passively in crevices on

drift, but left crevices to actively graze on standing kelp when drift kelp was scarce. To achieve this,

we modelled the proportion of exposed urchins as a declining function of drift biomass density

relative to urchin consumptive capacity (Rennick et al. 2022), building on Randell's (2022) approach.

We assumed a Type-II functional response for kelp grazing by urchins, and a Type-I response for

urchin predation by Sheephead, as a function of the total biomass of Sheephead large enough (>20

cm; Hamilton and Caselle 2015) to consume adult urchins. Exposed urchins had a higher rate of

predation mortality than cryptic urchins (Nichols et al. 2015). However, if there was no kelp for >3

months then Sheephead stopped consuming urchins, reflecting the reduced nutritional value of starved

urchins (Liebergesell 2022). We assume a spatial domain equivalent to a spatially isolated kelp forest,

at which scale the Sheephead and urchin populations would be open with external recruitment, and

the kelp population would be largely closed (Kinlan and Gaines 2003), with recruitment a function of

local adult biomass and density dependence through shading and competition (Nisbet and Bence

1989). This can be viewed as a worst-case scenario for kelp resilience without rescue effects from

external kelp populations. Stochasticity was captured through year-to-year variation in recruitment for

all species.

Modelling heatwaves & management strategies

All modelled scenarios began in an undisturbed kelp forest state. Reflecting the known effects of

heatwaves on kelp and urchins, we implemented a heatwave as a period of reduced kelp recruitment

(Hollarsmith et al. 2020) and growth (Zimmerman and Kremer 1986), and increased urchin grazing

rates (Spindel 2023). The modelled heatwave started in winter and lasted 2 years, following the recent

2014-2016 Californian heatwave (Michaud et al. 2022).

We considered three possible management actions: a temporary Sheephead fishery closure, urchin

removal (a set biomass of adult urchins removed each season), and kelp seeding (a set biomass of

juvenile kelp added each season). We focused on kelp seeding, rather than transplantation of adult

kelp, as seeding is more viable on a large scale (Eger et al. 2022). We modelled scenarios where

actions began up to two years before the heatwave (i.e., pre-emptive actions), during the heatwave, or

up to three years after the heatwave ended, and actions lasted one to five years. We also explored a

range of magnitudes for urchin removal and kelp seeding. Finally, we considered the implications of

simultaneously implementing combinations of two management actions.

We simulated 10^5 replicates of each scenario, running the model for 20 years prior to the heatwave

and then assessing the proportion of simulations in which a kelp forest persisted (defined as non-zero

kelp biomass for at least a year) 8 years after the heatwave ceased (enough time to complete all

potential management actions). Monitoring sooner only reduced the realized impact of a temporary

fishery closure (Supplemental Information). We present this response variable as *the percentage*

increase in probability of kelp forest persistence relative to the effect of the heatwave with no

management intervention, and to a baseline scenario with no heatwave (Figure 2), such that:

percentage increase in probability of persistence $= \frac{P_{MHW \& action} - P_{MHW \& no\ action}}{P_{no\ MHW \& no\ action} - P_{MHW \& no\ action}} * 100$

where P_i is the proportion of simulations with persistent kelp forests under scenario *i*. Note that even

without a heatwave, the proportion of kelp forests persisting declined over time (Figure 2) simply

because in a stochastic system (and without kelp or drift immigration) there is a small, constant

probability of switching from kelp forest to barren in any given time step.

Figure 2: Example scenario runs. The proportion of model simulations (n = 10,000) with kelp forests persisting over time, and our response variable, *the percentage increase in probability of kelp forest persistence*, calculated at the sampling year (eight years post-heatwave), for example scenario runs. In this example, the mitigation actions began the same year as the heatwave started, and lasted three years. For urchin removal and kelp seeding, 10% of the pre-heatwave biomass was removed/added yearly.

Results

The three standalone management actions considered were most effective at increasing the probability of a kelp forest persisting through the heatwave if initiated prior to, or during the first year of the heatwave (Figures 3-4). This was generally robust to changes in key parameter values (based on sensitivity analyses; see Supplemental Information). All three management actions had low to no efficacy (<25% increase in probability of persistence) if started after the heatwave finished (Figure 3). Fishery closures were more effective the earlier and longer they were implemented: for example, closures that began 6+ months before the heatwave and lasted a minimum of 3 years were highly effective (>100% increase in the relative probability of kelp persistence) (Figure 3a). In contrast, urchin removal and kelp seeding needed to overlap completely with the heatwave to be highly effective (Figure 2b-c, Figure 3). Both kelp and urchin actions had low to no efficacy (<25% increase in probability of persistence) if completed before the heatwave began (bottom left of Figure 3b-c), or started towards the end of the heatwave (Figure 3b-c).

