Projecting climate change and land-use scenarios of crop pollination risk

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Abstract

Recent studies have highlighted rapid ongoing changes in insect biodiversity, in part driven by the interaction of land-use and climate change. Insect pollinator biodiversity specifically has been shown to be undergoing rapid changes, with potential consequences for the provision of crop pollination. Here, we present a global assessment of the interactive effects of climate change and land use on pollinator abundance, and predictions of the risk to crop pollination worldwide from the inferred changes in pollinator abundance. We show that the combination of agricultural land use and recent climate change is associated with particularly large reductions in the abundance of insect pollinators. We further show that the tropics will likely experience the greatest risk of future pollination shortfalls, putting at risk the production of crops that depend on insect pollination. The risk to these crops is highest, and predicted to increase most rapidly, in regions of sub-Saharan Africa and northern South America, especially in areas used to produce cocoa, mango, pumpkin, melon, watermelon, and coffee. The health, well-being, and livelihoods of billions of people to some extent depends upon the availability and affordability of crops dependent on animal pollination. Climate change and agricultural land use could put these people's wellbeing at high risk.

One sentence summary

Climate change associated crop pollination risk is highest, and predicted to increase most rapidly, in regions of sub-Saharan Africa and northern South America, especially in areas used to produce cocoa, mango, pumpkin, melon, watermelon, and coffee.

Introduction

Recent studies have highlighted rapid ongoing changes in insect biodiversity (*1–3*). Some of these studies have reported net declines (*1, 3*), at least for terrestrial insects (*1*), while others have shown mixtures of

gains and losses (*2*). Pollinator biodiversity specifically has been shown to be undergoing rapid changes (*4–8*), with potential consequences for the provision of pollination to wild plants and crops. Importantly, evidence for insect biodiversity trends has been biased toward Western Europe and North America, with little coverage of tropical regions (*9, 10*). Although a few studies have shown steep declines of insects in the tropics (*3*), evidence about insect biodiversity trends there often remains anecdotal (*11*), with global syntheses (*1, 2*) having strong geographic biases towards non-tropical regions.

Among the drivers of insect and pollinator biodiversity changes, human land-use disturbance and climate change are prominent (*5, 10*). A large number of studies have shown substantial effects of human land use and agricultural intensification on insects, including pollinators (*12–15*). Climate change is emerging as an increasingly important driver of changes in insect biodiversity (*8, 12, 16–19*). Tropical insects are expected to be more susceptible to climate change, owing to their narrower physiological tolerance compared to non-tropical species (*20*), but evidence about the response of tropical insects to climate change remains very scarce (*19*). Evidence is now accumulating for strong interactive effects of land-use change and climate change on insects (*21–25*). A key mechanism underpinning this interactive effect is the altered microclimatic conditions in areas where vegetation has been modified for human land use (*21*). The interaction between the effects of climate change and land use is often synergistic, leading to greater declines in biodiversity than if the pressures acted alone (*21, 23–25*), and larger declines in tropical than non-tropical insect biodiversity (*25*).

Changes in the biodiversity and composition of pollinator communities will have large effects on the provision of pollination services. Pollen limitation from animal pollinator losses has already been shown to reduce the reproductive success of wild plants (*4, 26*), and the productivity of certain crops (*27–30*), although there is no evidence yet that pollen limitation is causing wholesale reductions in yields of crops that rely on animal pollination (*31*). Evidence that insect biodiversity responds to human pressures more strongly in the tropics than elsewhere (*15, 25*) is noteworthy, given that the majority of animal-pollination-dependent crops are grown there (*32*). However, it is not only tropical countries that will experience the effects of pollinator losses and subsequent pollen limitation, with highly developed countries benefiting from imports of animal-pollination-dependent foods from tropical areas (*33*). Abundance, species diversity and functional diversity of pollinators have all been implicated as determinants of the delivery of pollination service (*30, 34–38*). Previous attempts to model the provision of crop pollination service have been based on predictions of pollinator abundance, which bears a direct relation to pollination of plants, and has been shown to give a reasonable approximation of pollen deposition in at least some study systems (*39*).

Here, we present a global assessment of the interactive effects of climate change and land use on pollinator abundance, and predictions of the risk to crop pollination worldwide from the inferred changes in pollinator abundance (see Shiny app for full extended predictions, https://joemillard.shinyapps.io/pollinator_dependence_visualisation/). Our analyses are based on the PREDICTS database of biodiversity comparisons in different land uses (*40*), together with a list of species identified in the literature as likely pollinators (*15*). We use mixed-effects models to fit total pollinator community abundance as a function of land use (primary vegetation versus cropland) and a measure of historical climate change between a baseline period (XXXX-XXXX) and the X years prior to biodiversity sampling, standardized by monthly climate variability in the baseline period (*25*). Using estimated proportional reduction of pollinator total abundance as a proxy for relative risk to pollination services, together with estimates of where crops are grown (*41*), how dependent these crops are on animal pollination (*32*), projections of historical and future climate change, and a set assumptions for the relationship between local pollinator abundance and crop pollination, we predict which locations and crops are likely to be at greatest risk from pollination shortfalls. We then combine these projections with estimates of the trade in pollination dependent production, to predict regions of the world that may be vulnerable to the indirect consequences of crop pollination risk via trade routes.

