

## Characteristics, drivers and feedbacks of global greening

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**Abstract** | Vegetation greenness has been increasing globally since at least 1981, when satellite technology enabled large-scale vegetation monitoring. The greening phenomenon, together with warming, sea-level rise and sea-ice decline, represents highly credible evidence of anthropogenic climate change. In this Review, we examine the detection of the greening signal, its causes and its consequences. Greening is pronounced over intensively farmed or afforested areas, such as in China and India, reflecting human activities. However, strong greening also occurs in biomes with low human footprint, such as the Arctic, where global change drivers play a dominant role. Vegetation models suggest that CO<sub>2</sub> fertilization is the main driver of greening on the global scale, with other factors being notable at the regional scale. Modelling indicates that greening could mitigate global warming by increasing the carbon sink on land and altering biogeophysical processes, mainly evaporative cooling. Coupling high temporal and fine spatial resolution remote-sensing observations with ground measurements, increasing sampling in the tropics and Arctic, and modelling Earth systems in more detail will further our insights into the greening of Earth.

### Afforestation

The conversion of treeless lands to forests through planting trees.

Vegetation controls the exchange of carbon, water, momentum and energy between the land and the atmosphere, and provides food, fibre, fuel and other valuable ecosystem services<sup>1,2</sup>. Changes in vegetation structure and function are driven by climatic and environmental changes, and by human activities such as land-use change. Given that increased carbon storage in vegetation, such as through afforestation, could combat climate change<sup>3,4</sup>, quantifying vegetation change and its impact on carbon storage and climate has elicited considerable interest from scientists and policymakers.

However, it is not possible to detect vegetation changes at the global scale using ground-based observations due to the heterogeneity of change and the lack of observations that can detect these changes both spatially and temporally. While monitoring the changes in some vegetation properties (for example, stem-size distribution and below-ground biomass) at the global scale remains impossible, satellite-based remote sensing has enabled continuous estimation of a few important metrics, including vegetation greenness, since the 1980s (BOX 1).

In 1986, a pioneering study by Tucker et al.<sup>5</sup> on remotely sensed normalized difference vegetation index (NDVI; a radiometric measure of vegetation greenness) (BOX 1) revealed a close connection between vegetation canopy greenness and photosynthesis activity (as inferred from seasonal variations in atmospheric

CO<sub>2</sub> concentration). This index was successfully used to constrain vegetation primary production globally<sup>6</sup>. Using NDVI data from 1981 to 1991, Myneni et al.<sup>7</sup> reported an increasing trend in vegetation greenness in the Northern Hemisphere, which was subsequently observed across the globe<sup>8–13</sup>. This ‘vegetation greening’ is defined as a statistically significant increase in annual or seasonal vegetation greenness at a location resulting, for instance, from increases in average leaf size, leaf number per plant, plant density, species composition, duration of green-leaf presence due to changes in the growing season and increases in the number of crops grown per year.

There has also been considerable interest in understanding the mechanisms or drivers of greening<sup>11,14</sup>. Lucht et al.<sup>14</sup> and Xu et al.<sup>10</sup> revealed that warming has eased climatic constraints, facilitating increasing vegetation greenness over the high latitudes. Zhu et al.<sup>11</sup> further investigated key drivers of greenness trends and concluded that CO<sub>2</sub> fertilization is a major factor driving vegetation greening at the global scale. Subsequent studies based on fine-resolution and medium-resolution satellite data<sup>13</sup> have shown the critical role of land-surface history, including afforestation and agricultural intensification, in enhancing vegetation greenness. The large spatial scale of vegetation greening and the robustness of its signal have led the Intergovernmental Panel on Climate Change (IPCC) special report on climate change

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## Key points

- Long-term satellite records reveal a significant global greening of vegetated areas since the 1980s, which recent data suggest has continued past 2010.
- Pronounced greening is observed in China and India due to afforestation and agricultural intensification.
- Global vegetation models suggest that CO<sub>2</sub> fertilization is the main driver of global vegetation greening.
- Warming is the major cause of greening in boreal and Arctic biomes, but has negative effects on greening in the tropics.
- Greening was found to mitigate global warming through enhanced land carbon uptake and evaporative cooling, but might also lead to decreased albedo that could potentially cause local warming.
- Greening enhances transpiration, a process that reduces soil moisture and runoff locally, but can either amplify or reduce runoff and soil moisture regionally through altering the pattern of precipitation.

and land<sup>15</sup> to list it, together with global-scale warming, sea-level rise<sup>16</sup> and sea-ice decline<sup>16</sup>, as highly credible evidence of the environmental impact of anthropogenic climate change.

Greener vegetation not only results from climatic and atmospheric changes but also feeds back to the climate through biogeochemical and biogeophysical processes. These feedbacks are often studied with Earth system models (ESMs), in which vegetation is coupled with the atmosphere and the hydrologic cycle<sup>17</sup>. ESM-based studies have demonstrated that greening can accelerate the hydrologic cycle by increasing the amount of water transpired by plants, alter the energy exchange between land and the atmosphere, and affect atmospheric circulation patterns<sup>18,19</sup>.

In this Review, we synthesize past and recent efforts to characterize the spatiotemporal patterns of vegetation greening since the 1980s. We discuss how rising atmospheric CO<sub>2</sub> concentration, climate change, land-use change and nitrogen deposition are the key drivers of greenness changes on the global and regional scale. We assess the impacts of vegetation greening on carbon, water and energy balances, and conclude by identifying key challenges and perspectives for future research.

## Greenness changes

Global-scale vegetation greening has been demonstrated using nearly four decades of NDVI and leaf area index (LAI) greenness data derived from the Advanced Very-High-Resolution Radiometer (AVHRR) instrument

(FIG. 1a,b). While early studies primarily used the NDVI to detect changes in global greenness, recent studies widely use the LAI, since it has clear physical interpretation and is a fundamental variable in almost all land-surface models (BOX 1). An ensemble of LAI datasets has shown that 52% ( $P < 0.05$ ) to 59% ( $P < 0.10$ ) of global vegetated lands displayed an increasing trend in growing season LAI since the 1980s<sup>11</sup> (FIG. 1a). Although some studies reported a stalling, or even a reversal, of the greening trend since 2000 based on AVHRR<sup>20</sup> and collection 5 (C5) of the Moderate Resolution Imaging Spectroradiometer (MODIS) data<sup>21</sup>, this signal might be an artefact of sensor degradation and/or processing<sup>22–24</sup>. For example, using a revised calibration of the MODIS data in the most recent collection 6 (C6) dataset<sup>24</sup>, Chen et al.<sup>13</sup> showed that leaf area increased by 5.4 million km<sup>2</sup> over 2000–2017, an area equivalent to the areal extent of the Amazon rainforest<sup>13</sup>. Indeed, 34% of vegetated land exhibited greening ( $P < 0.10$ ), whereas only 5% experienced browning ( $P < 0.10$ ), that is, a loss of vegetation greening.

New satellite-based vegetation indices also support the global greening trend observed since 2000 (FIG. 1), including the enhanced vegetation index (EVI) and near-infrared reflectance of terrestrial vegetation (NIRv) (BOX 1). However, while vegetation greenness is increasing at the global scale, the changes vary considerably between regions and seasons.

**Regional trends.** In the high northern latitudes ( $>50^{\circ}\text{N}$ ), AVHRR and Landsat records indicate a widespread increase in vegetation greenness since the 1980s<sup>8,12,25</sup> (FIG. 2a–d). Regions with the greatest greening trend include northern Alaska and Canada, the low-Arctic parts of eastern Canada and Siberia, and regions of Scandinavia<sup>12,25,26</sup>. Dendrochronological data and photographic evidence further corroborate these findings<sup>27–30</sup>. In general, the LAI over high northern latitudes will continue to increase by the end of this century<sup>31</sup>, based on the results of an ensemble of ESMs (FIG. 2e–h). However, although only 3% of the high latitudes show browning during 1982–2014 (REF.<sup>25</sup>), there is a growing proportion of Arctic areas exhibiting a browning trend<sup>32</sup>. Such trends first emerged in boreal forests, where a multitude of disturbances (for example, fires, harvesting and insect defoliation) prevail<sup>9,33–37</sup>. The North American boreal forests in particular exhibit browning areas nearly 20 times larger than the Eurasian boreal forests, showing heterogeneous regional greenness change<sup>38</sup>.