While longer and earlier fishery closures were more beneficial, the same was not true for urchin
culling and kelp removal (Figures 3-4). So long as urchin removal and kelp seeding spanned the
heatwave, their effectiveness did not depend on timing or duration (Figure 3-4). For example, a 2-year
action begun at the onset of the 2-year heatwave and a 4-year action begun 2 years prior were equally
very effective (Figure 3). Figure 3 presents the most intensive management scenario in which 100%
of the pre-heatwave urchin and kelp biomass is removed or seeded each year, respectively. However,
that intense effort was not necessary: there was little additional benefit to removing more than 50% of
the pre-heatwave urchin biomass each season, or to seeding 50% of the pre-heatwave kelp biomass,
regardless of timing or length of the management action (vertical dashed lines; Figure 4).

Reflecting the different optimal timings for the management interventions, fishery closures that began
a year before the heatwave were generally more effective than urchin or kelp actions implemented at
the same time (Figure 4a-b). For actions that started concurrent to, or a year after, the heatwave,
fishery closures were generally more effective than urchin or kelp actions that removed or seeded less
than 20-60% of the pre-heatwave biomass, but equally or less effective than more intensive urchin or
kelp actions (dotted lines and dots in Figure 4c-f).

Combining management actions of varying degrees had additive effects, at best, but typically reached
a threshold beyond which additional management efforts provided little benefit (Figure 5). We
demonstrate this by focusing on the scenario where 3-yr long actions are implemented at the start of
the heatwave. When combining two actions (kelp seeding plus urchin removal, or fishery closure and
either kelp seeding or urchin removal), the highest gains for combining the actions occurred with low
degrees of kelp and urchin management actions (<10% additional or removal of kelp and urchins,
respectively; Figure 5). At this point, the effects of combined actions were essentially additive.
Beyond this point, however, the additional benefits of the second management action were less than
expected, based on the effect of that action alone (Figure 5).

Figure 3: Efficacy of individual heatwave mitigation actions. Percentage increase in the probability of kelp forest persistence through a 2-year marine heatwave with individual management actions, over length and timing of action. These modelled scenarios present the most intensive management scenario in which 100% of the pre-heatwave urchin and kelp biomass is removed or seeded each year, respectively. The orange dashed box indicates the action scenarios that began during the heatwave. Note that all actions are enacted temporarily.

Figure 4: Changes in efficacy with effort. Percentage increase in the probability of kelp forest persistence through a 2-year marine heatwave with varying degrees of urchin removal (a, c, e,) and keep seeding efforts (b, d, f), and lengths of actions (colors), at different times (*t,* in years) relative to the start of the heatwave (rows). Dotted lines with points indicate efficacy for fishery closures starting at the same time; lines without points indicate that temporary fishery closures were more effective than 100% kelp seeding or urchin removal efforts. Degrees of effort are measured as the percentage of pre-heatwave urchin and kelp biomass removed or seeded each year, respectively.

Figure 5: Efficacy of combining two heatwave mitigation actions. Modelled percentage increases
(±10 units) in the probability of a kelp forest persisting through a 2-year marine heatwave (numbers in
cells), and the proportional difference between the efficacy outcome of the combined actions
compared to the expected summed outcome of the actions based on their individual efficacy (grey
color scale), over a range of urchin culling and kelp seeding efforts. In this demonstrative scenario,
actions occur simultaneously at the start of the heatwave and last for the same period of time (3
years). Blue, purple, and green boxes show the efficacy of individual actions for comparison for
Sheephead, urchins and kelp respectively (see also Figure 3). As an example (boxes outlined in
orange), in the lower right corner of the figure, adding 100% kelp seeding and 1% urchin culling has
an *additive* effect (102% effect of kelp alone + 1% effect of urchin alone ≈ 105%, the combined effect
of both) but 100% kelp seeding plus a fishery closure is *less than additive* (102% effect of kelp alone
+ 84% effect of fish alone < 145%, the combined effect). The degree of urchin removal and kelp
seeding effort is the percentage of pre-heatwave urchin and kelp biomass removed or seeded each
year, respectively. Note that all actions are enacted temporarily.

