

REVIEW SUMMARY

CLIMATE CHANGE

Socioeconomic impacts of marine heatwaves: Global issues and opportunities

Kathryn E. Smith, Michael T. Burrows, Alistair J. Hobday, Alex Sen Gupta, Pippa J. Moore, Mads Thomsen, Thomas Wernberg, Dan A. Smale*

BACKGROUND: Anthropogenic climate change is altering the structure and functioning of ecosystems globally. The upper layers of the oceans have absorbed >90% of the excess heat arising from human activities and, as a result, surface waters have warmed at an average rate of 0.15°C per decade over the past 40 years, with global ocean heat content currently at a record high. Many marine species have shifted their distributions in response to this ubiquitous warming, causing community reconfigurations and changes to entire ecosystems. In addition to gradual warming, the frequency and severity of short-term oceanic warming events, marine heatwaves (MHWs), has increased substantially, largely as a consequence of rising anthropogenic carbon emissions and ocean warming. The global annual number of MHW days has risen by 54% over the past century, with eight of the 10 most extreme MHWs ever recorded occurring after 2010. MHWs are forceful agents of disturbance with wide-ranging impacts on marine ecosystems, including driving species range shifts and mass mortalities and altering foodwebs and species interactions.

Regional case studies have shown that ecological responses to MHWs can have socioeconomic implications such as loss of fisheries income, erosion of essential ecosystem services, mass mortalities of iconic species, and conflict. Although our understanding of the physical drivers and biological impacts from MHWs are becoming increasingly robust and predictive, only a few studies have examined their consequences for human-ocean interactions, a crucial step in understanding the associated costs and risks and in developing potential adaptation and mitigation measures.

ADVANCES: Using a large number of case studies, we provide a global perspective on the impacts of MHWs on the provision of ecosystem services and examine the socioeconomic implications. We show that these discrete warming events are altering ecosystem functioning with present-day ramifications for human-ocean interactions. Ecological impacts commonly include harmful algal blooms, species range shifts, mass mortality events, and ecosystem reconfiguration lasting from weeks

to years or longer. Such ecological impacts affect humans by changing provisioning, habitat, regulating, and cultural ecosystem services. Clear socioeconomic consequences have been linked to at least 34 MHWs occurring across a wide range of ecosystems in all major ocean basins globally. Impacts on provisioning services relating to fisheries and habitat services relating to loss of foundation species (e.g., corals, kelps, and seagrasses) have been particularly prevalent. Impacts on cultural services relating to mass mortality of charismatic (e.g., sea lions, sea birds, and corals) and spiritual (e.g., certain kelp) species, and on regulating services relating to carbon sequestration and altered nutrient cycling were also common. Biological responses to extreme warming events can, however, also increase human-ocean interactions and offer short-term socioeconomic opportunities through, for example, increased tourism or fisheries prospects. Economic losses of single MHW events to date exceed US\$800 million in direct losses and in excess of US\$3.1 billion per annum in indirect losses for multiple consecutive years. The true costs are, however, likely to be much greater because many socioeconomic effects likely remain unknown and underreported, particularly in lower-income countries.

OUTLOOK: The likelihood and intensity of MHWs have increased substantially in recent decades, largely due to broad-scale warming attributed to anthropogenic climate change, and are projected to intensify in coming decades. Concurrently, reported ecological impacts, changes in ecosystem services, and economic losses have also increased markedly. We discuss potential adaptation and mitigation measures to reduce risks and consequences for human-ocean interactions and examine knowledge gaps in our current understanding of MHW impacts, for example, in relation to the likely exacerbating effects of other anthropogenic stressors. Globally, marine ecosystems are threatened by a diverse range of anthropogenic stressors, and policy, management, and conservation measures must prioritize approaches to alleviating them to ensure continued provision of ecosystem services. We demonstrate the far-reaching impacts of MHWs on present-day human-ocean interactions and highlight the urgent need to develop a coherent understanding of the linked social-ecological impacts of MHWs and robust approaches to mitigation and adaptation. ■



MHWs have caused widespread loss of habitat-forming species, such as reef-building corals, severely compromising ecosystem service provision.

The list of author affiliations is available in the full article online.
*Corresponding author. Email: dansma@mba.ac.uk
Cite this article as K. E. Smith *et al.*, *Science* 374, eabj3593 (2021). DOI: 10.1126/science.abj3593

S READ THE FULL ARTICLE AT
<https://doi.org/10.1126/science.abj3593>

REVIEW

CLIMATE CHANGE

Socioeconomic impacts of marine heatwaves: Global issues and opportunities

Kathryn E. Smith¹, Michael T. Burrows², Alistair J. Hobday³, Alex Sen Gupta⁴, Pippa J. Moore⁵, Mads Thomsen^{6,7}, Thomas Wernberg^{8,9}, Dan A. Smale^{1*}

Extreme climatic events, including marine heatwaves (MHWs), are altering ecosystems globally, often with profound socioeconomic impacts. We examine how MHWs have affected the provision of ecosystem services and evaluate the socioeconomic consequences for human society. Ecological impacts range from harmful algal blooms and mass mortality events to reconfigurations of entire ecosystems, affecting provisioning, habitat, regulating, and cultural ecosystem services globally. Reported economic costs of individual MHW events exceed US\$800 million in direct losses or >US\$3.1 billion in indirect losses of ecosystem services for multiple years. However, biological responses to MHWs can also increase human-ocean interactions, providing opportunities for coastal societies. Our study provides a global perspective on the far-reaching impacts of MHWs on human societies and highlights the urgent need to develop robust approaches to mitigation and adaptation.

Anthropogenic climate change is altering the structure and functioning of ecosystems globally (1, 2). The upper layers of the oceans have absorbed >90% of the excess heat arising from human activities and, as a result, surface waters have warmed at an average rate of 0.15°C per decade over the past 40 years (3), with rates two to three times higher in some regions (4). Marine species have shifted their distributions in response to gradual warming trends, causing community reconfiguration and changes to entire ecosystems (5, 6). In addition to gradual long-term ocean warming, the frequency and severity of discrete warming events, marine heatwaves (MHWs), have increased substantially, with a rise in global annual MHW days of 54% over the past century (7). Of the 10 most severe MHWs ever recorded, eight occurred in the most recent decade (8). Although MHWs are driven by a range of complex oceanographic and atmospheric processes (9), their increasing severity over time can largely be explained by long-term increases in mean ocean temperature caused by anthropogenic greenhouse

gas emissions, and they are therefore unequivocally projected to intensify in coming decades (7, 10, 11).

High temperatures associated with MHWs have been shown to induce physiological stress as thermal thresholds are increasingly exceeded, leading to reduced viability or increased mortality, changes in population structure and species' distributions, and altered ecosystem structure and functioning (11). Indeed, ecosystem-wide impacts of MHWs have been reported from a number of regions where foundation species that support a wide range of associated biodiversity have been affected, including loss of kelp forests in the northeast and southwest Pacific and southeast Indian Oceans (12–15), coral bleaching in the Indo-Pacific and southwest Indian Ocean (16–18), widespread mortality of habitat-forming invertebrates in the Mediterranean Sea (19), and depleted extent and resilience of seagrass meadows in the southeast Indian Ocean (20, 21). MHWs have also been linked to mass dies-offs of invertebrates, fish, seabirds, and marine mammals (22–26); redistributions of commercially important fisheries species (27, 28); and disrupted food webs and carbon and nitrogen cycles (21, 29). An analysis of eight high-profile MHWs showed that all had negative impacts across a range of taxa and ecological functions (30). Responses to extreme climatic events are highly variable, however, with some species or populations showing little response and others even responding positively by extending their ranges (28, 31) or proliferating as a result of increasingly favorable conditions or competitive release (13, 28, 32, 33). Broadly, populations found toward their species' poleward range edge are more likely to respond positively to MHW events (30), and species exhibiting

certain life history traits (e.g., high mobility, generalist feeding, or wide distribution) may proliferate more rapidly with warming (34). Crucially, regional species losses or gains may be either temporary or persistent, depending on a variety of factors such as the characteristics of the MHW, the wider environmental and climatological context, and ecological interactions (11). Although our understanding of the physical drivers and biological impacts of MHWs is becoming increasingly robust and predictive, only a few scattered regional studies have examined their consequences on linked human-ocean interactions [e.g., (35–37)], and those that have typically reported regional losses in excess of hundreds of millions USD from individual MHW events, often with acute societal responses (28, 38, 39). Conversely, MHWs have also resulted in economic gains, for example, through increased fisheries or tourism opportunities (28, 31, 32, 37).

We draw on a large number of case studies to provide a global perspective on MHW-associated risks, costs, and opportunities, and use it to guide priorities and management decisions that counter or take advantage of more frequent and severe MHWs in the future. We review the linked impacts of MHWs globally to quantitatively examine associated changes in ecosystem service provision and evaluate socioeconomic consequences. Through analyses of global datasets, we identify a broad range of biological response variables related to ecosystem services and investigate variability across 10-year periods spanning prominent MHWs. We classify responses by associated ecosystem services using The Economics of Ecosystems and Biodiversity framework [TEEB; (40)] alongside a coherent framework to identify and describe MHWs (41, 42), allowing for direct comparisons of events, regions, and type of impacts. Specifically, an event is considered a MHW if it lasts for 5 or more days, with temperatures warmer than the 90th percentile based on a 30-year historical climatology (41). We then examine the impacts of MHWs in the context of United Nations Sustainable Development Goals (SDGs), because these extreme events will affect the achievement of several of these goals that seek improved socioeconomic outcomes for the planet. Finally, we explore possible adaptation measures and highlight future directions for research.

Impacts of MHWs on ecosystem services

We found compelling evidence for MHW impacts globally, on all major types of ecosystem services, with extensive socioeconomic consequences (Tables 1 and 2 and Fig. 1). Over the past 25 years, at least 34 MHW events have affected provisioning, regulating, habitat, and cultural ecosystem services (Fig. 1). For example, MHWs spanning the Indo Pacific associated with the extreme 1997/1998 El Niño caused

¹Marine Biological Association of the United Kingdom, The Laboratory, Citadel Hill, Plymouth, PL1 2PB, UK. ²Scottish Association for Marine Science, Oban, Argyll PA37 1QA, UK. ³CSIRO Oceans and Atmosphere, Hobart, Tasmania 7000, Australia. ⁴Climate Change Research Centre, University of New South Wales, Sydney, New South Wales 2052, Australia. ⁵School of Natural and Environmental Sciences, Newcastle University, Newcastle-Upon-Tyne NE1 7RU, UK. ⁶The Marine Ecology Research Group, Centre of Integrative Ecology, School of Biological Sciences, University of Canterbury, 8041 Christchurch, New Zealand. ⁷Department of Bioscience, Aarhus University, 4000 Roskilde, Denmark. ⁸University of Western Australia, Oceans Institute and School of Biological Sciences, Crawley, Western Australia 6009, Australia. ⁹Institute of Marine Research, Floedevigen, 4817 His, Norway.

*Corresponding author. Email: dansma@mba.ac.uk

Table 1. Examples of direct economic outcomes of MHWs.