The northern temperate region ( $25^{\circ}\text{S}$ – $50^{\circ}\text{N}$ ) is another vegetation greening hotspot, experiencing faster rates of greening than the high latitudes since 2000 (FIG. 2b,d). Indeed, ~14 million km<sup>2</sup> of the temperate region greened ( $P < 0.10$ ), contributing about one-half of the global net leaf area increase over this time period<sup>13</sup>. The increase of vegetation greenness is especially strong in agricultural regions (for example, India<sup>13</sup>) and recently afforested areas (for example, China<sup>13,39</sup>); collectively, China and India alone contribute more than 30% of the total net increase in the global LAI<sup>13</sup>.

Tropical regions ( $25^{\circ}\text{S}$ – $25^{\circ}\text{N}$ ) are also greening (FIG. 2b,d), contributing about a quarter of the net global

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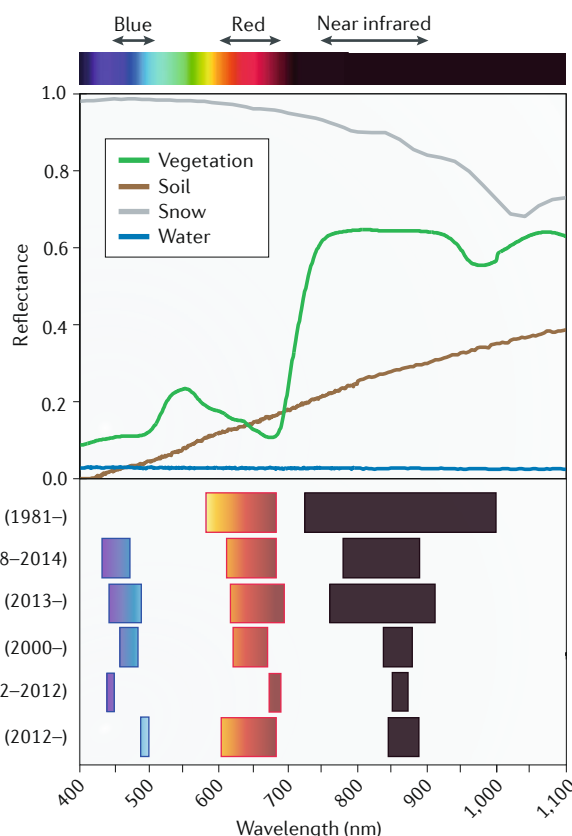
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# Box 1 | Remotely sensed vegetation greenness

Remotely sensed vegetation greenness generally refers to spectral vegetation indices (VIs) or the leaf area index (LAI). Photosynthetic pigments in plant leaves (mainly chlorophyll and carotenoids) strongly absorb photosynthetically active radiation, which largely overlaps with the visible spectrum (400–700 nm), particularly red wavelengths (620–700 nm). In the near-infrared (NIR) domain (700–1,300 nm), absorbance by leaf constituents is either small or absent; thus, scattering increases the likelihood that photons will exit the leaf. This is the biophysical basis for high leaf-level reflectance in the NIR region. At the canopy scale, structural properties such as LAI and leaf-angle distribution dominate variability in NIR reflectance<sup>176</sup>. This unique spectral signature of vegetation in the red and NIR channels, a characteristic not present in common non-vegetative features such as soil, snow and water<sup>177,178</sup>, has thus been utilized to derive numerical VIs measuring vegetation greenness<sup>176,179,180</sup> (Supplementary Table S1). For example, the normalized difference vegetation index, which is one of the most widely used VIs in assessing vegetation greenness and its changes from local to global scales (Supplementary Table S2), is useful for measuring canopy structural properties, such as leaf area, light interception and biomass<sup>41,181,182</sup>. Satellite sensors, such as the Advanced Very-High-Resolution Radiometer (AVHRR), Moderate Resolution Imaging Spectroradiometer (MODIS), Vegetation, Medium Resolution Imaging Spectrometer (MERIS) and Visible Infrared Radiometer Suite (VIIRS), have been deployed with varying temporal coverage, providing VI products based on a wide range of spectral-band specifications and data processing (Supplementary Table S3). For example, the AVHRR does not have a blue channel, so this sensor is unable to produce blue-band-based greenness indices like the enhanced vegetation index. These sensor differences make it a non-trivial challenge to produce consistent and continuing long-term greenness products<sup>183</sup>.

Compared with VIs, the LAI (the one-sided green leaf area per unit ground area in broadleaf canopies or one-half of the total needle surface area per unit ground area in coniferous canopies<sup>184,185</sup>) is a well-defined physical attribute of vegetation. The LAI is a state variable in all land models and key to quantifying the exchanges of mass, momentum and energy between the surface and the atmosphere. Multiple approaches for retrieving the LAI from remote sensing data have been developed — these can be conceptually categorized as: empirical approaches that are based on relationships between VIs and the LAI<sup>186,187</sup>; machine-learning approaches that train surface reflectance or VIs to given reference LAIs<sup>182,188,189</sup>; and physical approaches that are based on the physics of radiation interaction with elements of a canopy and transport within the vegetative medium<sup>184,190,191</sup>. See Supplementary Table S4 for currently available global LAI products.



increase in leaf area since 2000 (REF.<sup>13</sup>). However, the tropics also have areas where significant browning has been reported, for example, in the Brazilian Cerrado and Caatinga regions and Congolian forests<sup>13,40</sup>. It is worth noting that substantial uncertainties remain in the tropical vegetation greenness estimations due to the saturation effects of greenness indices in dense vegetation<sup>41</sup> and contamination by clouds and aerosols<sup>42</sup>. These uncertainties partly underlie the disagreement between the MODIS and AVHRR products<sup>13</sup> when measuring tropical greenness and the debate on whether the Amazonian forests have greened or browned in response to droughts<sup>42–44</sup>.

The extratropical Southern Hemisphere (>25°S) has experienced a general greening trend since the 1980s<sup>13,45</sup>, but it is lower than that in the temperate and high-latitude Northern Hemisphere<sup>13</sup> (FIG. 2a–d). Regional greening hotspots in southern Brazil and southeast Australia

mostly overlap with the intensive cropping areas<sup>13</sup>, highlighting the increasing contribution of managed ecosystems to vegetation greening. Note that most of this region is dominated by semi-arid ecosystems<sup>46</sup>, where vegetation coverage is generally sparse. Thus, satellite vegetation indices over this region are generally sensitive to change in soil background. For example, browning was detected from the AVHRR dataset since the 2000s<sup>20</sup> (FIG. 2b), but MODIS C6 data (which is better calibrated and can distinguish vegetation from background more accurately) instead showed an overall greening trend particularly since 2002 (REF.<sup>13</sup>; FIG. 2c,d).

**Seasonal changes of greenness.** In the northern temperate and high latitudes, greenness often shows distinctive seasonal patterns within a calendar year (FIG. 3). Several metrics of land-surface phenology have been developed

Land-surface phenology  
Cyclic phenomena in vegetated  
land surfaces observed from  
remote sensing.

## Carboxylation

The addition of CO<sub>2</sub> to ribulose 1,5-bisphosphate during photosynthesis.

to depict the seasonal cycle of greenness<sup>47</sup>, including the widely used start of the growing season (SOS) and end of the growing season (EOS)<sup>48</sup>. Although phenology dates can vary depending on the greenness product or algorithm used<sup>49–51</sup>, significant trends towards both earlier SOS (2–8 days decade<sup>–1</sup>) and later EOS (1–6 days decade<sup>–1</sup>) and, thus, longer lengths of the growing season (LOS) (2–10 days decade<sup>–1</sup>), have been observed in most Northern Hemisphere regions during the past four decades<sup>7,8,25,52–54</sup> (FIG. 3a–c). These trends are corroborated

by ground-based observation data in spring and autumn<sup>55–57</sup>. The increase in LOS is driven mainly by an advanced SOS in Eurasia (53–81% of LOS lengthening is due to SOS advance) and delayed EOS in North America (57–96% of LOS lengthening is due to EOS delay), with the more rapid total LOS increase seen in Eurasia<sup>25,58–60</sup>.