Discussion

Short-term, temporary management actions may be essential to supporting kelp forests through heatwaves. We found that timing is critical: mitigations needed to commence before a heatwave (for temporary fishery closures) or at the start of a heatwave (for urchin culling and kelp seeding) to be highly effective in reducing the probability that a kelp forest would shift to an urchin barren (measured 8 years after the heatwave ended; but see also Supplementary Information). As heatwave forecasts are currently available only months beforehand at best (Holbrook et al. 2020, Jacox et al. 2022), our results suggest that action plans (e.g., protocols, permits, and funding) need to be in place preemptively, so that mitigation actions can be enacted quickly. This also highlights the need to continue to develop longer, more robust forecast warnings so that effective actions can be implemented in time (Holbrook et al. 2020, Jacox et al. 2022). As forecasted heatwaves are not a certainty, cost-benefit analyses are needed to characterize potential trade-offs between the costs of undertaking an action that was not needed (if a potential heatwave does not occur) and failing to undertake an action that was needed. Such analyses need to consider the logistical and monetary costs as well as the local management goals and social acceptance of different actions, which should be assessed on a case-by-case basis.

Like others (Dunn et al. 2021, Arroyo-Esquivel et al. 2023), our findings reinforce the need to consider underlying ecological processes in management plans. To buffer against kelp forest collapse, fishery closures require years (depending on the life history of the species; Kaplan et al. 2019) to rebuild predator biomass to levels sufficient to suppress urchin densities and prevent overgrazing during a heatwave. Therefore, we found longer and earlier fishery closures more effective, such as could be achieved with a permanent no-take marine reserve (White et al. 2025). As preemptive management actions are unlikely, or potentially too costly/risky, to occur before a heatwave starts, and highly unlikely to occur earlier than 1 year prior to a heatwave (due to forecast limitations), the full effectiveness of a temporary fishery closure may be limited, especially compared to large-scale, highly effective urchin removal or kelp seeding efforts. However, temporary fishery closures that start at the beginning of the heatwave can be as effective as low-to-moderate urchin removal or kelp seeding actions. Managers would need to consider the costs, trade-offs, and feasibility at different spatial scales, of those alternatives when assessing options. Fishery adjustments (e.g., quotas, closures, and restocking) have already been implemented in response to heatwaves, although often retrospectively and with maintaining stock biomass rather than ecosystem persistence in mind (Smith et al. 2021). Nimble ecosystem-based fishery management could focus on regulations that maintain the desired level of urchin consumption by predators; the full closures we modeled could be viewed as a precautionary ecosystem-based intervention in cases where the amount of urchin predation needed for ecosystem persistence is uncertain.

In contrast to fishery closures, removing urchins or seeding juvenile kelp are relatively quick-acting

but short-lasting, logistically costly, and challenging at scale (Eger et al. 2020b, 2022). Our results

also show that these actions require ongoing intervention throughout the heatwave to be effective.

External larval supply for urchins can drive fast rebounds following removal, necessitating continuous

removal attempts through the heatwave. Likewise, kelp seeding success is dampened by density

dependence (here, we included shading of juveniles by the established kelp canopy; Nisbet and Bence

1989) and urchin grazing, quickly returning kelp biomass to pre-seeding levels if started and

completed too early. We also found that urchin removal and kelp seeding actions that occur prior to

and continue through heatwave are as effective as those that start the same year as the heatwave.

Consequently, starting these actions earlier may waste valuable resources. Although our model

included the effects of reduced kelp recruitment and growth during warmer temperatures, we assumed

kelp seeding efforts were always successful, even during heatwaves. More experimental work is

needed to test this assumption.

Interestingly, we found that combining two management actions did not always increase effectiveness

as much as could be assumed based on their individual outcomes, especially with large kelp seeding

and urchin removal efforts. The implications of this will depend on the per-kilogram cost of action,

but it suggests that there may be benefits to combining actions if increasing efforts of an individual

action yields diminishing returns, compared to undertaking two actions. There may also be benefits to

offsetting actions in time, such as an early fishery closure followed by urchin removal or kelp seeding.

However, testing all such possible combinations was outside the scope of this study. We recommend

that case-specific modelling and cost-benefits analysis be undertaken.