Results and Discussion

The abundance of insect pollinators responded strongly to the interaction of recent climate change and land use (Figure 1). Within cropland experiencing novel temperatures (Standardized Temperature Anomaly =

1), pollinator abundance is 38.9% of natural habitat that has not experienced temperature increases. Our results are qualitatively consistent with recent results for a sample of all insects (*25*), but importantly we show that responses to the interactive effects of climate change and land use are stronger for pollinating than non-pollinating insects (Figure 1). Whether the sensitivity of pollinating insects to the interaction of climate change and land use relates directly to their reliance on floral resources, or to other correlated traits such as dispersal ability, body size, voltinism, or specialism, is unclear. Most likely some combination of both is true. For example, insect-pollinated plants have been shown to respond more strongly to warming than wind-pollinated plants, suggesting that plant-pollinator interactions are highly sensitive to thermal changes (Fitter & Fitter, 2002). Pollinator pilosity (i.e. hairiness), on the other hand, has been shown to affect an insect's ability to adapt to changes in climate (Roquer-Beni et al., 2020), and given its nature as a trait typical of bees and hoverflies, tends to be correlated with a reliance on floral resources (Stavert et al., 2016).
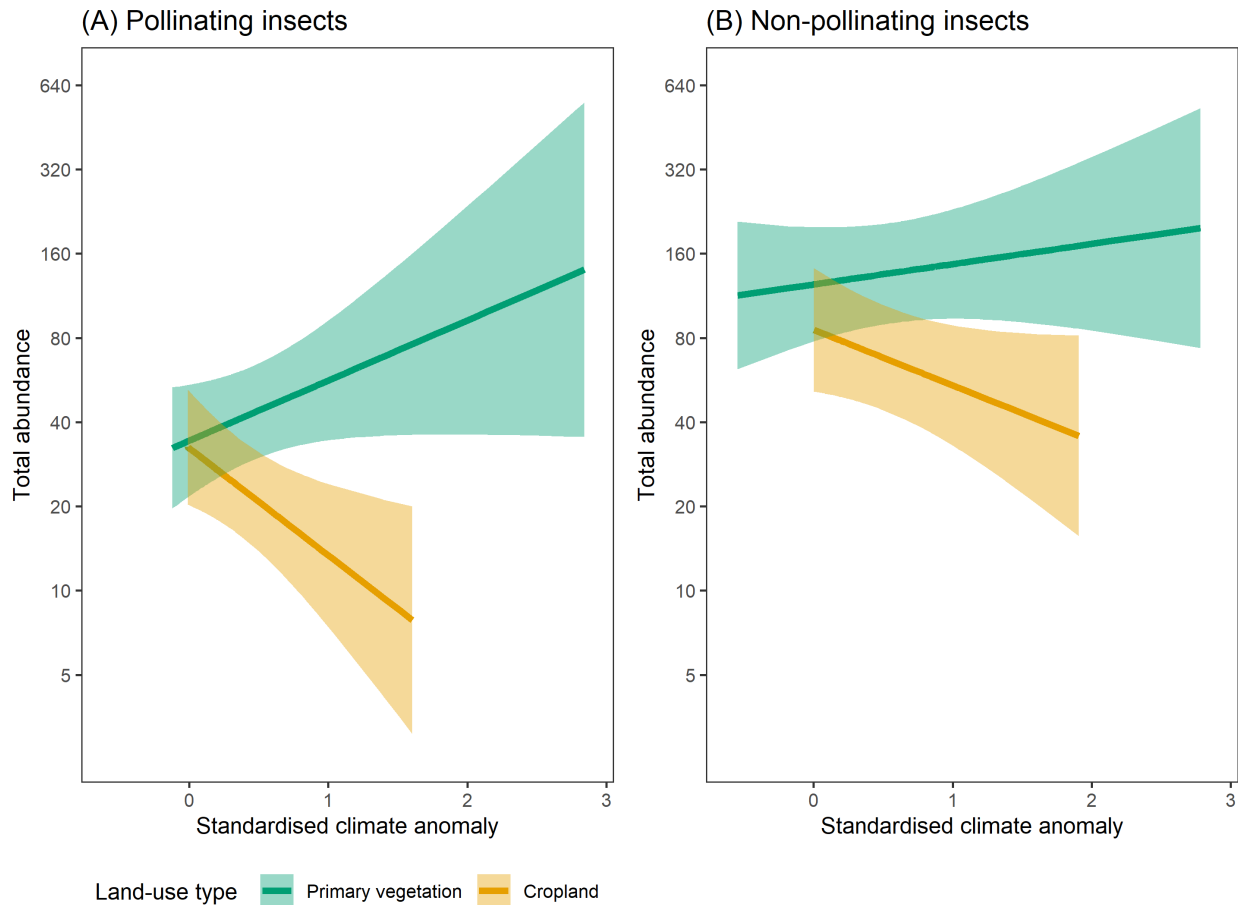

