

both pools independently during stress recovery to show that those mRNAs in granules remained translationally repressed, whereas the nonsequestered pool went on to initiate translation.

Halstead *et al.* also demonstrated that the method is applicable to whole-animal studies by constructing transgenic flies (*Drosophila*) bearing the requisite fluorescent protein fusions, as well as an *oskar* mRNA-based TRICK reporter. The maternally produced *oskar* mRNA is transported from nurse cells in which it is made into the oocyte, where it localizes to the posterior pole at later stages and is necessary for germ cell formation (9). Halstead *et al.* were able to confirm that *oskar* mRNA is translationally repressed until it localizes to the posterior pole.

By design, the TRICK method assays the first round of translation. The basis of the method, i.e., the ability of the ribosome to strip off proteins from newly synthesized mRNAs, is also used by cells as a means of quality control via nonsense-mediated mRNA decay (NMD). During pre-mRNA splicing, the exon-junction complex (EJC) is deposited upstream of exon-exon junctions (10). Generally, mRNAs containing premature termination codons (PTCs) are recognized as abnormal because during the first, or pioneer round of translation (11, 12), translating ribosomes do not strip off EJCs after the PTC. (13). Aside from clear utility as a sensor for the NMD-critical pioneer round of translation, the TRICK method will doubtless find useful applications in other areas in which translational repression and/or bursts of localized translation control important processes such as long-term potentiation in neurons during learning (14) and disruption of RNA granule formation during viral infection (15). ■

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## CLIMATE AND CONSERVATION

# Creating a safe operating space for iconic ecosystems

Manage local stressors to promote resilience to global change

By M. Scheffer<sup>1\*</sup>, S. Barrett<sup>2</sup>, S. R. Carpenter<sup>3</sup>, C. Folke<sup>4</sup>, A. J. Green<sup>5</sup>, M. Holmgren<sup>6</sup>, T. P. Hughes<sup>7</sup>, S. Kosten<sup>8</sup>, I. A. van de Leemput<sup>1</sup>, D. C. Nepstad<sup>9</sup>, E. H. van Nes<sup>1</sup>, E. T. H. M. Peeters<sup>1</sup>, B. Walker<sup>10</sup>

**A**lthough some ecosystem responses to climate change are gradual, many ecosystems react in highly nonlinear ways. They show little response until a threshold or tipping point is reached where even a small perturbation may trigger collapse into a state from which recovery is difficult (1). Increasing evidence shows that the critical climate level for such collapse may be altered by conditions that can be managed locally. These synergies between local stressors and climate change provide potential opportunities for proactive management. Although their clarity and scale make such local

**POLICY** approaches more conducive to action than global greenhouse gas management, crises in iconic UNESCO World Heritage sites illustrate that such stewardship is at risk of failing.

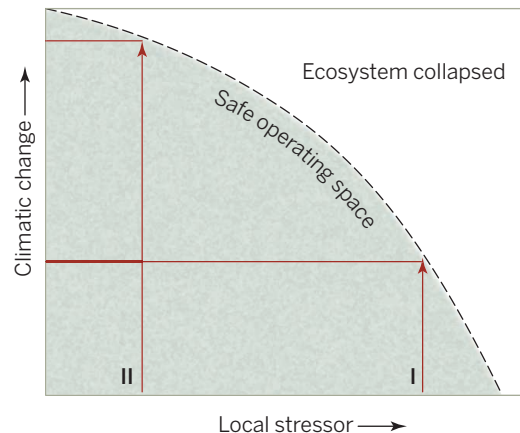
The term “safe operating space” frames the problem of managing our planet in terms of staying within acceptable levels or “boundaries” for global stressors (2). Uncertainty is accounted for by aiming to keep well on the safe side of such boundaries. The safe levels of different stressors at global scales are mostly considered independently. However, in ecosystems a safe level for one stressor is often strongly dependent on the level of other stressors. This implies that if such synergies are understood, local stressors may be effectively managed to enhance tolerance to global climate change (see first figure).