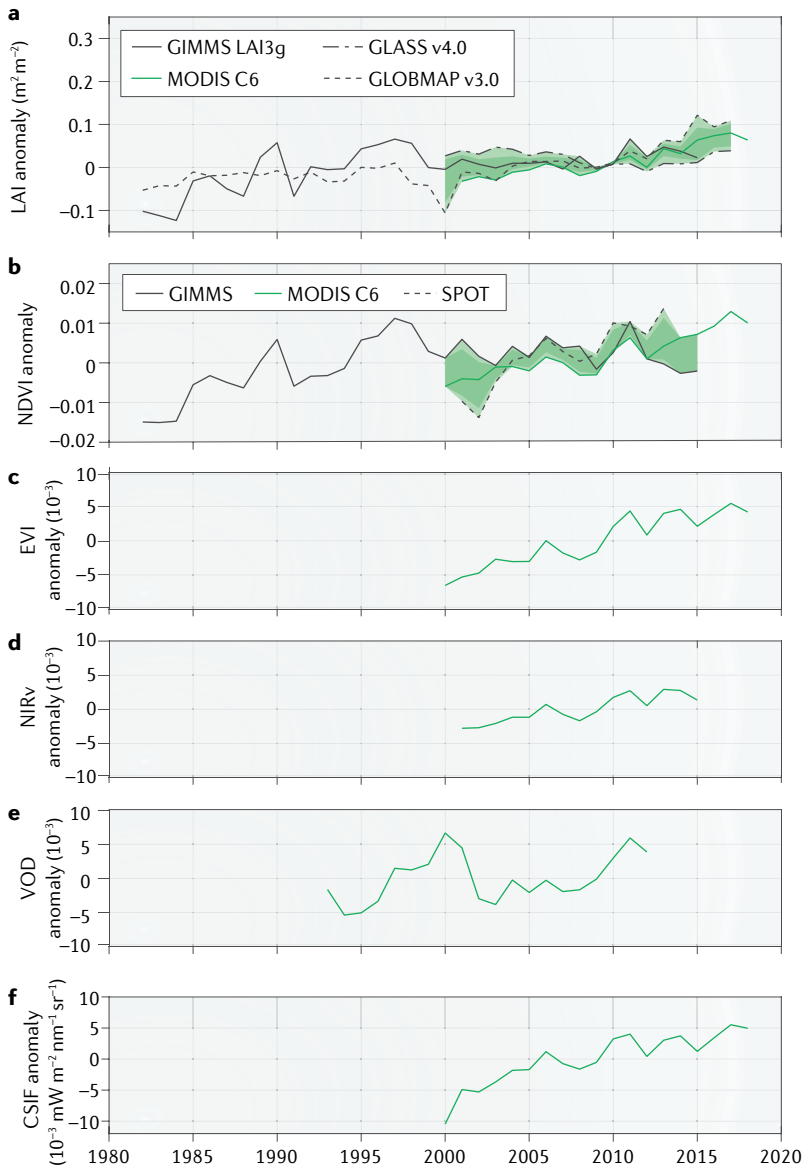
In addition to longer growing seasons, satellite greenness data also reveal important shifts in the timing and magnitude of the seasonal peak greenness<sup>47,61</sup>. For example, the timing of peak greenness has advanced by 1.2 days decade<sup>–1</sup> during 1982–2015 (REF.<sup>62</sup>) and 1.7 days decade<sup>–1</sup> during 2000–2016 (REF.<sup>61</sup>) over the extratropical Northern Hemisphere (FIG. 3a), with the boreal region peak greenness advancing twice as fast as the Arctic tundra and temperate ecosystem peaks<sup>61</sup>. Since the 1980s, the magnitude of the peak greenness has also increased over the extratropical Northern Hemisphere by ~0.1 standardized NDVI anomaly per year<sup>62</sup>, with a stronger signal in the pan-Arctic region<sup>63,64</sup>.

Phenology changes, including the SOS advancement, EOS delay and peak greenness enhancement, can significantly change the Earth's seasonal landscape. Northern high latitudes, which traditionally have high seasonality (that is, short and intense growing seasons), are exhibiting seasonality similar to that of their counterparts 6° to 7° south in the 1980s. In other words, the latitudinal isolines of northern vegetation seasonality have shifted southward since the 1980s. The diminished seasonality of the northern high-latitude vegetation<sup>10</sup> is consistent with changes in the velocity of vegetation greenness (defined as the ratio of temporal greenness change to its spatial gradient)<sup>65</sup>, which showed faster northward movement of the SOS ( $3.6 \pm 1.0$  km year<sup>–1</sup>) and the EOS ( $6.0 \pm 1.1$  km year<sup>–1</sup>) than the peak greenness ( $3.1 \pm 1.0$  km year<sup>–1</sup>) during 1982–2011 (REF.<sup>65</sup>).

## Drivers of greening

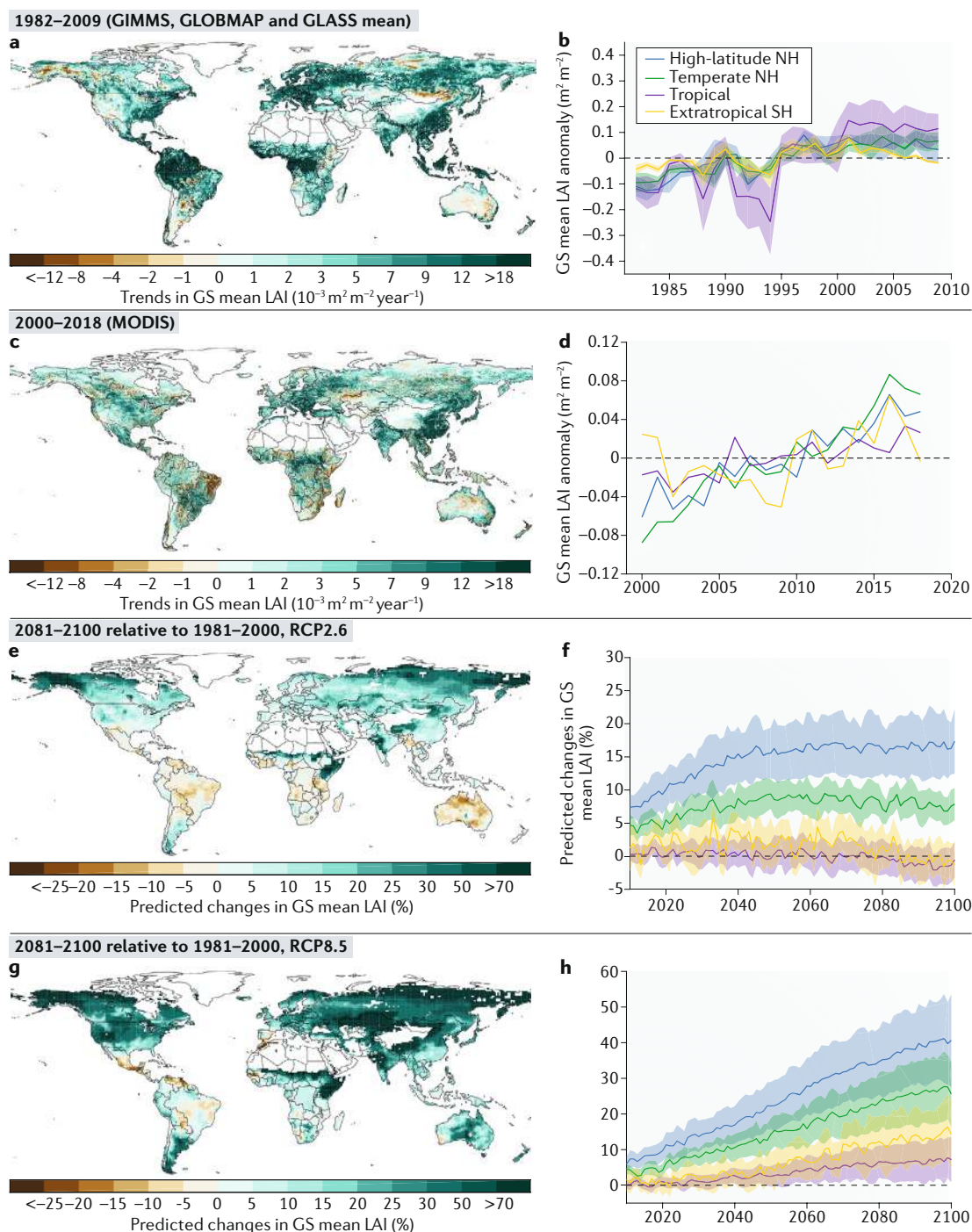
Several factors are thought to impact vegetation greening, including rising atmospheric CO<sub>2</sub> concentrations, climate change, nitrogen deposition and land-use changes. However, nonlinear impacts and interactions make it challenging to quantify the individual contribution of these factors to the observed greening trend. In this section, we review the contribution of several key drivers of vegetation greening and efforts to quantitatively attribute the observed greening trend to each of these factors.