Figure 1: Response of pollinating (A) and non-pollinating (B) insect total abundance to the interactive effect of standardised temperature anomaly and land use (primary vegetation versus cropland) (A: F = 23.3195, p < 0.001; B: F = 10.5764, p < 0.01). Note that abundance is plotted on a $\log_e$ scale. Results are based on linear mixed-effects models. Shading represents 95% confidence intervals around the mean fitted effect. Green = primary vegetation; yellow = cropland.

We predict that total pollination production risk will increase under all climate scenarios (Figure 2). For all scenarios, our projections use change in pollinating insect abundance as a proxy for the relative risk to the production of crops dependent on insects (incorporating information on where crops are grown worldwide (Earthstat ref) as well as the fractional dependence of crops on animal pollination (Klein ref)). Our estimates

of risk are based on the distribution of crops as grown in 2000. We also do not account for changes in the distribution of crops over time, which are likely to occur as a result of the direct impacts of climate change as well as indirect effects such as through the loss of pollinator biodiversity. Our estimates of both relative risk as well as absolute production risk should be interpreted as indices of risk, rather than predictions of the actual amount of production likely to be lost, given a very high uncertainty in how pollinator abundance changes will translate into actual production losses. Increases in risk are seen for all assumed relationships between abundance loss and production risk (linear and varying degrees of convexity), although the magnitude of changes in relative risk and especially absolute production risk varied widely. The predicted rate of increase in average production risk was substantially higher under RCP 6.0 than RCP 2.6, suggesting that efforts to mitigate climate change will reduce the risk to future crop production, alongside the many other benefits (refs). Relative production risk varied strongly between years under the convex abundance-production relationships. This volatility may be explained by way in which the non-linearity of abundance/production relationships interacts with inter-annual climate variability caused by the El Nino Southern Oscillation (*reference*), although at present this is unclear. While increasingly convex assumed relationships between insect abundance loss and crop production risk led to steeper increases in relative production risk with future climate change, absolute production risk was markedly lower owing to a lower baseline in the present day (Figure 2). Results were robust to variation in climate predictions under different individual climate models (Figure S06), do not change markedly when abundance loss is capped at the maximum model-fitted value (Figure S02), do not change markedly when tested for sensitivity to data quality (Figure S04), and likely hold across a set of plausible active season temperature thresholds, fitted through two different approaches (Figures S11 and S12).