Location	Year(s)	Socioeconomic impact	Location of impact	Value/loss/gain	Primary industry affected	Affected TEEB ecosystem service	Reference(s)
Gulf of Alaska	2014–2019	Population decline and low recruitment affects Gulf of Alaska commercial Pacific cod fishery catch limits	Alaska, USA	Fishery valued at US\$103 million per annum; loss unknown; funding relief: In 2018, Congress appropriated US\$24.4 million in federal disaster relief funding.*	Commercial fisheries	Provisioning and habitat services	(29)
	2020	Population decline and low recruitment closes Gulf of Alaska commercial Pacific cod fishery for the season	Alaska, USA	Fishery valued at US\$103 million per annum; comparative loss likely; federal disaster relief funding requested (pending as of 5 May 2021)*	Commercial fisheries	Provisioning and habitat services	(29, 48)
Northeast Pacific Ocean	2015	Harmful algal blooms cause closure of recreational razor clamming	Washington, USA	US\$40 million loss in tourist spending	Tourism and recreational fisheries	Regulating and cultural	(49, 66)
	2015–2016	Harmful algal blooms cause closure of commercial Dungeness crab fisheries	West coast, USA	US\$97.5 million loss; funding relief: Congress to appropriate US\$27.3 million in federal disaster relief funding including US\$1.5 million to the native American Quileute Tribe*	Commercial fisheries	Provisioning and regulating	(67, 103)
	2015–present	Population decline of red sea urchin due to kelp loss causes closure of commercial fishery	California, USA	Fishery valued at US\$3 million per annum; funding relief: Congress appropriated US\$3.3 million in federal disaster relief funding for 2016 and 2017*	Commercial fisheries	Provisioning, regulating, habitat, and cultural	(14)
	2018–present	Population decline of abalone due to kelp loss causes closure of recreational fishery	California, USA	US\$44 million loss per annum	Tourism and recreational fishing	Provisioning, regulating, habitat, and cultural	(14)
Southeast Pacific Ocean	2016–2017	Mass mortality of farmed salmon due to harmful algal blooms	Southern Chile	Export loss of US\$800 million	Aquaculture	Provisioning and regulating	(38)
Coral Sea	2016,–2018	Bleaching and mass mortalities of corals	Great Barrier Reef, Australia	The Great Barrier Reef is valued at US\$4.2 billion annually, with a total value estimated at US\$41 billion; loss related to bleaching unknown; gains related to “last chance tourism” also unknown	Tourism	Habitat and cultural	(69)

continued on next page

Location	Year(s)	Socioeconomic impact	Location of impact	Value/loss/gain	Primary industry affected	Affected TEEB ecosystem service	Reference(s)
Tasman Sea	2016	Pacific oyster mortality syndrome closed hatcheries and decimated juvenile stocks	East coast of Tasmania, Australia	Aquaculture valued at US\$19 million; loss unknown	Aquaculture	Provisioning and habitat	(104)
	2016	Poor performance of salmon limited production	East coast of Tasmania, Australia	Aquaculture valued at US\$545 million; loss unknown	Aquaculture	Provisioning and habitat	(104)
	2016	Mortality of wild caught abalone led to smaller catch and reduced quotas	South/southeast coast of Tasmania, Australia	Wild-caught fishery valued at US\$62 million; loss unknown	Fisheries	Provisioning and habitat	(104)
Southeast Asia seas	2010	Bleaching and mass mortalities of corals	Indonesia, Malaysia, Thailand	US\$49 to 74 million loss	Tourism	Habitat and cultural	(73)
Southeast Indian Ocean	2011–present	Closure of abalone fishery due to mass mortality event	West Coast, Australia	Estimated loss per annum ~US\$0.16 million	Commercial and recreational fisheries	Provisioning, habitat, and cultural	(51)
	2011–2013	Closure of commercial swimmer crab fishery for 18 months to protect breeding stock after low catch rates	Shark Bay, Australia	US\$3.1 million loss	Commercial fisheries	Provisioning and habitat	(105)
	2011–2016	Closure of commercial scallop fishery for 3 years (Shark Bay) or 5 years (Abrolhos Islands and Midwest) due to low recruitment	West Coast, Australia	Estimated losses per annum: Shark Bay ~US\$8.2 million [†] ; Abrolhos Islands and Midwest ~US\$3 million [†]	Commercial fisheries	Provisioning and habitat	(51)
	2011	Loss of carbon storage and other ecosystem services provided by seagrass	Shark Bay, Western Australia, Australia	US\$3.1 billion loss per annum for multiple consecutive years	Multiple ecosystem services	Provisioning, regulating, habitat, and cultural	(21)
	2012	Early lobster migration led to record landings and drop in lobster value	Gulf of Maine, USA	US\$38 million loss	Commercial fisheries	Provisioning and habitat	(77)
	2016	Proactive management of lobster fishery after 2012 MHW led to economic gains	Gulf of Maine, USA	US\$108 million gain	Commercial fisheries	Provisioning and habitat	(77)

*Funding relief values in the United States are from the National Oceanic and Atmospheric Administration (NOAA; <https://www.fisheries.noaa.gov/national/funding-and-financial-services/fishery-disaster-determinations>). †Values estimated from the Government of Western Australia State of the fisheries annual reports (<https://www.fish.wa.gov.au/About-Us/Publications/Pages/State-of-the-Fisheries-report.aspx>).

extensive coral bleaching and depleted zooplankton abundance, leading to mass mortalities of a range of marine mammals and seabirds. Ultimately, the disruption of ecosystem functioning had major ramifications for tourism and fishing across the Pacific (43). Similarly, the Mediterranean Sea has been repeatedly affected by strong MHWs, causing mass mortality of seagrass, corals, invertebrates, and fish and population explosions of invasive

species and pathogens (19, 24, 44, 45). These responses have markedly reduced provisioning (e.g., fisheries), cultural (e.g., mass mortalities of iconic species), regulating (e.g., carbon capture, water quality), and habitat (or supporting, e.g., restructuring, disease) ecosystem services (19, 24, 43–45) (Fig. 1), which, collectively, are of global socioeconomic importance.

The most pervasive socioeconomic impacts of MHWs reported to date relate to provision-

ing services, particularly fisheries, which directly influence markets, employment, food availability, and wider ecosystem services (Figs. 1 and 2). Impacts on provisioning services often closely align with cultural services by affecting tourism and recreational fisheries. In the northeast Pacific, “the blob” was associated with multiple regional MHWs between 2014 and 2016 that affected ecosystem services (46). In the Gulf of Alaska, the zooplankton community shifted

EMBARGOED UNTIL 2PM U.S. EASTERN TIME ON THE THURSDAY BEFORE THIS DATE:

Table 2. Reported impacts of MHW events. Responses in **bold** indicate positive socioeconomic consequences, those in *italics* are negative consequences, and those in roman are both negative and positive.

Location	Year(s)	Recorded response variable	Reference(s)
Southeast Atlantic (Benguela Niño)	1995	Mass mortalities of fish, southward displacement of fish species	(22)
South Indian Ocean	1997	Mortality and reproductive failure in king penguins, range shift in prey species	(106, 107)
East Pacific El Niño	1997–1998	Shift in fisheries species, mass mortalities and reproductive failure of Galapagos sea lions, decline in kelp forests, low zooplankton abundance, coral bleaching, ecosystem disruption	(26, 27, 43, 108, 109)
West Indian Ocean	1998	Coral bleaching, restructuring of ecosystems	(61, 110)
Red Sea	1998, 2007, 2010, 2012, 2015, 2017	Coral bleaching Mass mortalities of coral reef fish, fish disease	(111) (112)
Mediterranean Sea	1999	Mass mortalities of benthic organisms, habitat loss, reduced growth and reproduction, coral bleaching	(19, 24, 113, 114)
Mediterranean Sea	2003	Mortalities in mollusk fisheries (multiple life stages), mass mortalities of benthic organisms, coral bleaching, seagrass flowering and shoot mortality, reduced growth and reproduction, disease	(24, 44, 45, 114–116)
Caribbean Sea	2005	Coral bleaching, loss of fisheries	(16, 117, 118)
Mediterranean Sea	2006	Mortalities in mollusk fisheries (multiple life stages), mass mortalities of benthic organisms, shoot mortality in seagrass	(24, 44, 45)
Canadian Arctic	2009	Earlier sea ice breakup, decreased sea ice, increased productivity, increased cod reproduction	(32, 33)
Bay of Bengal	2010	Coral bleaching	(16, 17)
Southeast Indian Ocean (Ningaloo Niño)	2011	Loss of seagrass and kelp, coral bleaching, fisheries closures, reduced fisheries quotas, shifts in recruitment, range expansion of tropical fish	(15, 31, 36, 50, 51, 119–121)
Northwest Atlantic	2012	Spring boom in lobster landings, range shift of a range of commercially valuable fisheries species (e.g., squid, cod, flounder, hake)	(35, 39, 77, 84)
Great Australian Bight	2013	Mass mortalities of fish and abalone, harmful algal blooms, disease	(122)
Northeast Pacific (The Blob)	2014–2016	Mass mortalities of California sea lions, seals, seabirds, and marine invertebrates; sea star wasting disease; range shifts in a variety of species; harmful algal blooms; fisheries disruptions and closures; shifts in kelp forest ecosystems; increase in orca births; coral bleaching; increased abundance of tuna; unusual sightings of warm water species ; increases and decreases in fisheries recruitment; increased observations of whales; increased whale entanglement	(25, 28, 38, 123–125)
Northwest Atlantic	2015–2016	Loss of seagrass, early migration of lobsters	(58, 77)
Southwest Atlantic	2015	Harmful algal blooms, mass fish mortalities, closure of fisheries, flags flown on beaches to indicate potential health risks	(126)
Central Pacific	2015–2016	Mass coral bleaching and mortality, shift from coral to encrusting macroalgae and crustose coralline algae, decreased fish biomass, reduction in seabirds and seabird breeding	(61, 127, 128)

continued on next page

Location	Year(s)	Recorded response variable	Reference(s)
Southeast Indian Ocean	2015–2016	Coral bleaching	(121, 129)
Tasman Sea	2015–2016	Loss of fisheries, loss of kelp, POMS, mortality of abalone, fish species noted outside of their range	(74, 130)
Bering Sea	2016	Range expansion of bowhead whales, mass mortality of puffins, low sea ice, reduced ice-based fishing, low snow and unsafe river ice making winter transportation difficult, topicalization of copepods	(131)
Southern Ocean	2016	Decreased sea ice	(132)
Southeast Pacific	2016–2017	Mass mortalities of farmed salmon, harmful algal blooms	(38, 65)
Southeast Asia seas	2016–2017	Mass coral bleaching, reefs closed to tourism	(132)
Coral Sea	2016–2018	Mass coral bleaching, mass coral mortalities, failed coral recruitment, species range shifts, loss of seagrass, poor reproductive effort in seagrass, increased abundance and biomass of coral trout	(37, 53, 121, 133–138)
Tasman Sea	2017	Loss of kelp, range shift of fish and jellyfish, early spawning in snapper, salmon mortalities in fish farms	(13, 139)
East China Sea	2017	Harmful algal blooms, coral bleaching, mass mortalities of farmed fish	(140–142)
Southwest Atlantic	2017	Harmful algal blooms, mass mortalities of fish, closure of recreational beaches	(143, 144)
West Pacific	2017	Mass mortality of coral after successive bleaching events in 2013, 2014, 2016, and 2017	(61, 145)
Persian Gulf	2017	Mass coral bleaching, coral mortality, increase in turf algae	(146)
Northeast Pacific	2018	Loss of kelp	(12)
Northwest Atlantic	2018	Loss of kelp forest	(55)
North and Baltic Seas	2018	Loss of kelp forest	(55)
Northeast Pacific	2019–2021	Harmful algal blooms, Dungeness crab fisheries closures, recreational razor clam fisheries closures, fish mortalities at fish farms, olive Ridley sea turtles observed off British Columbia, coral bleaching	(38, 147–150)

from cold-water, lipid-rich copepods to less nutritious warm-water species, reducing food availability and thus the abundances of ground-fish, including Pacific cod (*Gadus macrocephalus*) (29) and Alaskan pollock (*Gadus chalcogrammus*) (47), and resulting in a massive die-off of fish-eating seabirds, including common murrelets (*Uria aalge*) (25) (Fig. 2A). The effect of the MHW on the life cycle of cod persisted for >5 years, as recruitment rates and spawning biomass remained well below pre-MHW levels (29, 48). This had major implications for the regional fishery and ecosystem services worth US\$103 million per year, with reduced quotas for several years and closure of the federal fishery in 2020 (49). Impacts of the blob were widespread: Throughout the West Coast, mass die-offs of finfish, shellfish, seabirds, sea lions, and seals were reported (25, 28); harmful algal blooms were extensive; and commercial and

recreational fisheries were closed (14, 49). Moreover, off the coast of California, a poleward range shift of market squid [*Doryteuthis (Loligo) opalescens*] resulted in higher and lower landings in northern and southern Californian counties, respectively, thereby altering supply chains and market values.