**CO<sub>2</sub> fertilization.** As CO<sub>2</sub> is the substrate for photosynthesis, rising atmospheric CO<sub>2</sub> concentration can enhance photosynthesis<sup>66</sup> by accelerating the rate of carboxylation; this process is known as the ‘CO<sub>2</sub> fertilization effect’. In addition, increased CO<sub>2</sub> concentrations can also enhance vegetation greenness by partially closing leaf stomata, leading to enhanced water-use efficiency<sup>67</sup>, which should relax water limitation to plant growth, particularly over semi-arid regions<sup>45,68,69</sup>. Analysis of the ‘Trends and drivers of the regional-scale sources and sinks of carbon dioxide’ (TRENDY) ensemble of dynamic global vegetation models (DGVMs)<sup>70</sup> suggests that rising CO<sub>2</sub> is the dominant driver of vegetation greening, accounting for nearly 70% of global

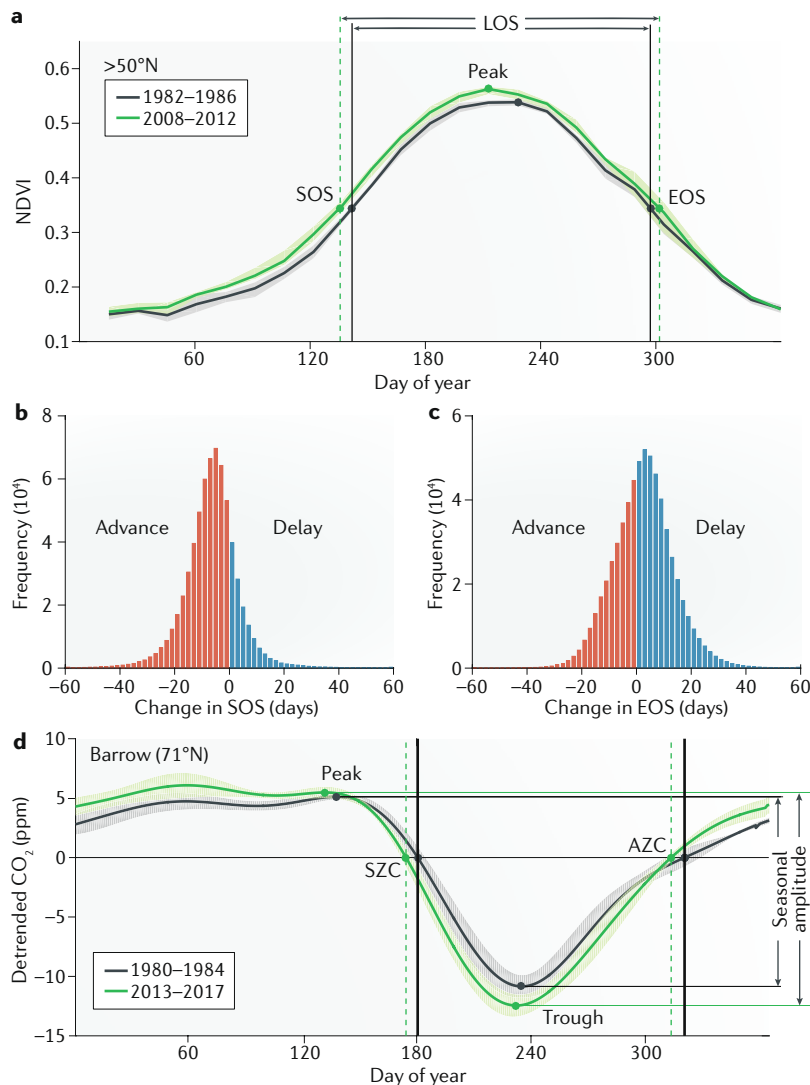


**Fig. 1 | Changes in satellite-derived global vegetation indices, vegetation optical depth and contiguous solar-induced fluorescence.** **a** | Leaf area index (LAI) from four products: GIMMS<sup>13</sup>, GLASS<sup>192</sup>, GLOBMAP<sup>23</sup> and Moderate Resolution Imaging Spectroradiometer (MODIS) C6 (REF.<sup>193</sup>). **b** | Normalized difference vegetation index (NDVI) from three products: GIMMS<sup>194</sup>, MODIS C6 (REF.<sup>195</sup>) and SPOT<sup>196</sup>. **c** | Enhanced vegetation index (EVI) from MODIS C6 (REF.<sup>195</sup>). **d** | Near-infrared reflectance of terrestrial vegetation (NIRv)<sup>197</sup>. **e** | Vegetation optical depth (VOD)<sup>119</sup>. **f** | Contiguous solar-induced fluorescence (CSIF)<sup>114</sup>. In parts **a** and **b**, the light-green shading denotes the range of LAI and NDVI across different products and the dark-green shading denotes the interquartile range (between the 25th and 75th percentiles). Only measurements during the growing season<sup>11</sup> were considered.





**Fig. 2 | Spatial patterns of changes in leaf area index.** **a** | Growing season (GS) mean Advanced Very-High-Resolution Radiometer (AVHRR) leaf area index (LAI) trend during 1982–2009. The AVHRR LAI dataset is the average of three different products (GIMMS<sup>13</sup>, GLOBMAP<sup>23</sup> and GLASS<sup>192</sup>). **b** | Change in the GS mean AVHRR LAI over four regions during 1982–2009. **c** | GS mean Moderate Resolution Imaging Spectroradiometer (MODIS) LAI during 2000–2018. **d** | Change in the GS mean MODIS LAI over four regions during 2000–2018. MODIS LAI is from collection 6 (REF.<sup>193</sup>). **e** | Relative change in GS mean LAI between 1981–2000 and 2081–2100 under the Representative Concentration Pathway 2.6 (RCP2.6), based on the Coupled Model Intercomparison Project Phase 5 (CMIP5) multi-model ensemble. **f** | Relative change in GS mean CMIP5 LAI 2018–2100 under RCP2.6, relative to 1981–2000. **g** | Relative change in GS mean LAI between 1981–2000 and 2081–2100 under RCP8.5, based on CMIP5. **h** | Relative change in GS mean CMIP5 LAI 2018–2100 under RCP8.5, relative to 1981–2000. The number of CMIP5 models used in the calculation of the multi-model mean is 16 and 19, for RCP2.6 and RCP8.5, respectively (Supplementary Table S5). In parts **a**, **c**, **e** and **g**, the white land areas depict barren lands, permanent ice-covered areas, permanent wetlands, built-up areas and water. In parts **b**, **d**, **f** and **h**, blue represents the high-latitude Northern Hemisphere (NH) (50–90°N), green represents the temperate NH (25–50°N), purple represents the tropical zone (25°S–25°N) and yellow represents the extratropical Southern Hemisphere (SH) (90–25°S). The shading shows the  $\pm 1$  inter-model standard deviation.



**Fig. 3 | Changes in the seasonality of vegetation greenness and atmospheric CO<sub>2</sub> concentration.** **a** | Five-year mean seasonal variations of the normalized difference vegetation index (NDVI) over Northern Hemisphere high latitudes (>50°N) during 1982–1986 (black line) and 2008–2012 (green line). Start of the growing season (SOS) and end of the growing season (EOS) are shown as 50% of the maximum NDVI. The length of the growing season (LOS) is the difference between the EOS and the SOS. **b** | Frequency distribution of SOS change in the Northern Hemisphere during 1982–2012. **c** | Frequency distribution of EOS change in the Northern Hemisphere during 1982–2012. **d** | Five-year mean detrended seasonal CO<sub>2</sub> variations at Barrow, AK, USA (71°N) (NOAA ESRL archive: <https://www.esrl.noaa.gov/gmd/ccgg/obspack/>) during 1980–1984 (black line) and 2013–2017 (green line). Vertical lines mark the spring zero-crossing date (SZC) and autumn zero-crossing date (AZC). Horizontal lines mark the seasonal amplitude as the difference between the maximum and the minimum of detrended seasonal CO<sub>2</sub> variations. Shaded areas show the range of interannual variations in the NDVI in part **a** and the standard deviation of the detrended CO<sub>2</sub> mole fraction (ppm) in part **d** at the day of year. NDVI data are the updated dataset from Tucker et al.<sup>194</sup>. Parts **b** and **c** are adapted with permission from REF.<sup>48</sup>, Wiley-VCH.

LAI trend since the 1980s<sup>11</sup> (FIG. 4). Statistical modelling also supports the important role of rising atmospheric CO<sub>2</sub> concentration in driving vegetation greening<sup>71,72</sup>. Free-air CO<sub>2</sub> enrichment (FACE) experiments show that elevating the CO<sub>2</sub> concentration by ~200 ppm above the ambient conditions significantly enhances vegetation productivity<sup>73</sup> and increases leaf area<sup>74</sup>. Different plant

species vary largely in the magnitude of LAI enhancement<sup>75</sup>, with the larger effect on forest stands having lower LAI at the ambient conditions<sup>76</sup>. In DGVMs, elevated CO<sub>2</sub> increases vegetation productivity more in tropical ecosystems than in temperate and boreal ecosystems<sup>11,77,78</sup> (FIG. 4b). However, the strength of the CO<sub>2</sub> fertilization effect can be limited by extreme weather events<sup>79,80</sup> and nutrient and water availability<sup>73,81,82</sup>. Indeed, nitrogen and phosphorus have been shown to regulate the global pattern of CO<sub>2</sub> fertilization effects<sup>83</sup>. Since nutrient processes were under-represented in the ESMs used in the IPCC Fifth Assessment Report (AR5), the predictions of continued greening trends through 2100 (REF.<sup>31</sup>) (FIGS 2e–h,5) might overestimate the CO<sub>2</sub> fertilization effects.