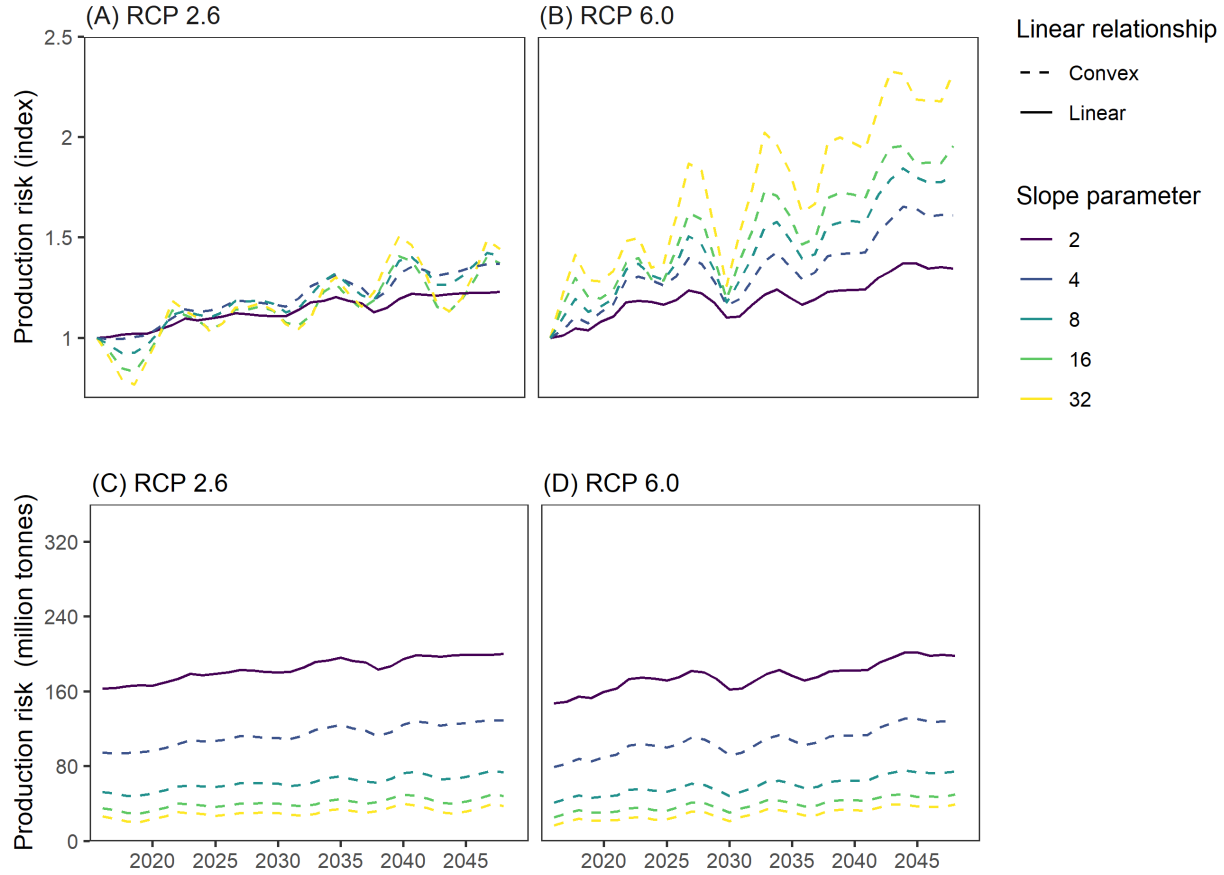
Figure 2: Projected change in total production risk under two RCP scenarios (6.0 and 2.6) and a set of different abundance-production relationships (linear and varying degrees of convexity). Results are shown both for an index of change in relative risk (A) and for absolute total production at risk (in tonnes) (B). For each year into the future, the standardised temperature anomaly was projected globally for all cells with production of crops dependent on animal pollination, using a 3-year rolling average of temperature anomaly estimates in each cell. Temperature projections are the mean across an ensemble of different climate models. We used data on crop production from the year 2000 (the latest year when such data are available for all crops). For each annual projection of standardised temperature anomaly, insect pollinator abundance on cropland was predicted according to the model shown in Figure 1A, and then expressed as proportional abundance loss compared to cropland that has experienced no warming (i.e. standardised temperature anomaly of 0). In each cell, the total production of crops (estimated according to (Earthstat ref)) was adjusted to account for dependence on animal pollination (using ratios from (Klein ref)), and then adjusted according to the predicted percentage reduction in insect pollinator abundance in that cell. These estimates of crop production at risk were summed across crops, and then across all terrestrial cells globally. Lines show different assumed relationships between insect pollinator abundance and crop production: dashed = convex relationships (of differing degrees, indicated by different colours: yellow = most convex; purple/blue = least convex); and solid = linear relationship.

Production risk from insect pollinator abundance losses is highest, and predicted to increase most rapidly, in regions of sub-Saharan Africa, northern South America, and southeast Asia (Figure 3B and Figure 3C). Somalia, Guinea Bissau, and Ivory Coast emerge as particularly vulnerable, given their high and increasing production risk, and the high dependence of their economy on agriculture (Figure 3C). A number of countries

in Asia, South America, and the Caribbean have high and increasing levels of production risk, but lower levels of agricultural dependence, making their economies less vulnerable overall. For example, the Philippines, Indonesia, Papua New Guinea, Puerto Rico, Haiti, and Suriname all have high and increasing levels of production risk (Figure 3C), but relatively low GDP adjusted pollination dependent production. Among crops, cocoa is estimated to be at highest risk by a large margin, followed by mango and watermelon (Figure 3A). Coffee is also expected to have a combination of relatively high production risk and high value, suggesting that regions in which it is grown may experience economic difficulties, unless the pollination service can be replaced cost effectively.