Other recent MHWs that have (positively or negatively) affected provisioning and cultural services include a MHW in the Canadian Arctic in 2009, which led to early sea ice breakup, resulting in increased primary productivity and enhanced polar cod (*Boreogadus saida*) recruitment (32). The Ningaloo Niño MHW in the southeast Indian Ocean during 2010–2011 (15, 50) caused poleward range extensions of tropical fish species, leading to a short-term boom in landings of valuable aquarium species (West Australian butterflyfish; *Chaetodon assarius*) off the coast of Perth, Western Australia

(Fig. 2B), and the opening of a new commercial rabbitfish fishery (*Siganus* spp.) in Cockburn Sound, Western Australia (31). At the same time, reduced growth rates of abalone (*Haliotis* spp.) led to a 50% drop in recreational landings near Perth (Fig. 2B), whereas farther north, mortalities and reduced recruitment of Roe's abalone (*Haliotis roei*), scallops (*Amusium balloti*), and blue swimmer crabs (*Portunus armatus*) necessitated closures affecting these fisheries for several years (36, 51). In the Gulf of Maine, a MHW in 2012 resulted in an early lobster (*Homarus americanus*) migration from deep to shallow water and an early molt to legal size (35). This led to high spring landings that were beyond processing capacity, causing a drop in lobster value (35, 39). Civil unrest followed as Canadian protests and blockades attempted to prevent imports of lobster from the United States into Canada for processing, and values

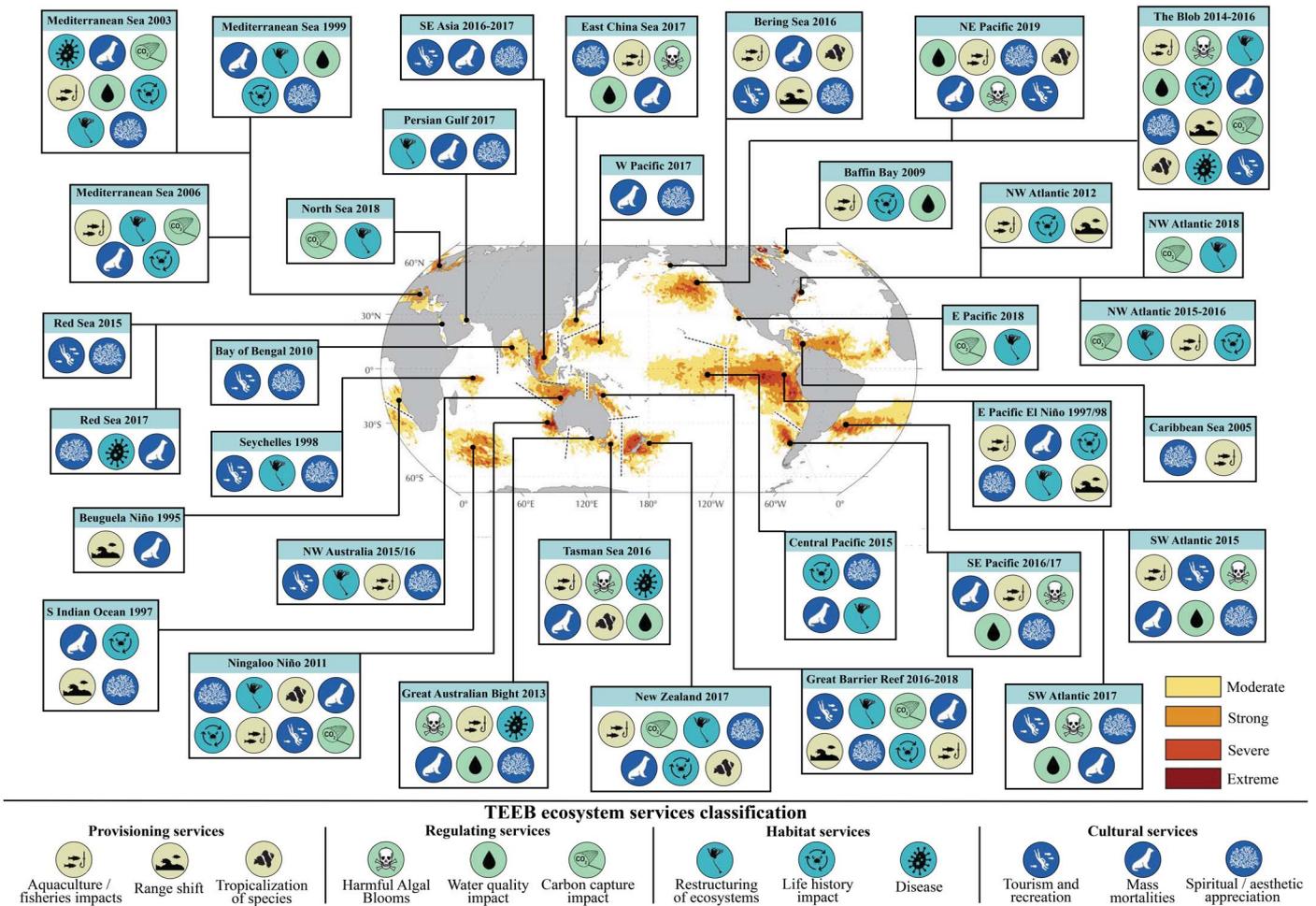


Fig. 1. Socioeconomic impacts of major MHW events since 1995. An event was considered a MHW if sea surface temperatures (SSTs) exceeded the 90th percentile for ≥ 5 days. The intensity scale of each event (moderate to extreme) represents conditions corresponding to the peak date of the event as defined by (41). Intensity categories are identified as sequential multiples of the difference between the 90th percentile and the mean climatology as defined by (42). For each event, biological responses have been linked to ecosystem services based on TEEB ecosystem services classifications [provisioning services, cultural services, regulating services, and habitat services; (40)], which recognize the economic value of nature. For further details on positive and negative biological responses and a full reference list, see Table 2.

fell on either side of the border (52). By contrast, the same MHW shifted longfin squid [*Doryteuthis (Amerigo) pealelli*] distribution poleward, enabling the Maine fishing fleet to develop a profitable new market within the season (35). Recent MHWs during 2016–2017 along the Great Barrier Reef have resulted in bleaching, and ultimately mortality, of coral (53), with the unexpected side effect of an increase in the presence of coral trout (*Plectropomus* and *Variola* spp.) near reefs, resulting in higher catch-per-unit effort by the commercial fishery (37).

Kelp forests, seagrass meadows, and coral reefs are globally distributed coastal marine habitats that deliver regulating services via nutrient cycling and carbon storage (blue carbon) and habitat services by offering biogenic structure for socioeconomically important species and by maintaining high levels of biodiversity (54). However, species that under-

pin these habitats are susceptible to MHWs (30), particularly where populations are situated toward upper thermal limits (i.e., equatorward range edges). Off the coast of Baja California, Mexico, giant kelp (*Macrocystis pyrifera*) populations were negatively affected by multiple MHWs between 2014 and 2018, although the magnitude of responses varied between locations within the species range (12) (Fig. 2C). The loss of giant kelp populations at the equatorward range edges after the MHWs affected several regulating (i.e., carbon capture and transfer, nutrient cycling, and storm protection) and habitat (i.e., ecosystem restructuring and habitat for associated species) services. Similarly, kelp populations on either side of the North Atlantic (the eastern United States and southern Norway) suffered mortalities during MHWs in 2018 (55), with similar implications for ecosystem service provision. Loss of southern bull kelp in New Zealand after the 2017–2018

Tasman Sea MHW (13) have, in addition to the loss of the aforementioned kelp-forest services, also affected traditional knowledge and spirituality because indigenous tribes on the South Island rely on abundant bull kelp to make “poha” bags for food storage. Indeed, ancient stories repeated across generations detail “sacred poha kelp bags” that symbolize the regions wealth, “mana,” and authority over food supplies. MHW-associated loss of bull kelp can thereby reduce tribal spiritual power and their long-term connection with the land and sea (56). Widespread loss of seagrass meadows as a result of MHWs have been reported off eastern and western Australia (21, 57), off the east coast of the United States (58), and in the Mediterranean Sea (44), reducing habitat for economically and culturally important associated species and compromising long-term carbon stocks (59). Reef-building corals are particularly sensitive to extreme heat (60), and between

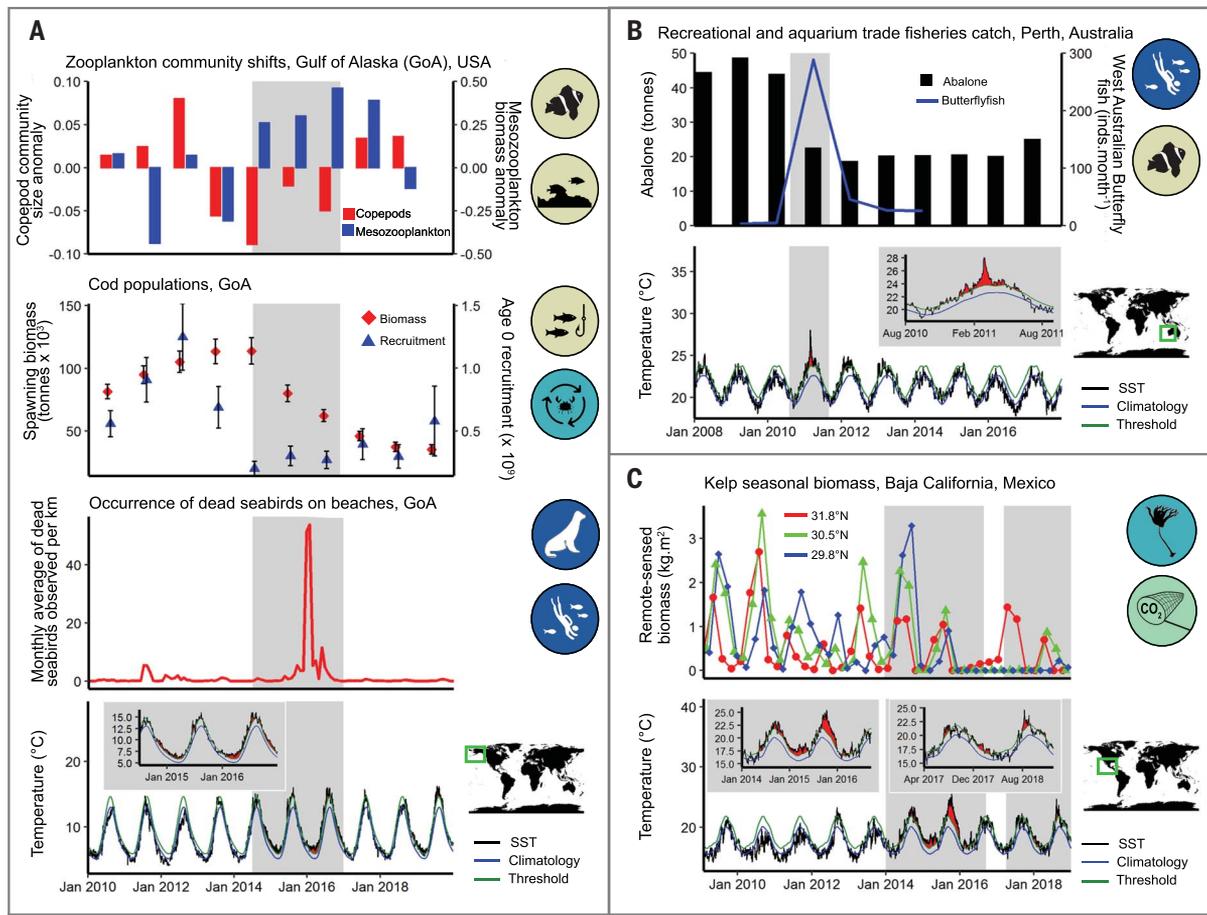


Fig. 2. Impacts of MHWs on ecosystem services. (A) Between 2014 and 2016, MHWs in the northeast Pacific led to a shift in zooplankton community in the Gulf of Alaska from lipid-rich, cold water species to smaller warm-water species [top plot; data extracted from (100)], reducing food supplies for groundfish. Concurrently, the spawning biomass and recruitment in cod (*Gadus macrocephalus*) decreased [second plot; data from (48)], and a mass die-off of fish-eating seabirds occurred due to starvation (third plot; data from <https://COASST.org>). Error bars represent 1 SD. (B) The 2011 MHW off Western Australia led to a short-term increase in butterflyfish [*Chaetodon assarius*, valuable to the aquarium trade; data extracted from (31)] and a long-term decrease in abalone growth, resulting in an ~50% reduction in recreational landings for at least 7 years (data from the Government of Western Australia Department of Fisheries

Annual reports; <https://www.fish.wa.gov.au/About-Us/Publications/Pages/State-of-the-Fisheries-report.aspx>). (C) Giant kelp (*Macrocystis pyrifera*) off the coast of Baja California has been affected by multiple MHWs since 2014, with equatorward populations being most severely affected [data extracted from (12)]. The SST climatology (blue), 90th percentile (green), SST time series (black), and periods of time with identified MHWs (red fill) for each location are shown in the bottom plots. Gray shading and inset climatologies indicate the timing of respective MHWs. An event was considered a MHW if SSTs exceeded the 90th percentile for periods of ≥ 5 days (41). All circled icons are as identified in Fig. 1. Climatology timelines were generated in RStudio (101) using the HeatwaveR package [R Core Team, <https://www.R-project.org>, R package version 0.4.4, <https://CRAN.R-project.org/package=heatwaver>] (102)].