**Climate change.** Although rising atmospheric CO<sub>2</sub> concentration is the main driver of global greening, climate change, such as anthropogenic warming and regional trends in precipitation, is a dominant driver of greening changes over 28% of the global vegetated area<sup>11</sup>. The global contribution of climate change to increasing greenness is only 8% (FIG. 4a), however, because impacts of climate change on vegetation greenness vary between regions<sup>11</sup>. For example, warming could reduce vegetation growth in the tropics<sup>84</sup>, where ambient temperature is close to vegetation optimal temperature<sup>85</sup>, but warming significantly increases vegetation greenness in the boreal and Arctic regions<sup>86</sup> by enhancing metabolism<sup>87</sup> and extending the growing season<sup>59,88,89</sup>. DGVM simulations show that the positive effects of climate change, primarily from warmer temperature<sup>14</sup>, dominate the greening trend over more than 55% of the northern high latitudes (FIG. 4b) and in the Tibetan Plateau<sup>11</sup>. However, this positive impact of anthropogenic warming on greenness appears to have weakened during the past four decades<sup>90,91</sup>, when the correlation coefficient between temperature and greenness decreased by more than 50%<sup>90,91</sup>, suggesting a possible saturation of future greening in response to warmer temperature.

In water-limited ecosystems, changes in precipitation — reflecting either decadal climate variability or trends from anthropogenic climate change — were suggested as the main driving factor of greening and browning<sup>45,92</sup>. Precipitation-driven greening is most evident in the African Sahel<sup>93,94</sup> and semi-arid ecosystems of southern Africa and Australia<sup>45,95</sup> (FIG. 4c). Both empirical models and DGVMs indicate that ‘the greening Sahel’, one of the early examples of vegetation greening detected by satellite measurements<sup>93,94</sup>, was primarily driven by increases in precipitation after a severe drought in the early 1980s<sup>96–98</sup>. This causal relationship between precipitation and greenness changes was further supported through analyses of recent microwave satellite measurements and long-term field surveys<sup>99,100</sup>.

**Land-use change.** Like climate change, land-use change exerts a considerable but highly spatially variable influence on greenness changes<sup>11,13</sup> (FIG. 4). Specifically, deforestation dominates the tropics<sup>101,102</sup>, while afforestation increases forest area over temperate regions,

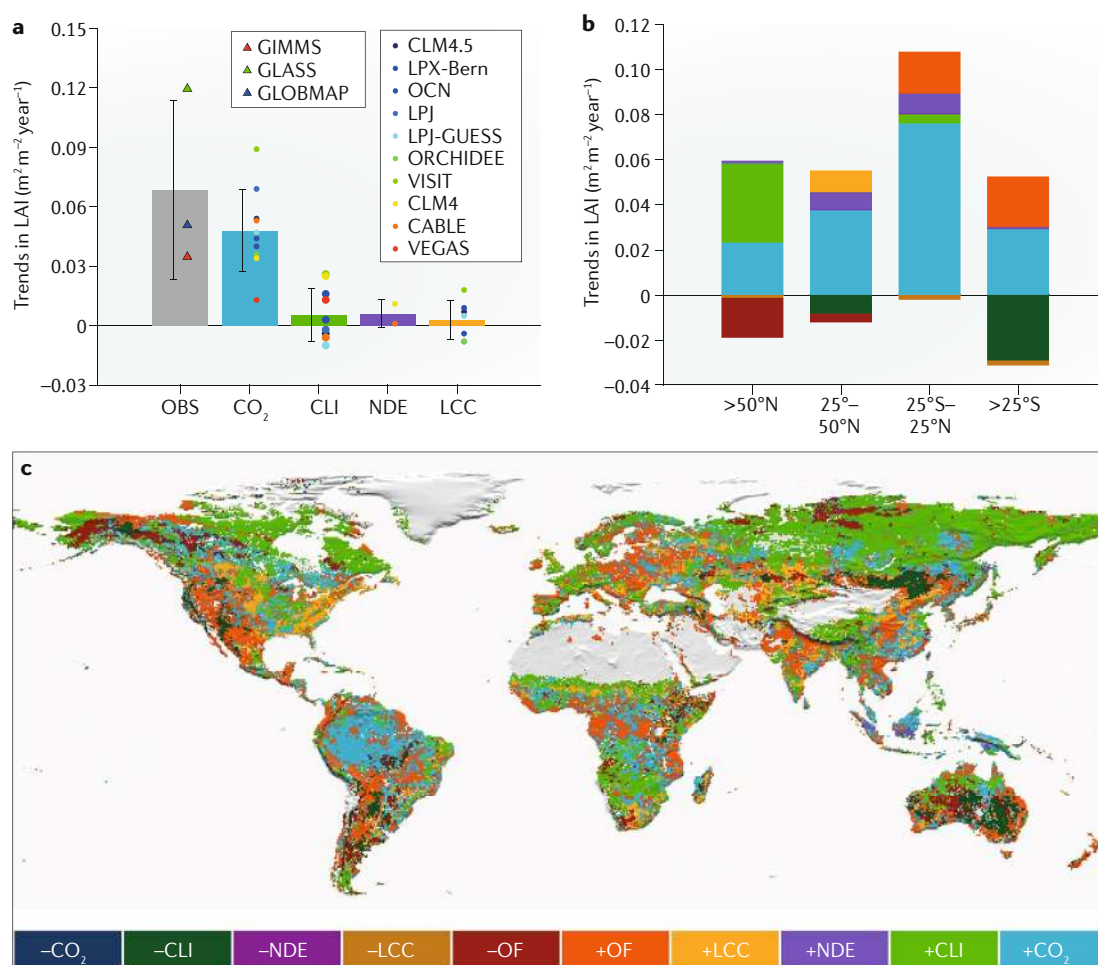
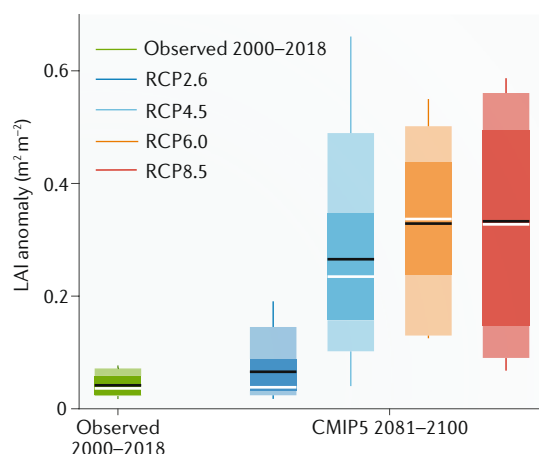


Fig. 4 | **Attribution of trends in growing season mean leaf area index.** **a** | Trends in the global-averaged leaf area index (LAI) derived from satellite observation (OBS) and attributed respectively to rising CO<sub>2</sub> (CO<sub>2</sub>), climate change (CLI), nitrogen deposition (NDE) and land cover change (LCC) during 1982–2009 (REF.<sup>11</sup>). The error bars show the standard deviation of trends derived from satellite data and model simulations. Individual model-estimated contributions of each driver to LAI trends are shown as coloured dots. **b** | Contribution of different drivers to LAI change in latitude bands (>50°N, 25°–50°N, 25°S–25°N, >25°S). **c** | Spatial distribution of the dominant driver of growing season mean LAI trend, defined as the driver that contributes the most to the increase (or decrease) in LAI in each vegetated grid cell. Other factors (OF) is defined by the fraction of the observed LAI trends not accounted for by modelled factors. Parts **b** and **c** share the same colour legend, where the '+' prefix indicates a positive effect from the corresponding driver on LAI trends and the '-' prefix indicates a negative effect. Data courtesy of Zhu et al.<sup>11</sup>. Part **c** adapted from REF.<sup>11</sup>, Springer Nature Limited.

particularly in China, where the forest area has increased by more than 20% since the 1980s<sup>103</sup>. The TRENDY ensemble of DGVMs<sup>70</sup> indicates that greenness changes over 19% of the northern temperate vegetation (25–50°N) are primarily driven by land-use change<sup>11</sup> (FIG. 4c). However, this might be an underestimate since critical land-use processes<sup>104,105</sup> are under-represented or missing in the current generation of DGVMs. For example, forest-age dynamics are not represented in most DGVMs, even though one-third of the global forests are younger than 20 years old<sup>106</sup>, implying that forest regrowth might contribute to global greening in the future. In addition, agricultural intensification with multiple cropping, irrigation and fertilizer usage must contribute considerably to vegetation greening, which is exemplified by the dominance of other unmodelled factors over agricultural lands of India, China and Eastern Europe (FIG. 4c).