**LOCAL AND FEASIBLE.** The feasibility of managing the climate sensitivity of ecosystems is becoming increasingly evident. Obviously, local interventions are no panacea for the threats of climatic change. For example, melting of arctic sea ice with its far-reaching ecological consequences cannot be arrested by local management. However, ways of

building climate resilience are emerging for a variety of ecosystems, ranging from control of local sources of ocean acidification (3) to management of grazing pressure on dry ecosystems (4). We focus here on lakes, coral reefs, and tropical forests.

In lakes, warming and nutrient loading have similar effects on the likelihood that the ecosystem will tip into encroachment by floating plants or into dominance by toxic cyanobacteria (5). Experiments and field studies on different scales revealed intricate mechanisms that drive the synergy between effects of warming and nutrient load—e.g., boosted nutrient cycling—and shifts in the competitive advantage that favor small, rapidly reproducing fish species, cyanobacteria, and floating plants (5, 6). Although the synergy of climate and nutrient stressors implies double jeopardy to many wetlands, the good news is that reducing the nutrient load can compensate for the effects of warming. For example, data from lakes across continents and climate zones suggest that a reduction in nutrient concentrations by one-third can compensate for the effect of a 1°C increase in water temperature when it comes to the risk of cyanobacterial dominance (6).

In coral reefs, resilience depends strongly on locally manageable stressors such as fish-



**Schematic representation of safe operating space.** In ecosystems at risk of collapse, safe boundaries for local stressors, such as harvest rates or pollution, often change with climate change. A local stressor that is currently at a safe level (I) needs to be adjusted to a lower value to keep the system within the safe operating space in a future climate (II).

ing pressure and water quality. For example, the takeover of most Caribbean reefs by seaweeds was triggered by sea-urchin mortality but was facilitated in many locations by high nutrient loading and overharvesting of fish functional groups that controlled the seaweeds (7). On the Great Barrier Reef, coral

recovery rates after the 1998 bleaching event were markedly suppressed by experimental exclusion of herbivorous fishes (8). Local conservation efforts can help in maintaining and enhancing resilience and in limiting longer-term damage from bleaching and other climate-related effects.

In tropical forests, resilience is under pressure from climate change as well as local stressors such as deforestation, logging, and fire (9). Forests become stressed by increases in temperature (10) and by greater rainfall variability (4). One important near-term risk from drought is a self-reinforcing shift to a contrasting fire-maintained state. Recent experiments confirm cascading effects of a decline in canopy cover, which favor invasion by flammable grasses (11). The removal of trees makes the forest more fire-prone, increasing the risk of further transition to open woodland in dry years (9, 11). In addition, there is a substantial positive feedback effect of forest cover on regional precipitation, implying that loss of forest contributes to overall reduction in rainfall (9). Thus, maintaining a critical mass of forested areas and preventing opening of the closed canopy structure are powerful tools to enhance the safe operating space of tropical forests in the face of rising drought risks.

**WORLD HERITAGE AT RISK.** Despite the solid scientific basis for managing climate resilience in such ecosystems, failure to do so is putting globally important ecosystems at risk. We highlight crises faced by three iconic World Heritage Areas (see second figure).

The Doñana wetlands in southern Spain provide the most important wintering site for waterfowl in Europe. They contain the largest temporary pond complex in Europe, with a diversity of amphibians and invertebrates. Despite the site's protected status, the marshes are threatened by eutrophication due to pollution and reduced flow of incoming streams, promoting toxic cyanobacterial blooms, and dominance by invasive floating plants that create anoxic conditions in the water. In addition, groundwater extraction for strawberry culture and beach tourism also has major effects (12). Little has been done to control these local stressors, leaving Doñana unnecessarily vulnerable to climate change. UNESCO has just rated this World Heritage Site as under "very high threat."

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Examples of iconic ecosystems where climate change may trigger transitions to a different state. From top to bottom: the Doñana wetlands, the Amazon rainforest, and the Great Barrier Reef.