A key assumption of our model was a closed kelp population without larval rescue effects or external

supplies of drift kelp. We also assumed a constant influx of urchin recruitment, independent of the

local urchin densities. These assumptions consider the worst-case scenario for kelp resilience, and

drive the observed low success of post-heatwave management actions. However, we expect that our

general findings will still hold true for areas that do experience notable external kelp recruitment, as

such recruitment events are patchy in space and time (Cavanaugh et al. 2013) and should not be relied

upon, although they may benefit kelp-forest restoration efforts (Eger et al. 2020a). Lastly, we focused

on Sheephead, but our results would be applicable to other fishery-targeted predators like the Spiny

Lobster (*Panulirus interruptus*) or Southern Rock Lobster (*Jasus edwardsii*), which are also mobile

benthic invertivores that suppress kelp herbivory when protected from harvest (e.g., Peleg et al. 2023).

Many climate-mitigation options are being explored for a range of marine systems (e.g., Ainsworth et

al. 2020), with increasing focus on optimizing these actions for success (e.g., through site-selection, or

synergistic actions; Eger et al. 2020a, Brown et al. 2024). Our study demonstrates that the timing of a

mitigation action, relative to a heatwave, critically affects its success in reducing the likelihood of a

kelp-forest to urchin-barren shift. Most heatwave responses to date have been reactive (Smith et al.
2021) and our results suggest that managers should begin to also consider analyzing options and
establishing protocols for preemptive heatwave mitigation actions.

Acknowledgements

We thank Pete Raimondi, Mark Novak, Andrés Pinos-Sánchez, and Iris Flores for helpful thoughts
during model development and analysis. J. K. Hopf thanks Cashie the wolfhound-cross for her
supportive snoring during the writing of this manuscript. This work was funded by the David and
Lucile Packard Foundation (award 2022-74730) and the California Ocean Protection Council
(agreement C0874012). This is publication XX of the Partnership for Interdisciplinary Study of
Coastal Oceans, funded primarily by the David and Lucile Packard Foundation.

References

Ainsworth, T. D., C. L. Hurd, R. D. Gates, and P. W. Boyd. 2020. How do we overcome abrupt
degradation of marine ecosystems and meet the challenge of heat waves and climate
extremes? Global Change Biology 26:343–354.

Amaya, D. J., M. G. Jacox, M. R. Fewings, V. S. Saba, M. F. Stuecker, R. R. Rykaczewski, A. C.
Ross, C. A. Stock, A. Capotondi, C. M. Petrik, S. J. Bograd, M. A. Alexander, W. Cheng, A.
J. Hermann, K. A. Kearney, and B. S. Powell. 2023. Marine heatwaves need clear definitions
so coastal communities can adapt. Nature 616:29–32.

Arroyo-Esquivel, J., M. L. Baskett, M. McPherson, and A. Hastings. 2023. How far to build it before
they come? Analyzing the use of the F ield of D reams hypothesis in bull kelp restoration.
Ecological Applications 33:e2850.

Brown, C. J., M. D. Campbell, C. J. Collier, M. P. Turschwell, M. I. Saunders, and R. M. Connolly.
2024. Speeding up the recovery of coastal habitats through management interventions that
address constraints on dispersal and recruitment. Proceedings of the Royal Society B:
Biological Sciences 291:20241065.

Cavanaugh, K. C., B. E. Kendall, D. A. Siegel, D. C. Reed, F. Alberto, and J. Assis. 2013. Synchrony

in dynamics of giant kelp forests is driven by both local recruitment and regional

environmental controls. Ecology 94:499–509.

CDFW. 2023, July. Marine Protected Area Decadal Management Review Report: Prioritized

Recommendations.

Dunn, R. P., J. F. Samhouri, and M. L. Baskett. 2021. Transient dynamics during kelp forest recovery

from fishing across multiple trophic levels. Ecological Applications 31:e02367.

Eger, A. M., E. Marzinelli, P. Gribben, C. R. Johnson, C. Layton, P. D. Steinberg, G. Wood, B. R.

Silliman, and A. Vergés. 2020a. Playing to the Positives: Using Synergies to Enhance Kelp

Forest Restoration. Frontiers in Marine Science 7:1–15.

Eger, A. M., E. M. Marzinelli, R. Beas-Luna, C. O. Blain, L. K. Blamey, J. E. K. Byrnes, P. E.