**Nitrogen deposition.** Anthropogenic changes in the amount, rate and distribution of nitrogen deposition can impact greening patterns, since insufficient nitrogen availability can stunt plant growth<sup>107–109</sup>, potentially slowing greening or causing browning, but excess nitrogen can enhance plant growth in nitrogen-limited systems<sup>109</sup>. However, the few DGVMs that include the nitrogen cycle do not indicate that nitrogen deposition plays a dominant driving role on the greening at either the global or regional scales (FIG. 4). Modelling studies differ on the contribution of increasing nitrogen deposition to the global LAI increase<sup>11</sup> (9 ± 12%), largely due to the incomplete representation of nitrogen-related processes<sup>110</sup>. A growing number of DGVMs are currently incorporating nitrogen processes<sup>111</sup>, though, and future research priorities include better measurement and representation of processes such as plant nitrogen uptake and allocation<sup>110</sup>.



**Fig. 5 | Current and predicted global leaf area index.** Current (observed 2000–2018) leaf area index (LAI) anomaly ( $\text{m}^2 \text{m}^{-2}$ ) from an average of satellite measurements based on GIMMS<sup>13</sup>, GLASS<sup>192</sup>, GLOBMAP<sup>23</sup> and Moderate Resolution Imaging Spectroradiometer (MODIS) C6 (REF.<sup>193</sup>). Predicted LAI anomalies from the Coupled Model Intercomparison Project Phase 5 (CMIP5) multi-model (Supplementary Table S5) projections during 2081–2100. The boxes and whiskers indicate the minimum, 10th, 25th, 50th, 75th and 90th percentiles and the maximum LAI of CMIP5 models; the black and white lines indicate the mean and median LAI of CMIP5 models, respectively. LAI anomalies were calculated against the average during 1980–2005.

### Impact of greening on the carbon cycle

Greening increases the amount of photosynthetically active sunlight that is absorbed by vegetation and, thus, enhances productivity<sup>112,113</sup>. There has been substantial evidence showing enhanced vegetation productivity from contiguous solar-induced fluorescence (CSIF; FIG. 1f) observations<sup>114</sup>, empirical models of vegetation productivity<sup>92,115</sup> and DGVM and ESM simulations<sup>70,116</sup> (FIG. 6). It should be noted, though, that CSIF is not fully independent from MODIS greenness indices, since its derivation relies on both solar-induced fluorescence measurements from Orbiting Carbon Observatory 2 and MODIS reflectance measurements<sup>114</sup>.

Enhanced vegetation productivity increases terrestrial carbon storage, slowing down anthropogenic climate warming<sup>117</sup>. For example, about 29% of anthropogenic  $\text{CO}_2$  emissions since the 1980s have been offset by the land carbon sink ( $2.5 \pm 1.0 \text{ PgC year}^{-1}$ )<sup>111</sup>. This vegetation-induced large land carbon sink was also inferred from forest inventories<sup>118</sup> and above-ground biomass estimated from the vegetation optical depth (FIG. 1e), a microwave-based satellite measurement of both woody and leaf biomass<sup>119</sup>. Multiple lines of evidence, including analyses from DGVMs, atmospheric inversion models and the residual land sink (the mass balance residual of anthropogenic  $\text{CO}_2$  emissions, atmospheric  $\text{CO}_2$  growth rate and ocean  $\text{CO}_2$  budget), confirm the increasing magnitude of the global land carbon sink since the 1980s<sup>111</sup> (FIG. 6). An ecosystem model driven by satellite LAI measurements estimated that increased LAI accounts for 36% ( $0.4 \text{ PgC year}^{-1}$ ) of the land carbon sink enhancement of 1981–2016 (REF.<sup>112</sup>). Recent studies

indicate that the trend in the land carbon sink has further accelerated since the late 1990s<sup>120,121</sup>. For example, the rate of update during 1998–2012 was three times that of 1980–1988 ( $0.17 \text{ PgC year}^{-2}$  in comparison with  $0.05 \text{ PgC year}^{-2}$ )<sup>121</sup>, attributed to afforestation-induced greening in the temperate Northern Hemisphere<sup>13,121</sup>. These hotspots of afforestation and forest regrowth are in accordance with the greening pattern observed since 2000 by MODIS (FIG. 2c). Recent DGVM studies<sup>122,123</sup> have further confirmed that the carbon sink during the 2000s was partly driven by afforestation and forest regrowth in East Asia and Europe<sup>124</sup>. The extensive greening over croplands, however, has probably contributed less to the carbon sink, because only a minor portion of assimilated carbon by crops remain sequestered due to crop harvest.

The impact of greening on the carbon cycle is also partly responsible for the increasing seasonality of atmospheric  $\text{CO}_2$  in the northern high latitudes<sup>125</sup>. The amplitude of the Northern Hemisphere  $\text{CO}_2$  seasonal cycle has increased by as much as 50% for latitudes north of  $45^\circ \text{N}$ <sup>126,127</sup> since the 1960s, indicating enhanced vegetation productivity in northern ecosystems during the carbon-uptake period<sup>128</sup>. The spring zero-crossing date — the time when the detrended seasonal  $\text{CO}_2$  variations down-cross the zero line in spring — is a phenological indicator of the timing of early season net carbon uptake<sup>125,129</sup>. From 1987 to 2009, the spring zero-crossing date has advanced at high-latitude stations<sup>130</sup> (from  $-0.5 \text{ days decade}^{-1}$  to  $-1.8 \text{ days decade}^{-1}$ ) (FIG. 3d), a trend that is consistent with the advancing SOS (FIG. 3b). At the end of the net carbon-uptake period, the autumn zero-crossing dates of detrended seasonal  $\text{CO}_2$  variations — the time when the detrended seasonal  $\text{CO}_2$  variations up-cross the zero line in autumn — have also advanced over eight of the ten Northern Hemisphere stations studied<sup>131</sup>. The observed autumn zero-crossing date advancement (FIG. 3d) is in contrast to the delayed EOS (FIG. 3a) in autumn. This divergence in the autumnal  $\text{CO}_2$  and greenness trends suggests that, unlike in spring, autumn vegetation greening does not lead to an increased carbon sink because respiration is increasing more quickly than photosynthesis in autumn<sup>131</sup>. Visual observations (for example, from the Pan European Phenology Project PEP725) and cameras (for example, PhenoCam datasets) are providing an increasing amount of ground-based phenological evidence of this process. In the future, these data can be paired with eddy covariance flux data, to further our mechanistic understanding of the climate-change-induced seasonal change in greenness and carbon balance.

### Biogeophysical impacts of greening

Greening has discernable impacts on the hydrologic cycle and climate through modifying surface biogeophysical properties (for example, albedo, evapotranspiration (ET) and surface roughness) on local to regional and global scales<sup>19,132</sup> (FIG. 7). Vegetation's biogeophysical feedbacks to climate are, thus, critical to understanding the potential of ecosystem management, such as afforestation, for climate change mitigation<sup>3,132,133</sup>. In this section, we present the feedbacks of vegetation

#### Evapotranspiration

The flux of water emitted from the Earth's surface to the atmosphere. It is the sum of evaporation by the soil, wet canopy, open-water surfaces and transpiration by plant stomata.



# Transpiration

The loss of water from plants to the atmosphere.

greening on the hydrologic cycle and land-surface air temperature.