For both countries and crops it is difficult to predict how the consequences of our production risk measure will play out in the real world. In part this is due to the push and pull of multiple uncertainties, some of which we do explore here (e.g. the relationship between pollinator abundance and crop production), but many of which we do not or cannot (e.g. the changing distribution of crops, the economic viability of hand pollination, the buffering effects of managed pollinators, the effects of climate change alone, and other technological solutions such as the breeding or engineering of pollinator independent cultivars). Regardless, at the global scale, it's not unreasonable to state that the distribution of standardised temperature anomaly and animal-pollinator dependent crop production should predict a substantial portion of the variation in pollinator-mediated losses to crop production. For cocoa, a large quantity of literature describes the direct effects of climate change on crop production (refs), often assuming direct effects on growth and overlooking those that might be pollinator-mediated. Given the difficulties in teasing these mechanisms apart, and that the direct effects of climate change are easier to intuit and likely stronger, it is probable that this trend continues. However, importantly solutions to direct and pollinator-mediated effects will differ. For example, shade trees might protect crops from extreme temperatures, but might not for the midge pollinators on which cocoa pollination depends.
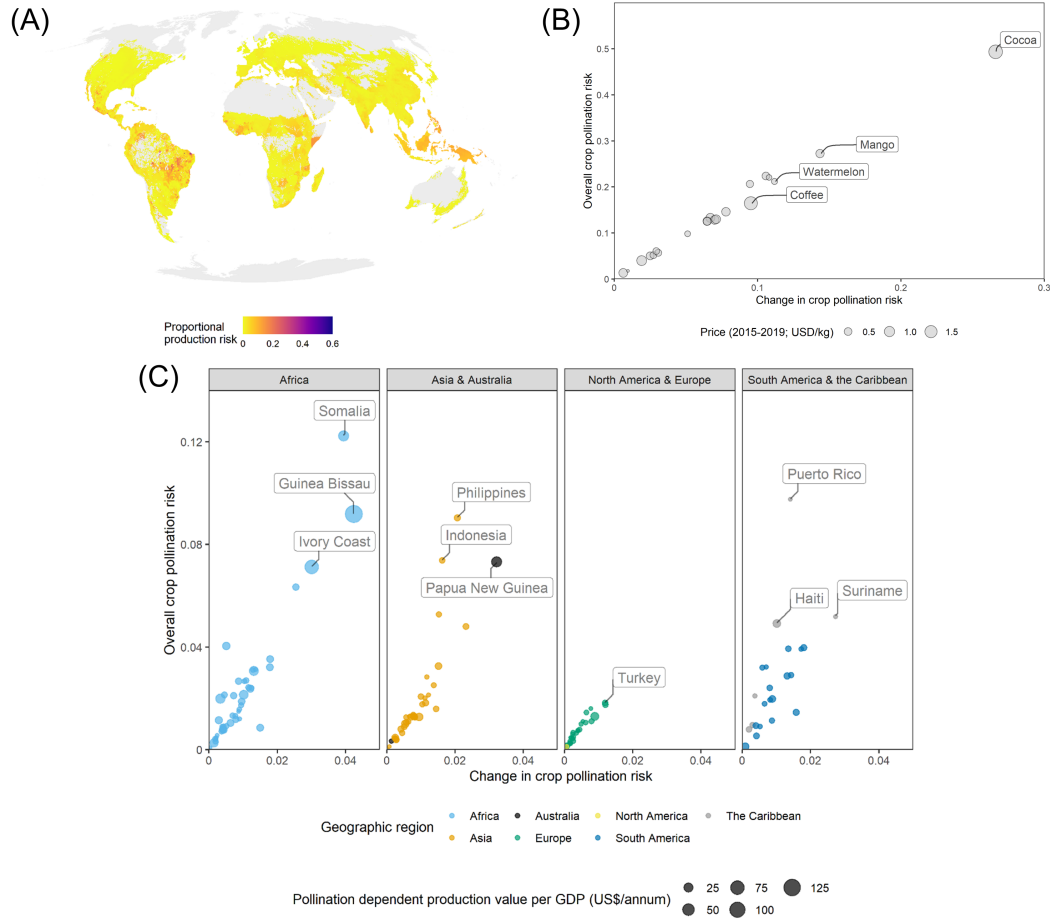
Figure 3: Projected change globally in the fraction of crop production estimated to be at risk in 2050 under the RCP 6.0 climate scenario, assuming a linear relationship between insect pollinator abundance loss and production loss for crops dependent on animal pollination. All projections are based on mean projections of the standardized temperature anomaly based on temperature estimates from an ensemble of individual climate models. (A) The sum of crop production at risk across all crops with some dependence on animal pollination (in each spatial cell), as a proportion of the production of all crops grown in that spatial cell (i.e. proportional production risk; see Figure S08 to see how cell-level risk differs among abundance-production scenarios). (B) Proportional production risk for the 20 crops with the highest total pollination dependent production globally (see Figure S01 for the top 20 crops). Overall risk here is the median of proportional production risk for all spatial cells in which that crop appears, whilst change in risk is the difference in overall risk between the start and the end of the series. Point size represents an estimation of the per kg value of each crop for the years 2015-2019, calculated from FAO data. (C) Proportional production risk at the level of each country. Here overall risk is the median of proportional production risk for all cells of that country, whilst change in risk is the difference in overall risk between the start and the end of the series. Point size here represents the total value of the pollination dependent production in that country adjusted for GDP, calculated from the product of total pollination dependent production per annum (according to Monfreda et al 2008 and Klein et al 2007) and the per kg value of each crop (according to FAO data). Colour represents the geographic region of each country, distinguishing between regions within a panel: Blue, Africa; orange, Asia; black, Australia; green, Europe; yellow, North America; dark blue, South America; grey, the Caribbean.