Carnell, C. G. Choi, M. Hessing-Lewis, K. Y. Kim, N. H. Kumagai, J. Lorda, P. Moore, Y.

Nakamura, A. Pérez-Matus, O. Pontier, D. Smale, P. D. Steinberg, and A. Vergés. 2023. The

value of ecosystem services in global marine kelp forests. Nature Communications 14:1894.

Eger, A. M., E. M. Marzinelli, H. Christie, C. W. Fagerli, D. Fujita, A. P. Gonzalez, S. W. Hong, J. H.

Kim, L. C. Lee, T. A. McHugh, G. N. Nishihara, M. Tatsumi, P. D. Steinberg, and A. Vergés.

2022. Global kelp forest restoration: past lessons, present status, and future directions.

Biological Reviews 97:1449–1475.

Eger, A. M., A. Vergés, C. G. Choi, H. Christie, M. A. Coleman, C. W. Fagerli, D. Fujita, M.

Hasegawa, J. H. Kim, M. Mayer-Pinto, D. C. Reed, P. D. Steinberg, and E. M. Marzinelli.

2020b. Financial and Institutional Support Are Important for Large-Scale Kelp Forest

Restoration. Frontiers in Marine Science 7:1–7.

Eisaguirre, J. H., J. M. Eisaguirre, K. Davis, P. M. Carlson, S. D. Gaines, and J. E. Caselle. 2020.

Trophic redundancy and predator size class structure drive differences in kelp forest

ecosystem dynamics. Ecology 101:1–11.

Eurich, J., R. Selden, and R. Warner. 2014. California spiny lobster preference for urchins from kelp

forests: implications for urchin barren persistence. Marine Ecology Progress Series 498:217–

225.

Filbee-Dexter, K., and R. E. Scheibling. 2014. Sea urchin barrens as alternative stable states of

collapsed kelp ecosystems. Marine Ecology Progress Series 495:1–25.

Frölicher, T. L., E. M. Fischer, and N. Gruber. 2018. Marine heatwaves under global warming. Nature

560:360–364.

Hamilton, S. L., and J. E. Caselle. 2015. Exploitation and recovery of a sea urchin predator has

implications for the resilience of southern California kelp forests. Proceedings of the Royal

Society B: Biological Sciences 282.

Hamilton, S. L., M. G. Gleason, N. Godoy, N. Eddy, and K. Grorud-colvert. 2022. Ecosystem-based

management for kelp forest ecosystems. Marine Policy 136:1–10.

Hobday, A. J., L. V. Alexander, S. E. Perkins, D. A. Smale, S. C. Straub, E. C. J. Oliver, J. A.

Benthuysen, M. T. Burrows, M. G. Donat, M. Feng, N. J. Holbrook, P. J. Moore, H. A.

Scannell, A. Sen Gupta, and T. Wernberg. 2016. A hierarchical approach to defining marine

heatwaves. Progress in Oceanography 141:227–238.

Holbrook, N. J., A. Sen Gupta, E. C. J. Oliver, A. J. Hobday, J. A. Benthuysen, H. A. Scannell, D. A.

Smale, and T. Wernberg. 2020. Keeping pace with marine heatwaves. Nature Reviews Earth

& Environment 1:482–493.

Hollarsmith, J. A., A. H. Buschmann, C. Camus, and E. D. Grosholz. 2020. Varying reproductive

success under ocean warming and acidification across giant kelp (Macrocystis pyrifera)

populations. Journal of Experimental Marine Biology and Ecology 522:151247.

Jacox, M. G., M. A. Alexander, D. Amaya, E. Becker, S. J. Bograd, S. Brodie, E. L. Hazen, M. Pozo

Buil, and D. Tommasi. 2022. Global seasonal forecasts of marine heatwaves. Nature

604:486–490.

Kaplan, K. A., L. Yamane, L. W. Botsford, M. L. Baskett, A. Hastings, S. Worden, and J. W. White.

2019. Setting expected timelines of fished population recovery for the adaptive management

of a marine protected area network. Ecological Applications 29:1202–1220.

Kinlan, B. P., and S. D. Gaines. 2003. Propagule dispersal in marine and terrestrial environments: a

community perspective. Ecology 84:2007–2020.

Kriegisch, N., S. E. Reeves, E. B. Flukes, C. R. Johnson, and S. D. Ling. 2019. Drift-kelp suppresses

foraging movement of overgrazing sea urchins. Oecologia 190:665–677.