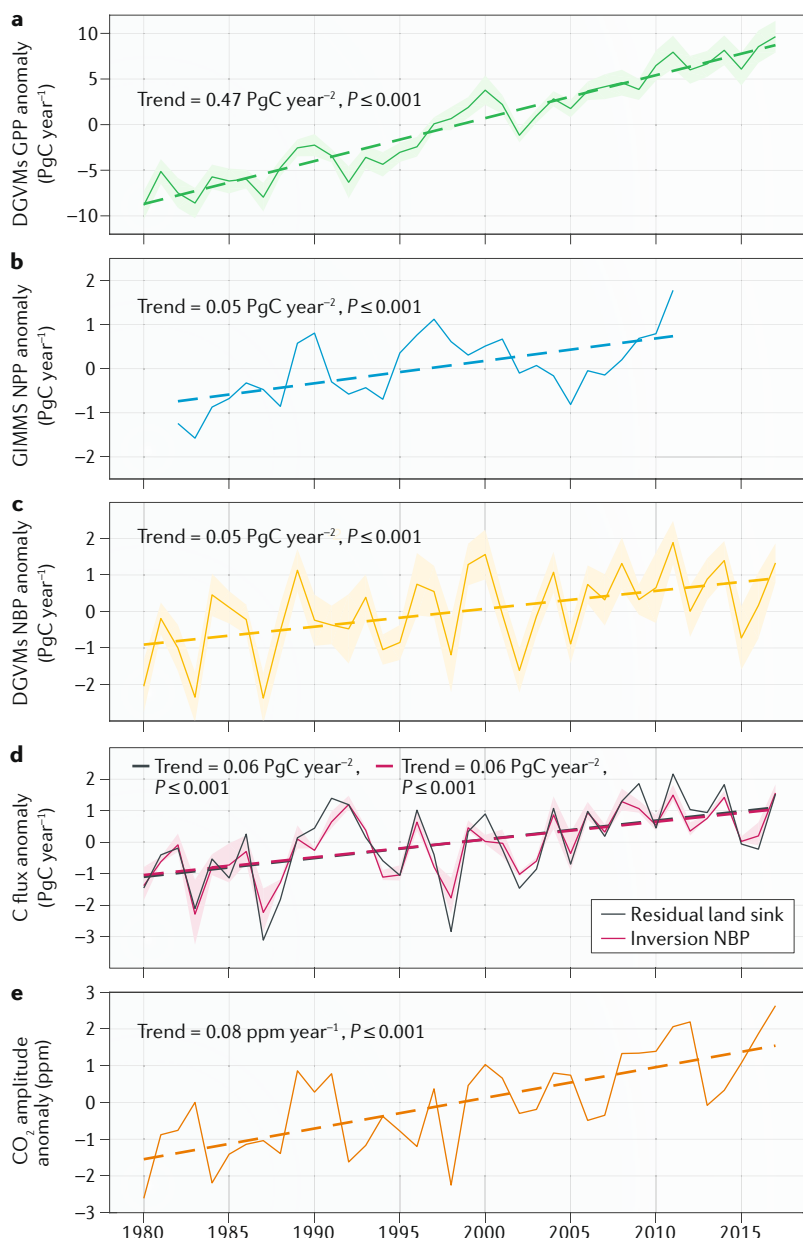
**The hydrologic cycle.** Vegetation greening modulates water cycling. Land water losses to the atmosphere occur through ET, which includes transpiration (60–90% of the total land ET<sup>134–136</sup>) and evaporation. Greening increases water losses through an extended area of leaves performing transpiration<sup>137</sup>. A larger foliage area

reduces the bare ground surface from which soil evaporation occurs, but increases the re-evaporation of rainfall intercepted by leaves<sup>138</sup>, so that greening can cause the net evaporation to either increase or decrease. Various remote-sensing-based ET estimates consistently point to a significant increase in global terrestrial ET over the past four decades, suggesting an intensified water exchange between the land and the atmosphere concurrent with the greening trend<sup>139</sup>. More than half of the global ET increase since the 1980s has been attributed to vegetation greening<sup>138,139</sup> (FIG. 7).

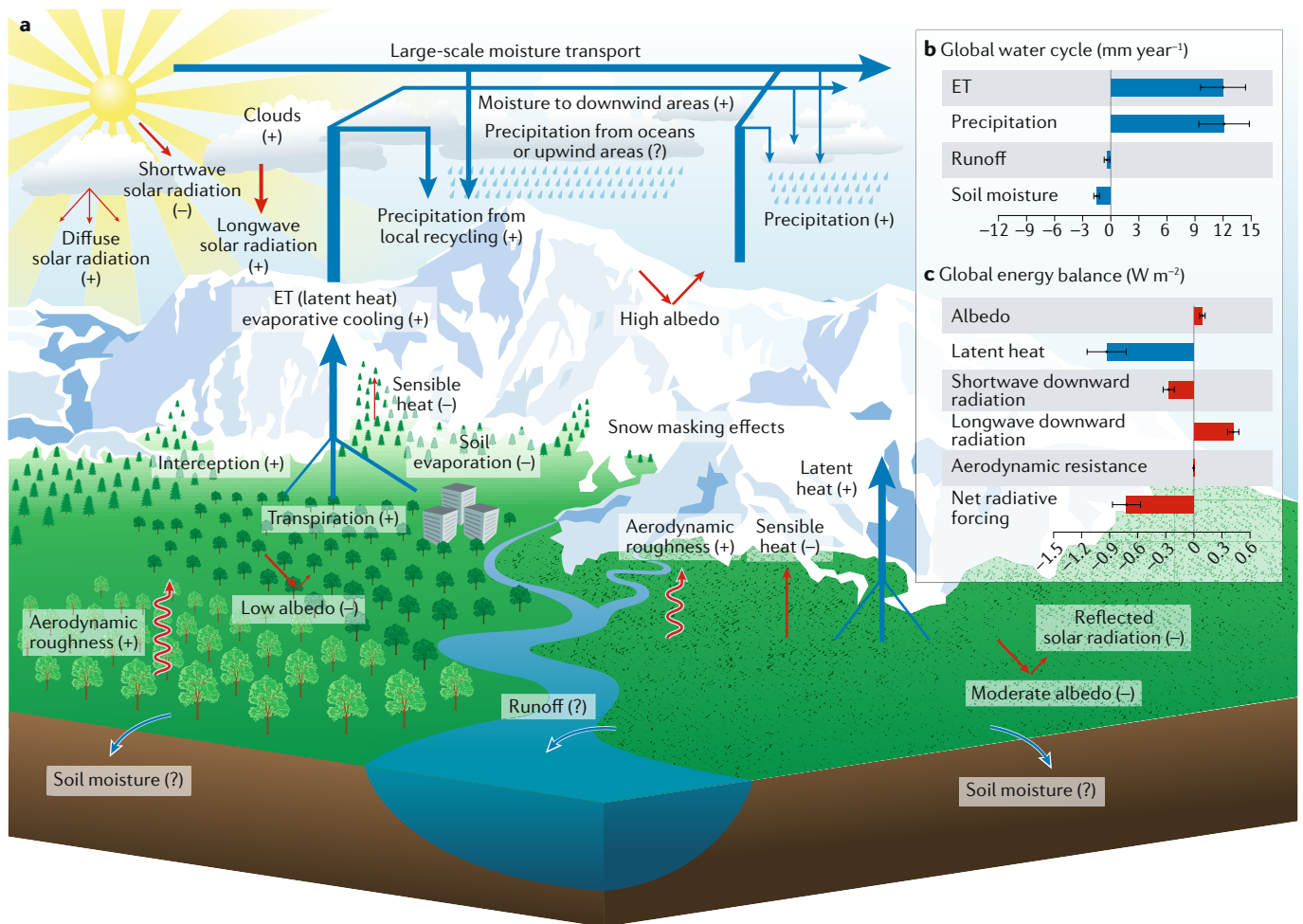
By controlling the changes in ET, vegetation greening also alters the water distribution between regions and water pools (for example, water in soil, rivers and the atmosphere). Assuming that precipitation does not change in response to vegetation greening, a greening-induced ET increase will reduce soil moisture and runoff, which can intensify droughts at the catchment scale<sup>140,141</sup>. In China's Loess Plateau for instance, where intensive afforestation is associated with a pronounced local greening, the river discharge has indeed decreased by a rate of 0.25 km<sup>3</sup> year<sup>-2</sup> over the past six decades<sup>142</sup>. However, when using ESMs that consider both the greening-induced ET increase and consequent changes in precipitation, simulations forced only with satellite-observed LAI trends do not generate dramatic changes in soil moisture or runoff at continental or global scales<sup>143,144</sup>. This is because greening-induced ET enhancement increases atmospheric water vapour content, which, in turn, promotes downwind precipitation<sup>145,146</sup>. The enhanced precipitation over transpiring regions is particularly evident in moist forests<sup>147</sup> like the Amazon or Congo, which are 'closed' atmospheric systems where 80% of the rainfall originates from upwind ET<sup>145</sup>. Such an efficient atmospheric water recycling mitigates water loss from the soil, sustains inland vegetation and maintains mesic and humid ecosystems.

In addition to intensifying water cycling at the annual scale, vegetation greening also induces seasonal hydrologic changes. There is emerging evidence that spring-greening-enhanced ET leads to a reduction in soil moisture content, which carries over into the following summer and likely suppresses vegetation growth and increases the risk of heatwaves<sup>148,149</sup>. The greening-induced water loss through ET is recycled as land precipitation in subsequent months, benefitting some remote regions through modulating large-scale atmospheric circulation patterns, despite often being insufficient to compensate for evaporative water loss locally<sup>149</sup>. Proposed climate-mitigation strategies, such as afforestation, therefore need to fully consider coupling between vegetation and other components of the Earth system.