Countries besides those with high production risk will feel the impact of losses of pollinator biodiversity

and the crops that depend on them, through disruption to imports. Total import risk per capita tends to be highest in northern countries, particularly continental western Europe, Canada, and parts of the Middle East. Some of the countries with the highest levels of import risk are those that have a high dependence on food imports, such as the Cayman Islands, Singapore, and Aruba. Interestingly, the Netherlands has the third highest per capita import risk, consistent with its status as the greatest importer of cocoa beans worldwide (reference). Our predictions of import risk are based on trade patterns from the present day (da Silva et al 2021), meaning we do not account for changes in trade flows that will likely occur in the future.
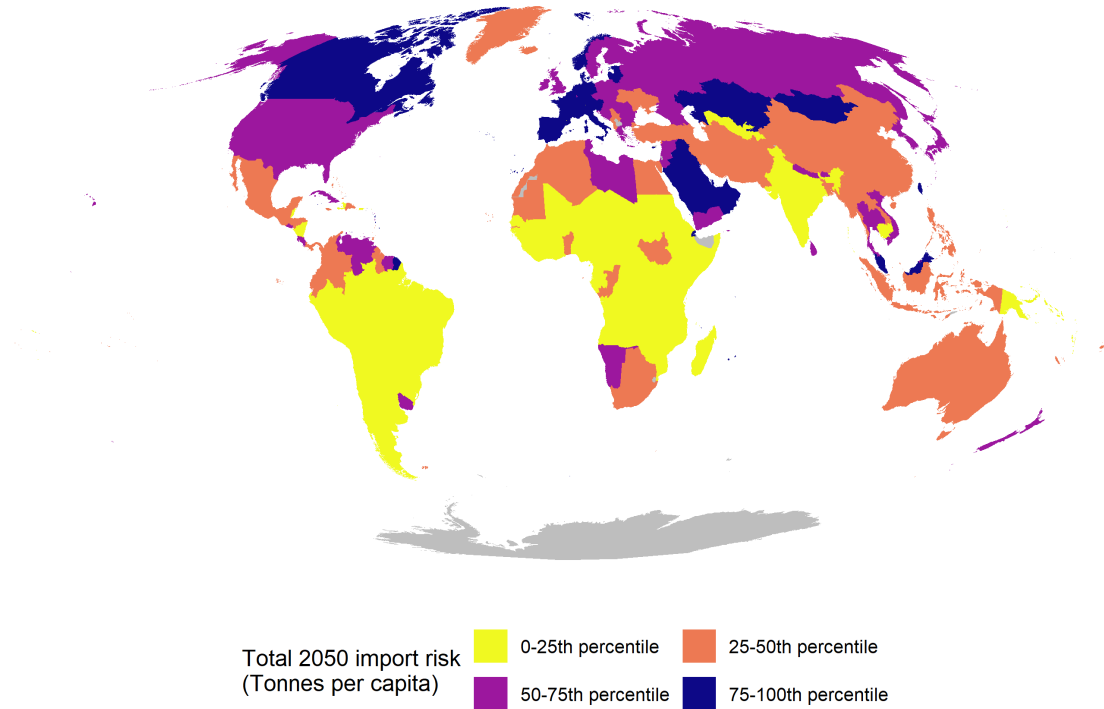

Figure 4: Projected total import risk for 2050 under the RCP 6.0 scenario, assuming a linear relationship between insect pollinator abundance loss and production loss. Projections are based on changes in the mean standardized temperature anomaly based on projections from an ensemble of individual climate models. Import risk is a measure of how the effects of localised production risk might be distributed among other countries, calculated using trade flow data from da Silva et al (2021). For each yearly time step into the future, local production risk in each spatial cell is attributed to importers according to the quantity of pollination trade imported from each country. For example, if an importer is dependent on 3 countries for imports, at a proportion of 30%, 50%, and 20%, then any change in import risk should scale as a function of the local production risk aggregated at those same proportions. Total import risk here has been adjusted for country population size and then converted to a percentile. Colours represent each percentile grouping: yellow, 0-25th percentile; orange, 25-50th; purple, 50-75th; dark blue, 75-100th.