2014 and 2017, MHWs were responsible for a global-scale coral-bleaching event (61) that caused habitat loss and ecosystem reconfiguration in many regions (53). As MHWs increase in frequency (7), the opportunity for recovery between successive events is further reduced, exacerbating ecosystem degradation and affecting regulating, provisioning, habitat, and cultural services, with wide-ranging socioeconomic consequences.

Economic consequences of MHWs

Conservative estimates value global ocean assets at ~US\$24 trillion (3). For example, coral reefs are worth ~US\$35.8 billion per year globally in tourism alone (62), Australian kelp

forests \geq US\$7.8 billion per year in fishing and tourism (63), and U.S. saltwater fishing industries generate >US\$210 billion in sales annually and support 1.7 million jobs (64). Where economic estimates of the impacts of MHWs have been made, reported gains are typically rare and relatively small, whereas losses are common and often extensive (Table 1). In Chile, MHWs associated with the El Niño in 2016 induced harmful algal blooms, which in turn caused the largest fish farm mortality ever recorded globally, resulting in an export loss of US\$800 million (38, 65). Similarly, off the west coast of the United States, a combination of harmful algal blooms and loss of kelp forest habitat (both caused by MHWs) led to closures

of shellfisheries in 2015, costing the economy in excess of US\$185 million in that year alone (14, 66, 67). The commercial tristate Dungeness crab (*Metacarcinus magister*) fishery recorded a loss of ~US\$97.5 million, affecting both tribal and nontribal fisheries (67), and Washington and Californian coastal communities lost a combined ~US\$84 million in tourist spending due to the closure of recreational razor clam (*Siliqua patula*) and abalone (*Haliotis* spp.) fisheries (14, 66). By contrast, the higher temperatures were associated with an increased abundance of warmer water species such as the Pacific bluefin tuna (*Thunnus orientalis*), resulting in an increase in sports fishing charters, and the whale-watching industry saw a surge

in wildlife sightings with associated economic gains from increased demand (28, 68). A surprising outcome of the 2016–2017 coral reef bleaching on the Great Barrier Reef was that, despite expectations, tourism in the area was not affected. Instead, “last chance tourism” motivated visitors, saw an increase in protective attitudes toward the reef, and illustrated the potential for the loss of iconic ecosystems to mobilize the public toward tackling climate change (69).

Loss of biogenic coastal marine habitats are particularly costly because of their role in blue carbon sequestration and as nursery habitat for economically and culturally important species, thereby underpinning provisioning and habitat services. Estimates place the annual per-hectare ecosystem service value of seagrass and coral reefs at ~US\$28,916, and US\$352,249, respectively (70). Scaling this up, the 106,900 hectares of seagrass completely lost in Shark Bay, Western Australia, after the Ningaloo Niño (57) is equivalent to ~US\$3.1 billion in lost ecosystem services per year, with little seagrass recovery observed after 5 years (57). In terms of climate regulation alone, the loss of seagrass habitat in Shark Bay, which was the largest seagrass mortality event in the world recorded to date, resulted in the potential release of between 2 and 9 Tg CO₂ from sedimentary carbon stocks (21).

Interpreting the impacts of MHWs in the context of the SDGs

Contextualizing biological responses within the framework provided by the SDGs further highlights the socioeconomic impacts of MHWs. Clearly, MHWs affect our ability to protect life below water (SDG 14), but there are multiple other SDGs that are interlinked with the impacts of these extreme warming events. Most responses relating to fisheries or aquaculture hinder our ability to achieve zero hunger (SDG 2) or decent work and economic growth (SDG 8), with local effects sometimes having global impacts. For example, the Peruvian anchovy (*Engraulis ringens*) is by far the most caught fish species globally (71). Anchovies are almost exclusively processed into fishmeal and fish oil, which are used extensively in aquaculture and livestock feeds, with Peru providing as much as 50 and 33% of total global fishmeal and fish oil production, respectively (72). An intense MHW associated with the 1997–1998 El Niño and lasting ~8 months caused a temporary range shift in anchovies, ultimately leading to a decrease in landings from 7.5 million tons in 1996 to 1.2 million tons in 1998 (data from FishStatJ, available at www.fao.org/fishery/statistics/software/fishstatj/en), affecting availability of fishmeal and fish oil worldwide and resulting in a reduction in global fishmeal production alone by ~25% (data from seafish, available at www.seafish.org/

document/?id=1b08b6d5-75d9-4179-9094-840195ceee4b).

Coastal and island communities globally rely heavily on human-ocean interactions for income, further highlighting the impacts of MHWs on SDGs 2 and 8. For example, in Southeast Asia, a MHW in 2010 caused a coral-bleaching event costing >US\$49 to 74 million (based on 2010 USD values) in SCUBA-related tourism, (73), and likely impacts on artisanal fisheries. Further, >10% of the Tasmanian Pacific oyster (*Magallana gigas*) industry workforce lost their jobs within months of Tasmania's first outbreak of Pacific oyster mortality syndrome (POMS), which was linked to the Tasman Sea 2015–2016 MHW (74). The Tasmanian POMS outbreak also affected SDG 12 (manage responsible consumption and production) by preventing the movement of spat (juvenile oysters) from Tasmania to seed the Australian oyster industry, affecting production in ~300 farms (75).

Mortality events associated with MHWs in the Mediterranean Sea have been widely recorded, particularly for habitat-forming filter- or suspension-feeding invertebrates such as cnidarians, sponges, and bivalves, thereby reducing water quality and local biodiversity (24). Widespread losses of vast seagrass habitats have been linked with MHWs in Western Australia and the Mediterranean (44, 57), reducing natural carbon sequestration capacity and, thus, the likelihood of meeting SDG13 (climate action). Similarly, kelp forests generally support a significant standing stock of carbon (76) and their loss exemplifies impacts of MHWs on SDG13.

Adaptation, mitigation, and resilience

Human societies are beginning to adapt to the impacts of MHWs to protect ecosystem services and limit socioeconomic consequences. Local industries in several areas have responded to limit economic losses from MHWs in the short-term. For example, during MHWs in North and South America, fisheries efforts were refocused to target transiently abundant species (27, 35, 77). Longer term, fishermen are developing aquaculture programs as a secondary source of income (78) and governments are encouraging diversification to focus on fisheries species expected to increase in abundance with climate warming (79). Approaches also include increasing demand for previously rare or undesirable species through marketing efforts such as renaming and rebranding (80). Although shifting to newly arrived species might ease short-term fishery disruption, it may also slow the population growth of these shifting species such that larger catches are delayed for many years. Different exploitation pathways should be explored with ecosystem models to aid fishery decision makers. Furthermore, aquaculture in-

dustries are responding globally to MHWs by considering climatic extremes when selecting new aquaculture sites, for example, by using remote sensing data (81, 82), by adjusting stock practices to reduce the likelihood of exposure to extreme climatic events (83), by regulating geographic production based on temperature-related lice and disease outbreaks (82), and by selective breeding for disease-resistant aquaculture strains (75).

Early management intervention and shifts in management approaches can also help to limit the impacts of MHWs. For example, after a US\$38 million loss from the lobster price crash caused by the 2012 MHW in the northwest Atlantic, processing capacity was expanded and a proactive management system was applied, which included quotas determined by climatological forecasts (35, 84). Such efforts to increase the resilience of the fishery proved effective during a subsequent MHW in 2016, when the shift in management approach led to a gain of US\$108 million (39, 77). Similarly, in many regions, the combination of stock assessments and flexible harvesting strategies allow management to intervene early to reduce MHW impacts, and a variety of approaches, including quotas, fisheries closures, and restocking, have been implemented to facilitate species recovery and stabilize stocks after MHWs (39, 51). For example, low female spawning biomass identified during stock assessments in 2019 led to the closure of the Gulf of Alaska Pacific cod federal fishery in 2020. Removing fishing pressure for a season enabled the cod population to begin recovering and helped to ensure that adequate prey were available for the endangered western stock of the Steller sea lion (48). However, consistent with increasing MHWs, the United States has seen a significant rise in federal fishery disasters related to climate warming over the past 30 years (85), suggesting that whereas proactive management can prove beneficial, it may be more effectively used in combination with other mitigation strategies.

Natural populations may also be showing resilience to MHWs, with emerging evidence suggesting that moderate climatic events affect population genetics (86). For example, the 2011 Ningaloo Niño caused substantial changes to the genetic structure of kelp forests in the southeast Indian Ocean, with a loss of genetic diversity (87) and an increase in warm-affiliated genotypes (88, 89), consistent with genomic evidence for adaptive changes to increased temperature tolerance (90). Therefore, moderate MHWs are likely to accelerate selection for heat tolerance. Similarly, coral populations exposed to previous MHWs and extreme temperature variation show increased resilience to warming events (91).

It remains uncertain whether marine protected areas (MPAs) can buffer the impacts of

extreme climatic events in a warming ocean (92). Evidence certainly indicates that warming events occurring inside MPAs have comparable, or sometimes even greater, impacts as events in nonprotected areas [(92) and references therein]. This is partly because, historically, climate change and extreme climatic events have not been considered during the design and implementation phases of MPAs (92). MPAs are more likely to buffer the impacts of MHWs if climate change responses are considered in their design. Planning larger networks of no-take MPAs may provide some protection for species tracking their thermal niches compared with smaller, isolated MPAs. Likewise, targeting locations with steep depth (and therefore often temperature) gradients or strong upwelling may offer climatic refugia. Furthermore, examining relevant scales of spatial variability in sea temperature may identify climatic refugia for the foundation species that underpin biodiversity (92, 93). Certainly, resilience to and recovery from MHW events vary spatially, and what drives this is not yet well understood but may be related to local environmental conditions (58, 94). For example, giant kelp populations off Bahia Tortugas, Mexico, close to the equatorward range edge of the species, have shown greater resilience to and more rapid recovery from multiple MHWs compared with populations both north and south of this location (94, 95), suggesting that this area could serve as an important refuge for giant kelp and its associated communities (and therefore should be prioritized in designs of future MPAs). Regardless, MPAs are likely to be more effective at mitigating the impacts of MHWs if combined with other management tools such as restoration or stock enhancement (96).

Future directions

Efforts to improve mechanistic understanding of the physical drivers of MHWs will undoubtedly lead to better climatological forecasting and predictions (39), which in turn will provide opportunities to implement mitigation measures. It is clear that MHWs induce both direct (e.g., coral reef bleaching reduces tourism) and indirect (e.g., loss of kelp forests results in lack of nursery grounds for fisheries species, which leads to reduced landings) impacts on ecosystem services, which can last from weeks to decades or more. The realization that heritable genotypic variation likely underpins responses to MHWs opens the possibility to harness “future-proof” populations through selective breeding and assisted evolution of resilience (97, 98). However, current knowledge of (direct or indirect) biological impacts of MHWs is lacking for many species and populations, which limits our capacity to safeguard marine resources against future extreme climatic events (99). Moreover, understanding of how MHWs

interact with other stressors, such as oxygen, nutrient, and light availability, to affect marine organisms, populations, and ecosystems remains limited. MHWs have intensified over the past century largely due to anthropogenic climate change and will continue to do so as warming intensifies further, with wide-ranging consequences for marine ecosystems globally (11). There is a pressing need to develop a toolbox of adaptation and mitigation measures, including improved climatological forecasting, proactive resource management, and enhanced resilience, in response to increasing MHW impacts in the coming decades.