Kumagai, J. A., M. C. Goodman, J. C. Villaseñor-Derbez, D. S. Schoeman, K. C. Cavanuagh, T. W.

Bell, F. Micheli, G. De Leo, and N. Arafeh-Dalmau. 2024. Marine Protected Areas That

Preserve Trophic Cascades Promote Resilience of Kelp Forests to Marine Heatwaves. Global

Change Biology 30:e17620.

Liebergesell, M. W. 2022. Evaluating mechanisms of prey selection by a fish predator and their sea

urchin prey linked to habitat-specific differences in kelp forests and urchin barrens.pdf.

Masters, San Diego State University.

Ling, S. D., R. E. Scheibling, A. Rassweiler, C. R. Johnson, N. Shears, S. D. Connell, A. K. Salomon,

K. M. Norderhaug, A. Pérez-Matus, J. C. Hernández, S. Clemente, L. K. Blamey, B. Hereu,

E. Ballesteros, E. Sala, J. Garrabou, E. Cebrian, M. Zabala, D. Fujita, and L. E. Johnson.

2015. Global regime shift dynamics of catastrophic sea urchin overgrazing. Philosophical

Transactions of the Royal Society B: Biological Sciences 370:20130269.

McPherson, M. L., D. J. I. Finger, H. F. Houskeeper, T. W. Bell, M. H. Carr, L. Rogers-Bennett, and

R. M. Kudela. 2021. Large-scale shift in the structure of a kelp forest ecosystem co-occurs

with an epizootic and marine heatwave. Communications Biology 4:1–9.

Michaud, K. M., D. C. Reed, and R. J. Miller. 2022. The Blob marine heatwave transforms California

kelp forest ecosystems. Communications Biology 5:1143.

Nichols, K. D., L. Segui, and K. A. Hovel. 2015. Effects of predators on sea urchin density and

habitat use in a southern California kelp forest. Marine Biology 162:1227–1237.

Nisbet, R. M., and J. R. Bence. 1989. Alternative Dynamic Regimes for Canopy-Forming Kelp: A

Variant on Density-Vague Population Regulation. The American Naturalist 134:377–408.

Peleg, O., C. O. Blain, and N. T. Shears. 2023. Long-term marine protection enhances kelp forest

ecosystem stability. Ecological Applications 33:e2895.

Pinsky, M. L., E. Fenichel, M. Fogarty, S. Levin, B. McCay, K. St. Martin, R. L. Selden, and T.

Young. 2021. Fish and fisheries in hot water: What is happening and how do we adapt?

Population Ecology 63:17–26.

Randell, Z. 2022. Grazer-resource Behavior, Interactions, and Feedbacks Control Kelp-forest

Dynamics and Stability.

Rennick, M., B. P. DiFiore, J. Curtis, D. C. Reed, and A. C. Stier. 2022. Detrital supply suppresses

deforestation to maintain healthy kelp forest ecosystems. Ecology 103:e3673.

Smith, K. E., M. T. Burrows, A. J. Hobday, N. G. King, P. J. Moore, A. Sen Gupta, M. S. Thomsen,

T. Wernberg, and D. A. Smale. 2023. Biological Impacts of Marine Heatwaves. Annual

Review of Marine Science 15:119–145.

Smith, K. E., M. T. Burrows, A. J. Hobday, A. Sen Gupta, P. J. Moore, M. Thomsen, T. Wernberg,

and D. A. Smale. 2021. Socioeconomic impacts of marine heatwaves: Global issues and

opportunities. Science 374:eabj3593.

Spindel, N. B. 2023. Ecophysiology of ectothermic ecosystem engineers: Bioenergetic effects of

climate and food on dominant consumers and their consequences for coastal ecosystems.

White, J. W., J. K. Hopf, N. Arafeh-Dalmau, N. C. Ban, A. E. Bates, J. Claudet, C. Lopazanski, J. M.

Sunday, and J. E. Caselle. 2025. Measurements, mechanisms, and management

recommendations for how marine protected areas can provide climate resilience. Marine

Policy 171:106419.

Zimmerman, R., and J. Kremer. 1986. In situ growth and chemical composition of the giant kelp,

Macrocystis pyrifera: response to temporal changes in ambient nutrient availability. Marine

Ecology Progress Series 27:277–285.

439