Interacting effects of climate change and anthropogenic land use are rapidly restructuring biodiversity (*refs*). Here we show that the combination of agricultural land use and recent climate change is associated with

particularly large reductions in the abundance of insect pollinators. We further show that the tropics will likely experience the greatest risk of future pollination shortfalls, putting at risk the production of crops that depend on insect pollination. The risk to these crops is highest, and predicted to increase most rapidly, in regions of sub-Saharan Africa and northern South America, especially in areas used to produce cocoa, mango, pumpkin, melon, watermelon, and coffee. Given the many factors that determine crop production and crop price, the likely effects of insect pollinator losses on crop production are unclear (*refs*), and even if they do occur conclusive attribution is likely to be challenging (Khanal et al., 2018; Santeramo et al., 2018). Such complications likely in part explain why identifying a strong effect of pollinator losses on global crop yield and price has thus far been so difficult (Aizen et al., 2008, 2009; Garibaldi et al.,2011). Regardless, there is sufficient evidence to suggest that declining insect pollinator abundance will influence crop production risk, particularly for those in the global south (Dicks et al., 2021). Such risk could manifest in the form of direct and immediate losses to crop production through pollinator shortfall, but also through decreased resilience to changes that will happen in conjunction (e.g. the effects of extreme temperature and drought on crop growth). The health, well-being, and livelihoods of billions of people to some extent depends upon the availability and affordability of crops dependent on animal pollination (Potts et al., 2016). Climate change and agricultural land use could put these people's wellbeing at very high risk.

References

1. R. van Klink, D. E. Bowler, K. B. Gongalsky, A. B. Swengel, A. Gentile, J. M. Chase, Meta-analysis reveals declines in terrestrial but increases in freshwater insect abundances. *Science.* **368**, 417–420 (2020).

2. M. Dornelas, N. J. Gotelli, H. Shimadzu, F. Moyes, A. E. Magurran, B. J. McGill, A balance of winners and losers in the anthropocene. *Ecology Letters.* **22**, 847–854 (2019).

3. B. C. Lister, A. Garcia, Climate-driven declines in arthropod abundance restructure a rainforest food web. *Proceedings of the National Academy of Sciences.* **115**, E10397–E10406 (2018).

4. J. C. Biesmeijer, S. P. M. Roberts, M. Reemer, R. Ohlemüller, M. Edwards, T. Peeters, A. P. Schaffers, S. G. Potts, R. Kleukers, C. D. Thomas, J. Settele, W. E. Kunin, Parallel declines in pollinators and insect-pollinated plants in britain and the netherlands. *Science.* **313**, 351–354 (2006).

5. S. G. Potts, J. C. Biesmeijer, C. Kremen, P. Neumann, O. Schweiger, W. E. Kunin, Global pollinator declines: Trends, impacts and drivers. *Trends in Ecology & Evolution.* **25**, 345–353 (2010).

6. T. H. Oliver, N. J. B. Isaac, T. A. August, B. A. Woodcock, D. B. Roy, J. M. Bullock, Declining resilience of ecosystem functions under biodiversity loss. *Nature Communications.* **6**, 10122 (2015).

7. G. D. Powney, C. Carvell, M. Edwards, R. K. A. Morris, H. E. Roy, B. A. Woodcock, N. J. B. Isaac, Widespread losses of pollinating insects in britain. *Nature Communications.* **10**, 1018 (2019).

8. P. Soroye, T. Newbold, J. Kerr, Climate change contributes to widespread declines among bumble bees across continents. *Science.* **367**, 685–688 (2020).

9. Y. Basset, G. P. A. Lamarre, Toward a world that values insects. *Science.* **364**, 1230–1231 (2019).

10. D. L. Wagner, Insect declines in the anthropocene. *Annual Review of Entomology.* **65**, 457–480 (2020).

11. D. H. Janzen, W. Hallwachs, Perspective: Where might be many tropical insects? *Biological Conservation.* **233**, 102–108 (2019).

12. M. S. Warren, J. K. Hill, J. A. Thomas, J. Asher, R. Fox, B. Huntley, D. B. Roy, M. G. Telfer, S. Jeffcoate, P. Harding, G. Jeffcoate, S. G. Willis, J. N. Greatorex-Davies, D. Moss, C. D. Thomas, Rapid responses of british butterflies to opposing forces of climate and habitat change. *Nature.* **414**, 65–69 (2001).