REFERENCES AND NOTES

- G. T. Pecl *et al.*, Biodiversity redistribution under climate change: Impacts on ecosystems and human well-being. *Science* **355**, eaai9214 (2017). doi: [10.1126/science.aai9214](https://doi.org/10.1126/science.aai9214); pmid: [28360268](https://pubmed.ncbi.nlm.nih.gov/28360268/)
- C. Nolan *et al.*, Past and future global transformation of terrestrial ecosystems under climate change. *Science* **361**, 920–923 (2018). doi: [10.1126/science.aan5360](https://doi.org/10.1126/science.aan5360); pmid: [30166491](https://pubmed.ncbi.nlm.nih.gov/30166491/)
- Intergovernmental Panel on Climate Change. “Climate change 2021: The physical science basis. Contribution of Working Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change.” V. Masson-Delmotte *et al.*, Eds. (Cambridge Univ. Press, 2021); https://www.ipcc.ch/report/ar6/wg1/downloads/report/IPCC_AR6_WGI_Full_Report_smaller.pdf.
- L. Wu *et al.*, Enhanced warming over the global subtropical western boundary currents. *Nat. Clim. Chang.* **2**, 161 (2012). doi: [10.1038/nclimate1353](https://doi.org/10.1038/nclimate1353)
- E. S. Poloczanska *et al.*, Global imprint of climate change on marine life. *Nat. Clim. Chang.* **3**, 919–925 (2013). doi: [10.1038/nclimate1958](https://doi.org/10.1038/nclimate1958)
- A. Vergés *et al.*, The tropicalization of temperate marine ecosystems: Climate-mediated changes in herbivory and community phase shifts. *Proc. Biol. Sci.* **281**, 201400846 (2014). doi: [10.1098/rspb.2014.0846](https://doi.org/10.1098/rspb.2014.0846); pmid: [25009065](https://pubmed.ncbi.nlm.nih.gov/25009065/)
- E. C. J. Oliver *et al.*, Longer and more frequent marine heatwaves over the past century. *Nat. Commun.* **9**, 1324 (2018). doi: [10.1038/s41467-018-03732-9](https://doi.org/10.1038/s41467-018-03732-9); pmid: [29636482](https://pubmed.ncbi.nlm.nih.gov/29636482/)
- A. Sen Gupta *et al.*, Drivers and impacts of the most extreme marine heatwaves events. *Sci. Rep.* **10**, 19359 (2020). doi: [10.1038/s41598-020-75445-3](https://doi.org/10.1038/s41598-020-75445-3); pmid: [33168858](https://pubmed.ncbi.nlm.nih.gov/33168858/)
- N. J. Holbrook *et al.*, A global assessment of marine heatwaves and their drivers. *Nat. Commun.* **10**, 2624 (2019). doi: [10.1038/s41467-019-10206-z](https://doi.org/10.1038/s41467-019-10206-z); pmid: [31201309](https://pubmed.ncbi.nlm.nih.gov/31201309/)
- E. C. J. Oliver *et al.*, Projected marine heatwaves in the 21st century and the potential for ecological impact. *Front. Mar. Sci.* **6**, 734 (2019). doi: [10.3389/fmars.2019.00734](https://doi.org/10.3389/fmars.2019.00734)
- M. Collins, M. Sutherland, L. Bouwer, “Extremes, abrupt changes and managing risk,” in: *IPCC Special Report on the Ocean and Cryosphere in a Changing Climate*, H. O. Portner, D. C. Roberts, V. Masson-Delmotte, P. Zhai, M. Tignor, E. Poloczanska, K. Minterbeck, A. Alegria, M. Nicolai, A. Okem, J. Petzold, B. Rama, N.M. Weyer, Eds. (IPCC, 2019), pp. 589–656.; https://www.ipcc.ch/site/assets/uploads/sites/3/2019/11/SROCC_Ch06_FINAL.pdf.
- N. Arafeh-Dalmau *et al.*, Extreme marine heatwaves alter kelp forest community near its equatorward distribution limit. *Front. Mar. Sci.* **6**, 499 (2019). doi: [10.3389/fmars.2019.00499](https://doi.org/10.3389/fmars.2019.00499)
- M. S. Thomsen *et al.*, Local extinction of bull kelp (*Durvillaea* spp.) due to a marine heatwave. *Front. Mar. Sci.* **6**, 84 (2019). doi: [10.3389/fmars.2019.00084](https://doi.org/10.3389/fmars.2019.00084)
- L. Rogers-Bennett, C. A. Cutton, Marine heat wave and multiple stressors tip bull kelp forest to sea urchin barrens. *Sci. Rep.* **9**, 15050 (2019). doi: [10.1038/s41598-019-51114-y](https://doi.org/10.1038/s41598-019-51114-y); pmid: [31636286](https://pubmed.ncbi.nlm.nih.gov/31636286/)
- T. Wernberg *et al.*, Climate-driven regime shift of a temperate marine ecosystem. *Science* **353**, 169–172 (2016). doi: [10.1126/science.aad8745](https://doi.org/10.1126/science.aad8745); pmid: [27387951](https://pubmed.ncbi.nlm.nih.gov/27387951/)
- S. D. Donner, G. J. M. Rickbeil, S. F. Heron, A new, high-resolution global mass coral bleaching database. *PLOS ONE* **12**, e0175490 (2017). doi: [10.1371/journal.pone.0175490](https://doi.org/10.1371/journal.pone.0175490); pmid: [28445534](https://pubmed.ncbi.nlm.nih.gov/28445534/)
- A. P. Krishnan *et al.*, Elevated sea surface temperature during May 2010 induces mass bleaching of corals in the Andaman. *Curr. Sci.* **100**, 111–117 (2011); www.curentscience.ac.in/Volumes/100/01/0111.pdf.
- J. A. Y. Moore *et al.*, Unprecedented mass bleaching and loss of coral across 12° of latitude in Western Australia in 2010–11. *PLOS ONE* **7**, e51807 (2012). doi: [10.1371/journal.pone.0051807](https://doi.org/10.1371/journal.pone.0051807); pmid: [23284773](https://pubmed.ncbi.nlm.nih.gov/23284773/)
- C. Cerrano *et al.*, A catastrophic mass-mortality episode of gorgonians and other organisms in the Ligurian Sea (North-western Mediterranean), summer 1999. *Ecol. Lett.* **3**, 284–293 (2000). doi: [10.1046/j.1461-0248.2000.00152.x](https://doi.org/10.1046/j.1461-0248.2000.00152.x)
- M. W. Fraser *et al.*, Extreme climate events lower resilience of foundation seagrass at edge of biogeographical range. *J. Ecol.* **102**, 1528–1536 (2014). doi: [10.1111/1365-1230.12300](https://doi.org/10.1111/1365-1230.12300)
- A. Arias-Ortiz *et al.*, A marine heatwave drives massive losses from the world’s largest seagrass carbon stocks. *Nat. Clim. Chang.* **8**, 338–344 (2018). doi: [10.1038/s41558-018-0096-y](https://doi.org/10.1038/s41558-018-0096-y)
- T. Gammelsrød, C. H. Bartholomae, D. C. Boyer, V. L. L. Filipe, M. J. O’Toole, Intrusion of warm surface water along the Angolan-Namibian coast in February–March 1995: The 1995 Benguela Niño. *S. Afr. J. Mar. Sci.* **19**, 41–56 (1998). doi: [10.2989/025776198784126719](https://doi.org/10.2989/025776198784126719)
- T. Jones *et al.*, Massive mortality of a planktivorous seabird in response to a marine heatwave. *Geophys. Res. Lett.* **45**, 3193–3202 (2018). doi: [10.1002/2017GL076164](https://doi.org/10.1002/2017GL076164)
- J. Garrabuou *et al.*, Collaborative database to track mass mortality events in the Mediterranean Sea. *Front. Mar. Sci.* **6**, 707 (2019). doi: [10.3389/fmars.2019.00707](https://doi.org/10.3389/fmars.2019.00707)
- J. F. Platt *et al.*, Extreme mortality and reproductive failure of common murrelets resulting from the northeast Pacific marine heatwave of 2014–2016. *PLOS ONE* **15**, e0226087 (2020). doi: [10.1371/journal.pone.0226087](https://doi.org/10.1371/journal.pone.0226087); pmid: [31940310](https://pubmed.ncbi.nlm.nih.gov/31940310/)
- S. P. Salazar, R. H. Bustamante, Effects of the 1997–1998 El Niño on population size and diet of the Galápagos sea lion (*Zalophus wollebaeki*). *Not. Galapagos* **62**, 40–45 (2003).
- M. Niquen, M. Bouchon, Impact of El Niño events on pelagic fisheries in Peruvian waters. *Deep. Res. Part II Top. Stud. Oceanogr.* **51**, 563–574 (2004). doi: [10.1016/j.jdsr.2004.03.001](https://doi.org/10.1016/j.jdsr.2004.03.001)
- L. M. Cavole *et al.*, Biological impacts of the 2013–2015 warm-water anomaly in the northeast Pacific: Winners, losers, and the future. *Oceanography (Wash. D.C.)* **29**, 273–285 (2016). doi: [10.5670/oceanog.2016.32](https://doi.org/10.5670/oceanog.2016.32)
- S. J. Barbeaux, K. Holsman, S. Zador, Marine heatwave stress test of ecosystem-based fisheries management in the Gulf of Alaska Pacific cod fishery. *Front. Mar. Sci.* **7**, 703 (2020). doi: [10.3389/fmars.2020.00703](https://doi.org/10.3389/fmars.2020.00703)
- D. A. Smale *et al.*, Marine heatwaves threaten global biodiversity and the provision of ecosystem services. *Nat. Clim. Chang.* **9**, 306–312 (2019). doi: [10.1038/s41558-019-0412-1](https://doi.org/10.1038/s41558-019-0412-1)
- R. C. J. Lenanton, C. E. Dowling, K. A. Smith, D. V. Fairclough, G. Jackson, Potential influence of a marine heatwave on range extensions of tropical fishes in the eastern Indian Ocean—Invaluable contributions from amateur observers. *Reg. Stud. Mar. Sci.* **13**, 19–31 (2017). doi: [10.1016/j.jrsm.2017.03.005](https://doi.org/10.1016/j.jrsm.2017.03.005)
- C. Bouchard *et al.*, Climate warming enhances polar cod recruitment, at least transiently. *Prog. Oceanogr.* **156**, 121–129 (2017). doi: [10.1016/j.pocean.2017.06.008](https://doi.org/10.1016/j.pocean.2017.06.008)
- M. LeBlanc *et al.*, Pelagic production and the recruitment of juvenile polar cod *Boreogadus saida* in Canadian Arctic seas. *Polar Biol.* **43**, 1043–1054 (2020). doi: [10.1007/s00300-019-02565-6](https://doi.org/10.1007/s00300-019-02565-6)
- J. M. Sunday *et al.*, Species traits and climate velocity explain geographic range shifts in an ocean-warming hotspot. *Ecol. Lett.* **18**, 944–953 (2015). doi: [10.1111/ele.12474](https://doi.org/10.1111/ele.12474); pmid: [26189556](https://pubmed.ncbi.nlm.nih.gov/26189556/)
- K. E. Mills *et al.*, Fisheries management in a changing climate. *Oceanography (Wash. D.C.)* **26**, 191–195 (2013). doi: [10.5670/oceanog.2013.27](https://doi.org/10.5670/oceanog.2013.27)
- N. Caputi *et al.*, Management adaptation of invertebrate fisheries to an extreme marine heat wave event at a global warming hot spot. *Ecol. Evol.* **6**, 3583–3593 (2016). doi: [10.1002/ece3.2137](https://doi.org/10.1002/ece3.2137); pmid: [28725352](https://pubmed.ncbi.nlm.nih.gov/28725352/)
- C. J. Brown, C. Mellin, G. J. Edgar, M. D. Campbell, R. D. Stuart-Smith, Direct and indirect effects of heatwaves on a coral reef fishery. *Glob. Chang. Biol.* **27**, 1214–1225 (2021). doi: [10.1111/gcb.15472](https://doi.org/10.1111/gcb.15472); pmid: [33340216](https://pubmed.ncbi.nlm.nih.gov/33340216/)
- V. L. Trainer *et al.*, Pelagic harmful algal blooms and climate change: Lessons from nature’s experiments with extremes.