The Great Barrier Reef is the largest coral system in the world. In response to multiple threats, fishing has been prohibited since 2004 over 33% of the Great Barrier Reef Marine Park, and efforts have begun to reduce runoff of nutrients, pesticides, herbicides, and sediments from land. However, these interventions may be too little, too late. Approximately half of the coral cover has been lost in recent decades (13), and the outlook is “poor, and declining,” with climate change, coastal development, and dredging as major future threats (14). The World Heritage Committee has warned that in the absence of a solid long-term plan, it would consider listing the reef as “in danger” in 2015 (15).

The Amazon rainforest is one of the world’s great biological treasures and a vital component of Earth’s climate system. Yet this ecosystem is under increasing pressure from climate change as well as local stressors such as logging and forest fire (9). Brazil has shown leadership by slowing down Amazon deforestation by 70% (16) and by creating the largest protected area (PA) network in the world. Yet these successes are now being partially undermined by major infrastructure and natural resource extraction projects and by shifts in legislation (17).

**FRAMING FOR ACTION.** The evidence we have for enhancing climate resilience of ecosystems places direct responsibility on governments to ensure implementation. However, investment will only happen if costs of refraining from activities that undermine resilience are distributed in ways that lead to effective action. Realizing such incentive schedules may be challenging. However, there are three specific reasons that building a safe operating space for ecosystems by controlling local stressors is more conducive to immediate action than global control of greenhouse gases.

*From global to local commons.* Potential incentives for local protection are much stronger than those to supply the global public good of abating greenhouse gas emissions (18), for the same reason that countries tend to favor adaptation over mitigation. Mitigation requires global collective action and is vulnerable to free riding, whereas adaptation can be done unilaterally, with benefits accruing almost exclusively to the country doing the adaptation. However, iconic ecosystems also provide a global public good. This is why they are on the World Heritage list in the first place. In some cases, the local interventions can result in substantial global mitigation. For instance, slowing down Amazon deforestation made Brazil a global leader in climate change mitigation (16).

*From high to low uncertainty.* Perceived uncertainty has often paralyzed policy (19), and experimental evidence suggests that uncertainty about climate change tipping points undermines efforts to avoid crossing a dangerous threshold (20). There is less uncertainty on the ecosystem level than on the global level when it comes to effects of management options.

*From negative to positive framing.* Gloom-and-doom perceptions may backfire to block action. Terms such as “extreme events” and “catastrophic transitions” may express the urgency of the matter. However, social experiments reveal that accounts of disastrous future effects of climate change can invoke cognitive dissonance that causes many people to disbelieve climate change altogether. This response disappears if a feasible approach to take action and abate the problems is presented simultaneously (21). A positive, action-oriented framing of a safe operating space for the world’s iconic ecosystems may help stimulate societal consensus that climate change is real and should be addressed. ■

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**STEM CELLS**

# Holding your breath for longevity

A nutrient-sensing protein is important for the health of hematopoietic stem cells during aging

By Alejandro Ocampo and Juan Carlos Izpisua Belmonte

**A**ging is a complex process. Progressive molecular changes lead to a decline in the ability of living beings to maintain homeostasis and overcome cellular stress, protein damage, and disease (1). At the organismal level, stem cells play a fundamental role in maintaining tissue integrity, and their functional and proliferative exhaustion is a major cause of aging (2). Hematopoietic stem cells, which reside in the bone marrow and give rise to all blood cell types, are a favored model for studying stem cell aging.

*“...decline of stem cell function with age due to intrinsic factors...and extrinsic factors...could potentially be reverted...”*

However, the exact molecular mechanisms underlying their aging remain unknown. SirTuins, a family of nutrient-sensing proteins (SIRT1 to SIRT7) that regulate gene expression and protein function in mammalian cells, orchestrate multiple pathways that are associated with age-related processes and longevity. On page 1374 of this issue, Mohrin *et al.* (3) connect SIRT7 to a metabolic checkpoint that controls aging in hematopoietic stem cells.

SIRT7 is a nicotinamide adenine dinucleotide (NAD)-dependent deacetylase that senses the nutrient status of a cell and adjusts cell function accordingly. Thus, under nutrient deprivation, SIRT7 alters transcription to reduce cell metabolism and decrease cell growth, thereby promoting cell survival. By analyzing all proteins that