**Land-surface air temperatures.** Greening impacts the exchange of energy between the land and the atmosphere, which ultimately leads to modifications in surface air temperature<sup>150</sup>. Greening increases ET, which cools the surface through evaporative cooling<sup>19,150</sup>, but greener canopies have a lower albedo than bare ground and absorb more sunlight, which can result in a larger sensible heat flux. This enhanced sensible heat warms the land surface, an effect called albedo warming<sup>151</sup>.



**Fig. 6 | Changes in global carbon fluxes and seasonal CO<sub>2</sub> amplitude.** Graphs depict changes in Barrow, AK, USA, since 1980. **a** | Global gross primary production (GPP). **b** | Net primary production. **c** | Net biome production (NBP). **d** | Residual land sink. **e** | Seasonal CO<sub>2</sub> amplitude. The GPP is from the ensemble mean of 16 dynamic global vegetation models (DGVMS)<sup>111</sup>. The NPP is from greenness-based modelling by Smith et al.<sup>198</sup>. The NBP is from the ensemble mean of 16 DGVMS and two atmospheric inversions<sup>111</sup>. Residual land sink is the mass balance residual of anthropogenic CO<sub>2</sub> emissions, the atmospheric CO<sub>2</sub> growth rate and the ocean CO<sub>2</sub> budget<sup>111</sup>. The shaded areas indicate the standard deviation of the GPP, NPP or NBP across models. The dashed lines indicate linear trends.



**Fig. 7 | Biogeophysical feedbacks of recent vegetation greening to the climate system. a** | Schematic diagram summarizing land-surface and atmospheric processes through which changes in vegetation greenness feed back into the climate system. For each process or flux, the corresponding symbols ‘–’, ‘+’ and ‘?’ in brackets represent an increasing, decreasing and unknown trend, respectively, in response to vegetation greening, and the colour of arrows represents impacts on water (blue) or energy balance (red, except the latent heat in blue). **b** | Summary of greening-induced changes in major global water cycle fluxes in mm year<sup>-1</sup> from 1982 to 2011. Data courtesy of Zeng et al.<sup>19</sup>. **c** | Summary of greening-induced changes in global surface energy balance in W m<sup>-2</sup> from 1982 to 2011. Data courtesy of Zeng et al.<sup>144</sup>. The error bars show the standard error of the estimates. The bar colours are the same as the corresponding fluxes shown in part **a**. ET, evapotranspiration.

The net effect of greening on surface air temperature in many cases can be viewed as the balance between evaporative cooling and albedo warming<sup>152,153</sup>, which was estimated globally to be  $-0.9 \text{ W m}^{-2}$  from evaporative cooling and  $+0.1 \text{ W m}^{-2}$  from albedo warming<sup>19</sup> (FIG. 7c).

Greening can also trigger a series of changes through atmospheric circulation that indirectly affect the surface temperature<sup>154</sup>. For example, the additionally transpired water enhances atmospheric water vapour content, which results in more longwave solar radiation entrapment and re-emission in the atmosphere, but reduces the amount of shortwave solar radiation reaching the Earth’s surface through increased cloud formation<sup>19,155,156</sup> (FIG. 7). When all the aforementioned impacts of vegetation greening on near-surface air temperature were simulated in coupled ESMs driven by the satellite-based greening since the 1980s, the results suggested a net cooling trend by  $12\% \pm 3\%$  of the concurrent observed warming rate<sup>19</sup>.

In warm regions such as the tropics and subtropics, evaporative cooling effects are generally larger than albedo warming effects, leading to a net cooling effect when vegetation greenness increases<sup>19,157,158</sup>. However, the net effect of greening on surface air temperature over the Northern Hemisphere extratropical regions is still subject to debate. Studies based on idealized afforestation and/or deforestation experiments<sup>1,159</sup> or comparisons of the energy budget differences between paired forest and short vegetation sites<sup>132,153</sup> suggested that the albedo warming effect plays a dominant role. These studies, though, assumed complete land cover changes, whereas greening can be gradual. By integrating satellite observations with ESMs, several studies provided an alternative approach that more realistically simulated the effects of vegetation greenness changes and isolated the signal of climate response to greening. These studies found that greening slowed down warming through

evaporative cooling in Arctic and boreal regions<sup>19</sup>, the Tibetan Plateau<sup>160</sup> and temperate regions like East Asia<sup>161</sup>. Nonetheless, current state-of-the-art modelling efforts are still inconclusive, as some processes are not yet well represented in ESMs, such as snow masking by greener canopies during cold seasons<sup>162–164</sup> and the partitioning of transpiration and evaporation that is sensitive to vegetation greenness change<sup>136</sup>. Since most ESMs underestimate the ratio of transpiration to ET<sup>136</sup>, evaporative cooling by greening could have been underestimated<sup>19,133</sup>.

## Conclusions

Widespread vegetation greening since the 1980s is one of the most notable characteristics of biosphere change in the Anthropocene. Greening has significantly enhanced the land carbon sink, intensified the hydrologic cycle and cooled the land surface at the global scale. A mechanistic understanding of the underlying drivers shows how anthropogenic forcing has fundamentally altered today's Earth system through a set of feedback loops. Improved knowledge of greenness changes, together with recent progress in observing technology and modelling capacity, has resulted in major advances in understanding global vegetation dynamics. Nonetheless, we still face many challenges ahead.

One key challenge is to continue developing the capacity of remote sensing to measure vegetation structure and functions. Although the vegetation greenness indices described in this Review have proved highly reliable, contemporary satellite greenness products still suffer from limitations, such as inadequate sensitivity to detect changes in dense vegetation, aliasing between snow cover decrease and leaf area increase in cold ecosystems (such as boreal forests), atmospheric contamination, orbital drift and sensor replacements. Compared with the AVHRR, the new moderate-resolution spectral bands and spatial resolutions of 250 m to 1 km of the MODIS sensors on board the Terra (operating since 1999) and Aqua (operating since 2002) satellites have provided global datasets that largely improved the long-term monitoring of vegetation greenness<sup>13</sup>. The current scientific community needs to include Earth observations with higher temporal, richer spectral and finer spatial resolutions to capture various ecosystem functions and processes responding to different parts of the

electromagnetic spectrum<sup>165</sup>. We expect the development of next-generation satellite missions and vegetation indices to better fulfil these needs. For example, ongoing efforts on developing hyperspectral remote sensing such as the EnMAP, FLEX and HypsIRI missions will improve the richness and specificity of spectral information on vegetation structure and functioning.

Another equally important challenge is to validate satellite-based greenness changes with ground observations. Currently, the lack of systematic long-term ground observations covering a large spatial gradient from the high Arctic to the tropics has led to few available ground truths<sup>166</sup> to confirm greenness changes detected through satellite products. Therefore, expanding existing observational networks (such as PhenoCam and FLUXNET) is a high priority. For example, the mismatch between the spatial distribution of vegetation productivity and the density of FLUXNET sites<sup>167</sup> highlights the need to expand the current network from the mid-latitudes to the tropics, where the most photosynthesis takes place. Also, growing crowd-sourced observations by citizen scientists, such as the CrowdCurio phenology observations over the eastern USA<sup>168</sup>, can provide valuable data that complement the more expensive professional ground observation networks. These increasing types and amounts of data, together with the rapid development of deep learning<sup>169</sup> and process modelling<sup>11</sup>, offer promising tools for improving our understanding of vegetation greening<sup>169</sup>.

Considerable uncertainties remain in ESM projections on if and where vegetation greening will occur. Recent studies have identified several processes causing vegetation browning in some regions, including forest diebacks<sup>170</sup>, insect<sup>35</sup> and disease outbreaks<sup>171</sup>, thermokarst development<sup>172</sup>, human mismanagement<sup>36,173</sup>, destructive logging<sup>174</sup> and industrial development<sup>175</sup>. These emerging threats could lead to unexpected changes in vegetation greenness relative to our current projections (such as the projections shown in FIGS 2e–h,5), since these processes are under-represented in ESMs. Thus, integrating continued space and ground monitoring and advancing ESM developments is a critical cross-sectoral research priority.

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## Author contributions

S.P., X.W., T.P., C.C., X.L., Y.H., J.W.B., A.C., P.C., H.T. and R.B.M. wrote the first draft of the manuscript. S.P., X.W. and R.B.M. reviewed and edited the manuscript before submission. All authors made substantial contributions to the discussion of content.

## Competing interests

The authors declare no competing interests.

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## Supplementary information

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