- Harmful Algae* **91**, 101591 (2020). doi: [10.1016/j.hal.2019.03.009](https://doi.org/10.1016/j.hal.2019.03.009); pmid: [32057339](https://pubmed.ncbi.nlm.nih.gov/32057339/)
39. N. J. Holbrook *et al.*, Keeping pace with marine heatwaves. *Nat. Rev. Earth Environ.* **1**, 482–493 (2020). doi: [10.1038/s43017-020-0068-4](https://doi.org/10.1038/s43017-020-0068-4)
 40. R. De Groot, B. Fisher, M. Christie, “Integrating the ecological and economic dimensions in biodiversity and ecosystem service valuation,” in *The Economics of Ecosystems and Biodiversity: The Ecological and Economic Foundations*, P. Kumar, Ed. (Cambridge Univ. Press, 2010); pp. 9–40.
 41. A. J. Hobday *et al.*, A hierarchical approach to defining marine heatwaves. *Prog. Oceanogr.* **141**, 227–238 (2016). doi: [10.1016/j.pcean.2015.12.014](https://doi.org/10.1016/j.pcean.2015.12.014)
 42. A. J. Hobday *et al.*, Categorizing and naming marine heatwaves. *Oceanography (Wash. D.C.)* **31**, 162–173 (2018). doi: [10.5670/oceanog.2018.205](https://doi.org/10.5670/oceanog.2018.205)
 43. D. B. Enfield, Evolution and historical perspective of the 1997–1998 El Niño–Southern Oscillation event. *Bull. Mar. Sci.* **69**, 7–25 (2001).
 44. N. Marbà, C. M. Duarte, Mediterranean warming triggers seagrass (*Posidonia oceanica*) shoot mortality. *Glob. Change Biol.* **16**, 2366–2375 (2010). doi: [10.1111/j.1365-2486.2009.02130.x](https://doi.org/10.1111/j.1365-2486.2009.02130.x)
 45. L. C. Rodrigues *et al.*, Sensitivity of Mediterranean Bivalve Mollusc Aquaculture to Climate Change, Ocean Acidification, and Other Environmental Pressures: Findings from a Producer Survey. *J. Shellfish Res.* **34**, 1161–1176 (2015). doi: [10.2983/035.034.0341](https://doi.org/10.2983/035.034.0341)
 46. E. Di Lorenzo, N. Mantua, Multi-year persistence of the 2014/15 North Pacific marine heatwave. *Nat. Clim. Chang.* **6**, 1042–1047 (2016). doi: [10.1038/nclimate3082](https://doi.org/10.1038/nclimate3082)
 47. L. A. Rogers, M. T. Wilson, J. T. Duffy-Anderson, D. G. Kimmel, J. F. Lamb, Pollock and “the Blob”: Impacts of a marine heatwave on walleye pollock early life stages. *Fish. Oceanogr.* **30**, 142–158 (2020). doi: [10.1111/fog.12508](https://doi.org/10.1111/fog.12508)
 48. S. J. Barbeaux, T. A. mar, W. A. Palsson, “Assessment of the Pacific cod stock in the Gulf of Alaska,” in: *North Pacific Fishery Management Council Gulf of Alaska Stock Assessment and Fishery Evaluation Report* (NPFMC, 2020); https://www.researchgate.net/publication/311733967_Assessment_of_the_Pacific_cod_stock_in_the_Gulf_of_Alaska
 49. R. M. McCabe *et al.*, An unprecedented coastwide toxic algal bloom linked to anomalous ocean conditions. *Geophys. Res. Lett.* **43**, 10366–10376 (2016). doi: [10.1002/2016GL070023](https://doi.org/10.1002/2016GL070023); pmid: [27917011](https://pubmed.ncbi.nlm.nih.gov/27917011/)
 50. T. Wernberg *et al.*, An extreme climatic event alters marine ecosystem structure in a global biodiversity hotspot. *Nat. Clim. Chang.* **3**, 78–82 (2013). doi: [10.1038/nclimate1627](https://doi.org/10.1038/nclimate1627)
 51. N. Caputi *et al.*, Factors affecting the recovery of invertebrate stocks from the 2011 Western Australian extreme marine heatwave. *Front. Mar. Sci.* **6**, 484 (2019). doi: [10.3389/fmars.2019.00484](https://doi.org/10.3389/fmars.2019.00484)
 52. K. Sgueglia, “Too much Maine lobster prompts protests and court order in Canada,” *CNN*, 10 August 2012; <https://edition.cnn.com/2012/08/09/us/canada-lobster-injection/index.html>
 53. T. P. Hughes *et al.*, Global warming transforms coral reef assemblages. *Nature* **556**, 492–496 (2018). doi: [10.1038/s41586-018-0041-2](https://doi.org/10.1038/s41586-018-0041-2); pmid: [29670282](https://pubmed.ncbi.nlm.nih.gov/29670282/)
 54. A. Himes-Cornell, L. Pendleton, P. Atiyah, Valuing ecosystem services from blue forests: A systematic review of the valuation of salt marshes, sea grass beds and mangrove forests. *Ecosyst. Serv.* **30**, 36–48 (2018). doi: [10.1016/j.ecoser.2018.01.006](https://doi.org/10.1016/j.ecoser.2018.01.006)
 55. K. Filbee-Dexter *et al.*, Marine heatwaves and the collapse of marginal North Atlantic kelp forests. *Sci. Rep.* **10**, 13388 (2020). doi: [10.1038/s41598-020-70273-x](https://doi.org/10.1038/s41598-020-70273-x); pmid: [32770015](https://pubmed.ncbi.nlm.nih.gov/32770015/)
 56. Te Rūnanga o Kaikōura, “Te Poha o Tohu Raumati (Environmental Management Plan)” (Te Rūnanga o Kaikōura, 2007); <https://ngaitahu.iwi.nz/wp-content/uploads/2013/08/Te-Runanga-o-Kaikoura-Environmental-Management-Plan.pdf>
 57. S. Strydom *et al.*, Too hot to handle: Unprecedented seagrass death driven by marine heatwave in a World Heritage Area. *Glob. Change Biol.* **26**, 3525–3538 (2020). doi: [10.1111/gcb.15065](https://doi.org/10.1111/gcb.15065); pmid: [32129909](https://pubmed.ncbi.nlm.nih.gov/32129909/)
 58. L. R. Aoki *et al.*, Seagrass recovery following marine heat wave influences sediment carbon stocks. *Front. Mar. Sci.* **7**, 576784 (2021). doi: [10.3389/fmars.2020.576784](https://doi.org/10.3389/fmars.2020.576784)
 59. R. J. Orth *et al.*, A global crisis for seagrass ecosystems. *Bioscience* **56**, 987–996 (2006). doi: [10.1641/0006-3568\(2006\)56\[987:AGCFSE\]2.0.CO;2](https://doi.org/10.1641/0006-3568(2006)56[987:AGCFSE]2.0.CO;2)
 60. T. P. Hughes *et al.*, Coral reefs in the Anthropocene. *Nature* **546**, 82–90 (2017). doi: [10.1038/nature22901](https://doi.org/10.1038/nature22901); pmid: [28569801](https://pubmed.ncbi.nlm.nih.gov/28569801/)
 61. C. M. Eakin, H. P. A. Sweatman, R. E. Brainard, The 2014–2017 global-scale coral bleaching event: Insights and impacts. *Coral Reefs* **38**, 539–545 (2019). doi: [10.1007/s00338-019-01844-2](https://doi.org/10.1007/s00338-019-01844-2)
 62. M. Spalding *et al.*, Mapping the global value and distribution of coral reef tourism. *Mar. Policy* **82**, 104–113 (2017). doi: [10.1016/j.marpol.2017.05.014](https://doi.org/10.1016/j.marpol.2017.05.014)
 63. S. Bennett *et al.*, The “Great Southern Reef”: Social, ecological and economic value of Australia’s neglected kelp forests. *Mar. Freshw. Res.* **67**, 47–56 (2016). doi: [10.1071/MF15232](https://doi.org/10.1071/MF15232)
 64. National Marine Fisheries Service, Fisheries Economics of the United States, 2016 (NOAA, 2018); <https://www.fisheries.noaa.gov/resource/document/fisheries-economics-united-states-report-2016>
 65. A. Clement *et al.*, Exceptional summer conditions and HABs of *Pseudoacanthionella* in Southern Chile create record impacts on salmon farms. *Harmful Algae News* **53**, 1–3 (2016); www.e-pages.dk/ku/1217/
 66. B. E. Ferriss, D. J. Marcinek, D. Ayres, J. Borchert, K. A. Lefebvre, Acute and chronic dietary exposure to domoic acid in recreational harvesters: A survey of shellfish consumption behavior. *Environ. Int.* **101**, 70–79 (2017). doi: [10.1016/j.envint.2017.01.006](https://doi.org/10.1016/j.envint.2017.01.006); pmid: [28109640](https://pubmed.ncbi.nlm.nih.gov/28109640/)
 67. S. K. Moore *et al.*, An index of fisheries closures due to harmful algal blooms and a framework for identifying vulnerable fishing communities on the U.S. West Coast. *Mar. Policy* **110**, 103543 (2019). doi: [10.1016/j.marpol.2019.103543](https://doi.org/10.1016/j.marpol.2019.103543)
 68. J. A. Santora *et al.*, Habitat compression and ecosystem shifts as potential links between marine heatwave and record whale entanglements. *Nat. Commun.* **11**, 536 (2020). doi: [10.1038/s41467-019-14215-w](https://doi.org/10.1038/s41467-019-14215-w); pmid: [31988285](https://pubmed.ncbi.nlm.nih.gov/31988285/)
 69. M. I. Curnock *et al.*, Shifts in tourists’ sentiments and climate risk perceptions following mass coral bleaching of the Great Barrier Reef. *Nat. Clim. Chang.* **9**, 535–541 (2019). doi: [10.1038/s41558-019-0504-y](https://doi.org/10.1038/s41558-019-0504-y)
 70. R. Costanza *et al.*, Changes in the global value of ecosystem services. *Glob. Environ. Change* **26**, 152–158 (2014). doi: [10.1016/j.gloenvcha.2014.04.002](https://doi.org/10.1016/j.gloenvcha.2014.04.002)
 71. Food and Agriculture Organization of the United Nations, “FACT SHEET: The international fish trade and world fisheries” (FAO, 2010); http://www.fao.org/fileadmin/user_upload/newsroom/docs/fact_sheet_fish_trade_en.pdf
 72. P. Frón *et al.*, Harvesting for food versus feed: A review of Peruvian fisheries in a global context. *Rev. Fish Biol. Fish.* **24**, 381–398 (2014). doi: [10.1007/s11160-013-9336-4](https://doi.org/10.1007/s11160-013-9336-4)
 73. A. Doshi *et al.*, “Loss of economic value from coral bleaching in SE Asia,” in 12th International Coral Reef Symposium, Cairns, Australia, 9–13 July 2012; https://www.academia.edu/12494037/Loss_of_economic_value_from_coral_bleaching_in_SE_Asia
 74. E. C. J. Oliver *et al.*, The unprecedented 2015/16 Tasman Sea marine heatwave. *Nat. Commun.* **8**, 16101 (2017). doi: [10.1038/ncomms16101](https://doi.org/10.1038/ncomms16101); pmid: [28706247](https://pubmed.ncbi.nlm.nih.gov/28706247/)
 75. I. Catzone, “National impact from Tasmanian POMS outbreak” (Fisheries Research and Development Corporation, 2016); <https://www.frdc.com.au/fish-vol-24-2/national-impact-tasmanian-poms-outbreak>
 76. K. Filbee-Dexter, T. Wernberg, Substantial blue carbon in overlooked Australian kelp forests. *Sci. Rep.* **10**, 12341 (2020). doi: [10.1038/s41598-020-69258-7](https://doi.org/10.1038/s41598-020-69258-7)
 77. A. Pershing, K. Mills, A. Dayton, B. Franklin, B. Kennedy, Evidence for adaptation from the 2016 marine heatwave in the Northwest Atlantic Ocean. *Oceanography (Wash. D.C.)* **31**, 152–161 (2018). doi: [10.5670/oceanog.2018.213](https://doi.org/10.5670/oceanog.2018.213)
 78. F. Bever, “Maine fishermen prepare for losses and gains in a climate changed ocean,” *WBUR*, 23 September 2019; <https://www.wbur.com/news/2019/09/23/maine-fishermen-prepare-for-losses-and-gains-in-a-climate-changed-ocean>
 79. Oxford Business Group, “Peru Agriculture: Diversification efforts are up as instability over anchovy supply continues” (Oxford Business Group, 2016); <https://oxfordbusinessgroup.com/analysis/choppy-waters-diversification-efforts-are-instability-over-anchovy-supply-continues>
 80. T. Yusuke, N. Masato, “Marine heatwaves driving change in Hokkaido’s fishing industry,” *NHK World-Japan*, 10 December 2020; <https://www3.nhk.or.jp/nhkworld/en/news/backstories/1404/>
 81. Y. Liu, S. I. Saitoh, H. Igarashi, T. Hirawake, The regional impacts of climate change on coastal environments and the aquaculture of Japanese scallops in northeast Asia: Case studies from Dalian, China, and Funka Bay, Japan. *Int. J.*
 - Remote Sens. **35**, 4422–4440 (2014). doi: [10.1080/01431161.2014.916435](https://doi.org/10.1080/01431161.2014.916435)
 82. H. T. Sanderson, J. Olsen, G. K. Hovelsrud, A. Gjertsen, “Climate change and Norwegian arctic aquaculture: Perception, relevance, and adaptation,” in *Arctic Yearbook 2020*, L. Heininen, H. Exner-Pireot, J. Barnes, Eds. (2020); pp. 316–327.
 83. L. Lebel *et al.*, Innovation, practice, and adaptation to climate in the aquaculture sector. *Rev. Fish. Sci. Aquacult.* (2020); doi: [10.1080/23308249.2020.1869695](https://doi.org/10.1080/23308249.2020.1869695)
 84. A. Le Bris *et al.*, Climate vulnerability and resilience in the most valuable North American fishery. *Proc. Natl. Acad. Sci. U.S.A.* **115**, 1831–1836 (2018). doi: [10.1073/pnas.1711221115](https://doi.org/10.1073/pnas.1711221115); pmid: [29358389](https://pubmed.ncbi.nlm.nih.gov/29358389/)
 85. L. Bellquist, V. Saccomanno, B. X. Semmens, M. Gleason, J. Wilson, The rise in climate change-induced federal fishery disasters in the United States. *PeerJ* **9**, e11186 (2021). doi: [10.7717/peerj.11186](https://doi.org/10.7717/peerj.11186); pmid: [33981495](https://pubmed.ncbi.nlm.nih.gov/33981495/)
 86. M. A. Coleman, T. Wernberg, The silver lining of extreme events. *Trends Ecol. Evol.* **35**, 1065–1067 (2020). doi: [10.1016/j.tree.2020.08.013](https://doi.org/10.1016/j.tree.2020.08.013); pmid: [32958366](https://pubmed.ncbi.nlm.nih.gov/32958366/)
 87. T. Wernberg *et al.*, Genetic diversity and kelp forest vulnerability to climatic stress. *Sci. Rep.* **8**, 1851 (2018). doi: [10.1038/s41598-018-20009-9](https://doi.org/10.1038/s41598-018-20009-9); pmid: [293282916](https://pubmed.ncbi.nlm.nih.gov/293282916/)
 88. M. A. Coleman, A. J. P. Minne, S. Vranken, T. Wernberg, Genetic tropicalisation following a marine heatwave. *Sci. Rep.* **10**, 12726 (2020). doi: [10.1038/s41598-020-69665-w](https://doi.org/10.1038/s41598-020-69665-w); pmid: [32728196](https://pubmed.ncbi.nlm.nih.gov/32728196/)
 89. C. F. D. Gurgel, O. Camacho, A. J. P. Minne, T. Wernberg, M. A. Coleman, Marine heatwave drives cryptic loss of genetic diversity in underwater forests. *Curr. Biol.* **30**, 1199–1206.e2 (2020). doi: [10.1016/j.cub.2020.01.051](https://doi.org/10.1016/j.cub.2020.01.051); pmid: [32109397](https://pubmed.ncbi.nlm.nih.gov/32109397/)
 90. S. Vranken *et al.*, Genotype–Environment mismatch of kelp forests under climate change. *Mol. Ecol.* **30**, 3730–3746 (2021). doi: [10.1111/mec.15993](https://doi.org/10.1111/mec.15993); pmid: [34018645](https://pubmed.ncbi.nlm.nih.gov/34018645/)
 91. J. R. Guest *et al.*, Contrasting patterns of coral bleaching susceptibility in 2010 suggest an adaptive response to thermal stress. *PLOS ONE* **7**, e33353 (2012). doi: [10.1371/journal.pone.0033353](https://doi.org/10.1371/journal.pone.0033353); pmid: [22428027](https://pubmed.ncbi.nlm.nih.gov/22428027/)
 92. A. E. Bates *et al.*, Climate resilience in marine protected areas and the “Protection Paradox”. *Biol. Conserv.* **236**, 305–314 (2019). doi: [10.1016/j.biocon.2019.05.005](https://doi.org/10.1016/j.biocon.2019.05.005)
 93. J. Verdura *et al.*, Local-scale climatic refugia offer sanctuary for a habitat-forming species during a marine heatwave. *J. Ecol.* **109**, 1758–1773 (2021). doi: [10.1111/1365-2745.13599](https://doi.org/10.1111/1365-2745.13599)
 94. K. C. Cavanaugh, D. C. Reed, T. W. Bell, M. C. N. Castorani, R. Beas-Luna, Spatial variability in the resistance and resilience of giant kelp in southern and Baja California to a multiyear heatwave. *Front. Mar. Sci.* **6**, 413 (2019). doi: [10.3389/fmars.2019.00413](https://doi.org/10.3389/fmars.2019.00413)
 95. M. S. Edwards, J. A. Estes, Catastrophe, recovery and range limitation in NE Pacific kelp forests: A large-scale perspective. *Mar. Ecol. Prog. Ser.* **320**, 79–87 (2006). doi: [10.3354/meps320079](https://doi.org/10.3354/meps320079)
 96. R. M. Freedman, J. A. Brown, C. Caldwell, J. E. Caselle, Marine protected areas do not prevent marine heatwave-induced fish community structure changes in a temperate transition zone. *Sci. Rep.* **10**, 21081 (2020). doi: [10.1038/s41598-020-77885-3](https://doi.org/10.1038/s41598-020-77885-3); pmid: [33273514](https://pubmed.ncbi.nlm.nih.gov/33273514/)
 97. M. J. H. van Oppen, J. K. Oliver, H. M. Putnam, R. D. Gates, Building coral reef resilience through assisted evolution. *Proc. Natl. Acad. Sci. U.S.A.* **112**, 2307–2313 (2015). doi: [10.1073/pnas.1422301112](https://doi.org/10.1073/pnas.1422301112); pmid: [25646461](https://pubmed.ncbi.nlm.nih.gov/25646461/)
 98. M. A. Coleman *et al.*, Restore or redefine: Future trajectories for restoration. *Front. Mar. Sci.* **7**, 237 (2020). doi: [10.3389/fmars.2020.00237](https://doi.org/10.3389/fmars.2020.00237)
 99. J. A. Benthuisen, E. C. J. Oliver, K. Chen, T. Wernberg, Editorial: Advances in understanding marine heatwaves and their impacts. *Front. Mar. Sci.* **7**, 147 (2020). doi: [10.3389/fmars.2020.00147](https://doi.org/10.3389/fmars.2020.00147)
 100. S. Zador, E. Yasumiishi, G. A. Whitehouse, eds., “Ecosystem Status Report 2019 Gulf of Alaska” (National Oceanic and Atmospheric Administration, 2019); <https://www.fisheries.noaa.gov/resource/data/ecosystem-status-report-2019-gulf-alaska>
 101. RStudio Team, “RStudio: Integrated development for R” (2020); <https://www.rstudio.com/>
 102. R. W. Schlegel, A. J. Smit, heatwaveR: A central algorithm for the detection of heatwaves and cold-spells. *J. Open Source Softw.* **3**, 821 (2018). doi: [10.21105/joss.00821](https://doi.org/10.21105/joss.00821)
 103. M. C. Fisher, S. K. Moore, S. L. Jardine, J. R. Watson, J. F. Samhuri, Climate shock effects and mediation in fisheries. *Proc. Natl. Acad. Sci. U.S.A.* **118**, e2014379117 (2021). doi: [10.1073/pnas.2014379117](https://doi.org/10.1073/pnas.2014379117); pmid: [3397723](https://pubmed.ncbi.nlm.nih.gov/3397723/)

104. Australian Government Department of Aquaculture, Water and the Environment, "Australian fisheries and aquaculture statistics 2018" (ABARES, 2020); https://daff.ent.sirsidynix.net.au/client/en_AU/search/asset/1030241/0.
105. A. Chandrapavan, N. Caputi, M. I. Kangas, The decline and recovery of a crab population from an extreme marine heatwave and a changing climate. *Front. Mar. Sci.* **6**, 510 (2019). doi: [10.3389/fmars.2019.00510](https://doi.org/10.3389/fmars.2019.00510)
106. C. A. Bost *et al.*, Large-scale climatic anomalies affect marine predator foraging behaviour and demography. *Nat. Commun.* **6**, 8220 (2015). doi: [10.1038/ncomms9220](https://doi.org/10.1038/ncomms9220); pmid: [26506134](https://pubmed.ncbi.nlm.nih.gov/26506134/)
107. R. Cristofari *et al.*, Climate-driven range shifts of the king penguin in a fragmented ecosystem. *Nat. Clim. Chang.* **8**, 245–251 (2018). doi: [10.1038/s41558-018-0084-2](https://doi.org/10.1038/s41558-018-0084-2)
108. J. M. Alonso Vega, J. A. Vásquez, A. H. Buschmann, Population biology of the subtidal kebs *Macrocystis integrifolia* and *Lessonia trabeculata* (Laminariales, Phaeophyceae) in an upwelling ecosystem of northern Chile: Interannual variability and El Niño 1997–1998. *Rev. Chil. Hist. Nat.* **78**, 33–50 (2005). doi: [10.4067/S0716-078X2005000100004](https://doi.org/10.4067/S0716-078X2005000100004)
109. F. Trillmich *et al.*, On the challenge of interpreting census data: Insights from a study of an endangered pinniped. *PLOS ONE* **11**, e0154588 (2016). doi: [10.1371/journal.pone.0154588](https://doi.org/10.1371/journal.pone.0154588); pmid: [27148735](https://pubmed.ncbi.nlm.nih.gov/27148735/)
110. T. Spencer, K. A. Teleki, C. Bradshaw, M. D. Spalding, Coral bleaching in the Southern Seychelles during the 1997–1998 Indian Ocean warm event. *Mar. Pollut. Bull.* **40**, 569–586 (2000). doi: [10.1016/S0025-326X\(00\)00026-6](https://doi.org/10.1016/S0025-326X(00)00026-6)
111. L. G. C. Genevier, T. Jamil, D. E. Raitso, G. Krokos, I. Hoteit, Marine heatwaves reveal coral reef zones susceptible to bleaching in the Red Sea. *Glob. Chang. Biol.* **25**, 2338–2351 (2019). doi: [10.1111/gcb.14652](https://doi.org/10.1111/gcb.14652); pmid: [30974020](https://pubmed.ncbi.nlm.nih.gov/30974020/)
112. A. Genin, L. Levy, G. Sharon, D. E. Raitso, A. Diamant, Rapid onsets of warming events trigger mass mortality of coral reef fish. *Proc. Natl. Acad. Sci. U.S.A.* **117**, 25378–25385 (2020). doi: [10.1073/pnas.2009748117](https://doi.org/10.1073/pnas.2009748117); pmid: [32958634](https://pubmed.ncbi.nlm.nih.gov/32958634/)
113. C. N. Bianchi *et al.*, Consequences of the marine climate and ecosystem shift of the 1980–90s on the Ligurian Sea biodiversity (NW Mediterranean). *Eur. Zool. J.* **86**, 458–487 (2019). doi: [10.1080/24750263.2019.1687765](https://doi.org/10.1080/24750263.2019.1687765)
114. R. Cupido *et al.*, Sexual structure of a highly reproductive, recovering gorgonian population: Quantifying reproductive output. *Mar. Ecol. Prog. Ser.* **469**, 25–36 (2012). doi: [10.3354/meps09976](https://doi.org/10.3354/meps09976)
115. J. Garrabou *et al.*, Mass mortality in Northwestern Mediterranean rocky benthic communities: Effects of the 2003 heat wave. *Glob. Change Biol.* **15**, 1090–1103 (2009). doi: [10.1111/j.1365-2486.2008.01823.x](https://doi.org/10.1111/j.1365-2486.2008.01823.x)
116. E. Diaz-Almela, N. Marbà, C. M. Duarte, Consequences of Mediterranean warming events in seagrass (*Posidonia oceanica*) flowering records. *Glob. Change Biol.* **13**, 224–235 (2007). doi: [10.1111/j.1365-2486.2006.01260.x](https://doi.org/10.1111/j.1365-2486.2006.01260.x)
117. C. M. Eakin *et al.*, Caribbean corals in crisis: Record thermal stress, bleaching, and mortality in 2005. *PLOS ONE* **5**, e13969 (2010). doi: [10.1371/journal.pone.0013969](https://doi.org/10.1371/journal.pone.0013969); pmid: [21125021](https://pubmed.ncbi.nlm.nih.gov/21125021/)
118. Food and Agriculture Organization of the United Nations, "Climate change adaptation in the eastern Caribbean fisheries sector (CC4FISH) launching workshop, Barbados, 7–9 February 2017" (FAO Fisheries and Aquaculture Report No. 1189, 2017); <http://www.fao.org/3/i8281en/i8281EN.pdf>.
119. G. A. Kendrick *et al.*, A systematic review of how multiple stressors from an extreme event drove ecosystem-wide loss of resilience in an iconic seagrass community. *Front. Mar. Sci.* **6**, 455 (2019). doi: [10.3389/fmars.2019.00455](https://doi.org/10.3389/fmars.2019.00455)
120. D. A. Smale, T. Wernberg, Extreme climatic event drives range contraction of a habitat-forming species. *Proc. Biol. Sci.* **280**, 20122829 (2013). doi: [10.1098/rspb.2012.2829](https://doi.org/10.1098/rspb.2012.2829); pmid: [23325774](https://pubmed.ncbi.nlm.nih.gov/23325774/)
121. R. C. Babcock *et al.*, Severe continental-scale impacts of climate change are happening now: Extreme climate events impact marine habitat forming communities along 45% of Australia's coast. *Front. Mar. Sci.* **6**, 411 (2019). doi: [10.3389/fmars.2019.00411](https://doi.org/10.3389/fmars.2019.00411)
122. S. D. Roberts, P. D. Van Ruth, C. Wilkinson, S. S. Bastianello, M. S. Bansemer, Marine heatwave, harmful algae blooms and an extensive fish kill event during 2013 in South Australia. *Front. Mar. Sci.* **6**, 610 (2019). doi: [10.3389/fmars.2019.00610](https://doi.org/10.3389/fmars.2019.00610)
123. C. D. Harvell *et al.*, Disease epidemic and a marine heat wave are associated with the continental-scale collapse of a pivotal predator (*Pycnopodia helianthoides*). *Sci. Adv.* **5**, eaau7042 (2019). doi: [10.1126/sciadv.aau7042](https://doi.org/10.1126/sciadv.aau7042); pmid: [30729157](https://pubmed.ncbi.nlm.nih.gov/30729157/)
124. E. Sanford, J. L. Sones, M. García-Reyes, J. H. R. Goddard, J. L. Largier, Widespread shifts in the coastal biota of northern California during the 2014–2016 marine heatwaves. *Sci. Rep.* **9**, 4216 (2019). doi: [10.1038/s41598-019-40784-3](https://doi.org/10.1038/s41598-019-40784-3); pmid: [30862867](https://pubmed.ncbi.nlm.nih.gov/30862867/)
125. R. M. Suryan *et al.*, Ecosystem response persists after a prolonged marine heatwave. *Sci. Rep.* **11**, 6235 (2021). doi: [10.1038/s41598-021-83818-5](https://doi.org/10.1038/s41598-021-83818-5); pmid: [33737519](https://pubmed.ncbi.nlm.nih.gov/33737519/)
126. S. M. Mendez, A. Martinez, L. Ortego, A. Fabre, High sea surface temperature, the potential trigger of mass mortality of fish, exceptional toxin producing HABs, and other socio-economic impacts in Uruguay. *Harmful Algae News* **51**, 1–2 (2015); www.e-pages.dk/ku/114/.
127. B. Vargas-Ángel *et al.*, El Niño-associated catastrophic coral mortality at Jarvis Island, central Equatorial Pacific. *Coral Reefs* **38**, 731–741 (2019). doi: [10.1007/s00338-019-01838-0](https://doi.org/10.1007/s00338-019-01838-0)
128. R. E. Brainard *et al.*, Perspective, Special Supplement to the Bulletin of the American Meteorological Society, S. C. Herring, N. Christidis, A. Hoell, J. P. Kossin, C. J. Schreck III, P. A. Stott, Eds. (American Meteorological Society, 2018), vol. 99, pp. S21–S26; <https://doi.org/10.1175/BAMS-ExplainingExtremeEvents2016.1>
129. M. Le Nohaïc *et al.*, Marine heatwave causes unprecedented regional mass bleaching of thermally resistant corals in northwestern Australia. *Sci. Rep.* **7**, 14999 (2017). doi: [10.1038/s41598-017-14794-y](https://doi.org/10.1038/s41598-017-14794-y); pmid: [29101362](https://pubmed.ncbi.nlm.nih.gov/29101362/)
130. K. Mathiesen, "Ocean heatwave destroys Tasmania's unique underwater jungle." *Climate Home News*, 14 October 2016; <https://www.climatechangenews.com/2016/10/14/ocean-heatwave-destroys-tasmanias-unique-underwater-jungle/>.
131. J. E. Walsh *et al.*, 8. The high latitude marine heat wave of 2016 and its impacts on Alaska. *Bull. Am. Meteorol. Soc.* **99**, S39–S43 (2018). doi: [10.1175/BAMS-D-17-0105.1](https://doi.org/10.1175/BAMS-D-17-0105.1)
132. G. A. Mehl *et al.*, Sustained ocean changes contributed to sudden Antarctic sea ice retreat in late 2016. *Nat. Commun.* **10**, 14 (2019). doi: [10.1038/s41467-018-07865-9](https://doi.org/10.1038/s41467-018-07865-9); pmid: [30600315](https://pubmed.ncbi.nlm.nih.gov/30600315/)
133. C. M. Eakin *et al.*, Ding, dong, the witch is dead (?)—Three years of global coral bleaching 2014–2017. *Reef Encounter* **32**, 33–38 (2017); https://www.ncei.noaa.gov/data/oceans/coris/library/NOAA/CRCP/NESDIS/STAR/Project/915/Eakin2017_2014-2017_Coral_Bleaching.pdf.
134. T. P. Hughes *et al.*, Global warming and recurrent mass bleaching of corals. *Nature* **543**, 373–377 (2017). doi: [10.1038/nature21707](https://doi.org/10.1038/nature21707); pmid: [28300113](https://pubmed.ncbi.nlm.nih.gov/28300113/)
135. R. D. Stuart-Smith, C. J. Brown, D. M. Ceccarelli, G. J. Edgar, Ecosystem restructuring along the Great Barrier Reef following mass coral bleaching. *Nature* **560**, 92–96 (2018). doi: [10.1038/s41586-018-0359-9](https://doi.org/10.1038/s41586-018-0359-9); pmid: [30046108](https://pubmed.ncbi.nlm.nih.gov/30046108/)
136. L. J. McKenzie *et al.*, "Marine Monitoring Program: Annual report for inshore seagrass monitoring 2016–2017 (Great Barrier Reef Marine Park Authority, 2018); <https://elibrary.gbrmpa.gov.au/jspui/handle/11017/3398>.
137. A. Thompson *et al.*, Australian Institute of Marine Science, "Annual Report for inshore coral reef monitoring 2016–2017" (Great Barrier Reef Marine Park Authority, 2018); <https://elibrary.gbrmpa.gov.au/jspui/handle/11017/3397>.
138. A. Thompson, P. Costello, J. Davidson, M. Logan, G. Coleman, Australian Institute of Marine Science, "Annual Report for inshore coral reef monitoring 2017–2018" (Great Barrier Reef Marine Park Authority, 2019); <https://elibrary.gbrmpa.gov.au/jspui/handle/11017/3491>.
139. M. J. Salinger *et al.*, The unprecedented coupled ocean-atmosphere summer heatwave in the New Zealand region 2017/18: Drivers, mechanisms and impacts. *Environ. Res. Lett.* **14**, 044023 (2019). doi: [10.1088/1748-9326/ab012a](https://doi.org/10.1088/1748-9326/ab012a)
140. G. Gao *et al.*, Drivers of marine heatwaves in the East China Sea and the South Yellow Sea in three consecutive summers during 2016–2018. *J. Geophys. Res. Oceans* **125**, 1–19 (2020). doi: [10.1029/2020JC016518](https://doi.org/10.1029/2020JC016518)
141. T. Singh, M. Iijima, K. Yasumoto, K. Sakai, Effects of moderate thermal anomalies on Acropora corals around Sesoko Island, Okinawa. *PLOS ONE* **14**, e0210795 (2019). doi: [10.1371/journal.pone.0210795](https://doi.org/10.1371/journal.pone.0210795); pmid: [30699163](https://pubmed.ncbi.nlm.nih.gov/30699163/)
142. Fish Information & Services, "Strong heatwave causes mortality in aquaculture farms." *Seafood Media Group*, 10 August 2017; <https://seafood.media/fis/worldnews/worldnews.asp?e=e&id=93266&ndb=1>.
143. N. Phongsuwan *et al.*, Status and changing patterns on coral reefs in Thailand during the last two decades. *Deep. Res. Part II Top. Stud. Oceanogr.* **96**, 19–24 (2013). doi: [10.1016/j.jdsr.2013.02.015](https://doi.org/10.1016/j.jdsr.2013.02.015)
144. G. Manta, S. de Mello, R. Trinchin, J. Badagian, M. Barreiro, The 2017 record marine heatwave in the Southwestern Atlantic shelf. *Geophys. Res. Lett.* **45**, 12449–12456 (2018). doi: [10.1029/2018GL081070](https://doi.org/10.1029/2018GL081070)
145. L. J. Raymundo *et al.*, Successive bleaching events cause mass coral mortality in Guam, Micronesia. *Coral Reefs* **38**, 677–700 (2019). doi: [10.1007/s00338-019-01836-2](https://doi.org/10.1007/s00338-019-01836-2)
146. J. A. Burt, F. Paparella, N. Al-Mansoori, A. Al-Mansoori, H. Al-Jailani, Causes and consequences of the 2017 coral bleaching event in the southern Persian/Arabian Gulf. *Coral Reefs* **38**, 567–589 (2019). doi: [10.1007/s00338-019-01767-y](https://doi.org/10.1007/s00338-019-01767-y)
147. D. Y. Ige, S. D. Case, "11/5/19 – Coral bleaching not as severe as predicted but still widespread; extensive surveys show bleaching event now abating" (Department of Land and Natural Resources, 2019); <https://dlnr.hawaii.gov/blog/2019/11/05/nr19-186/>.
148. E. Rardon, "Rare tropical sea turtle rescued from Alberni Inlet." *Alberni Valley News*, 3 October 2019; <https://www.albernivalleynews.com/news/tropical-sea-turtle-rescued-from-alberni-inlet/>.
149. D. Hammock, "Commercial Dungeness crab season opener delayed through at least mid-December." *The Daily World*, 30 November 2020; <https://www.thedailyworld.com/news/commercial-dungeness-crab-season-opener-delayed-through-at-least-mid-december/>.
150. J. L. Boldt, A. Javorski, P. C. Chandler, *State of the Physical, Biological and Selected Fishery Resources of Pacific Canadian Marine Ecosystems in 2019: Canadian Technical Report of Fisheries and Aquatic Sciences 377* (Fisheries and Oceans Canada, 2020); https://publications.gc.ca/collections/collection_2020/mpo-dfo/Fs97-6-3377-eng.pdf.

ACKNOWLEDGMENTS

We thank the Coastal Observation and Seabird Survey Team (<https://COASST.org>) for providing the seabird mortality data presented in Fig. 2. **Funding:** D.A.S. was supported by a UK Research and Innovation (UKRI) Future Leaders Fellowship (MR/S032827/1). T.W. was supported by the Australian Research Council (DP200100201). M.T. was supported by The Brian Mason Trust, New Zealand. P.J.M. and M.T.B. were supported by NERC-Newton Fund grant NE/S011692/2. **Author contributions:** A.S.G., A.J.H., M.T.B., M.T., P.J.M., T.W., and D.A.S. conceived the research. K.E.S. conducted the systematic analysis of the literature. K.E.S. and D.A.S. wrote the original draft of the manuscript. All authors contributed to reviewing and editing the manuscript. **Competing interests:** The authors declare no competing financial interests. **Data and materials availability:** All data are available in the main text or the supplementary materials.

10.1126/science.abj3593