13. A. De Palma, S. Abrahamczyk, M. A. Aizen, M. Albrecht, Y. Basset, A. Bates, R. J. Blake, C. Boutin, R. Bugter, S. Connop, L. Cruz-López, S. A. Cunningham, B. Darvill, T. Diekötter, S. Dorn, N. Downing, M. H. Entling, N. Farwig, A. Felicioli, S. J. Fonte, R. Fowler, M. Franzén, D. Goulson, I. Grass, M. E. Hanley, S. D. Hendrix, F. Herrmann, F. Herzog, A. Holzschuh, B. Jauker, M. Kessler, M. E. Knight, A. Kruess, P. Lavelle, V. Le Féon, P. Lentini, L. A. Malone, J. Marshall, E. M. Pachón, Q. S. McFrederick, C. L. Morales, S. Mudri-Stojnic, G. Nates-Parra, S. G. Nilsson, E. Öckinger, L. Osgathorpe, A. Parra-H, C. A. Peres, A. S. Persson, T. Petanidou, K. Poveda, E. F. Power, M. Quaranta, C. Quintero, R. Rader, M. H. Richards, T. Roulston, L. Rousseau, J. P. Sadler, U. Samnegård, N. A. Schellhorn, C. Schüepp, O. Schweiger, A. H. Smith-Pardo, I. Steffan-Dewenter, J. C. Stout, R. K. Tonietto, T. Tscharntke, J. M. Tylianakis, H. A. F. Verboven, C. H. Vergara, J. Verhulst, C. Westphal, H. J. Yoon, A. Purvis, Predicting bee community responses to land-use changes: Effects of geographic and taxonomic biases. *Scientific Reports.* **6**, 31153 (2016).

14. I. Oliver, J. Dorrough, H. Doherty, N. R. Andrew, Additive and synergistic effects of land cover, land use and climate on insect biodiversity. *Landscape Ecology.* **31**, 2415–2431 (2016).

15. J. Millard, C. L. Outhwaite, R. Kinnersley, R. Freeman, R. D. Gregory, O. Adedoja, S. Gavini, E. Kioko, M. Kuhlmann, J. Ollerton, Z.-X. Ren, T. Newbold, Global effects of land-use intensity on local pollinator biodiversity. *Nature Communications.* **12**, 2902 (2021).

16. R. Hickling, D. B. Roy, J. K. Hill, R. Fox, C. D. Thomas, The distributions of a wide range of taxonomic groups are expanding polewards. *Global Change Biology.* **12**, 450–455 (2006).

17. B. Martay, M. J. Brewer, D. A. Elston, J. R. Bell, R. Harrington, T. M. Brereton, K. E. Barlow, M. S. Botham, J. W. Pearce-Higgins, Impacts of climate change on national biodiversity population trends. *Ecography.* **40**, 1139–1151 (2017).

18. J. Lenoir, R. Bertrand, L. Comte, L. Bourgeaud, T. Hattab, J. Murienne, G. Grenouillet, Species better track climate warming in the oceans than on land. *Nature Ecology & Evolution.* **4**, 1044–1059 (2020).

19. C. A. Halsch, A. M. Shapiro, J. A. Fordyce, C. C. Nice, J. H. Thorne, D. P. Waetjen, M. L. Forister, Insects and recent climate change. *Proceedings of the National Academy of Sciences.* **118**, e2002543117 (2021).

20. C. A. Deutsch, J. J. Tewksbury, R. B. Huey, K. S. Sheldon, C. K. Ghalambor, D. C. Haak, P. R. Martin, Impacts of climate warming on terrestrial ectotherms across latitude. *Proceedings of the National Academy of Sciences.* **105**, 6668–6672 (2008).

21. A. J. Suggitt, R. J. Wilson, N. J. B. Isaac, C. M. Beale, A. G. Auffret, T. August, J. J. Bennie, H. Q. P. Crick, S. Duffield, R. Fox, J. J. Hopkins, N. A. Macgregor, M. D. Morecroft, K. J. Walker, I. M. D. Maclean, Extinction risk from climate change is reduced by microclimatic buffering. *Nature Climate Change.* **8**, 713–717 (2018).

22. P. J. Platts, S. C. Mason, G. Palmer, J. K. Hill, T. H. Oliver, G. D. Powney, R. Fox, C. D. Thomas, Habitat availability explains variation in climate-driven range shifts across multiple taxonomic groups. *Scientific Reports.* **9**, 15039 (2019).

23. T. H. Oliver, S. Gillings, J. W. Pearce-Higgins, T. Brereton, H. Q. P. Crick, S. J. Duffield, M. D. Morecroft, D. B. Roy, Large extents of intensive land use limit community reorganization during climate warming. *Global Change Biology.* **23**, 2272–2283 (2017).

24. M. K. Peters, A. Hemp, T. Appelhans, J. N. Becker, C. Behler, A. Classen, F. Detsch, A. Ensslin, S. W. Ferger, S. B. Frederiksen, F. Gebert, F. Gerschlauer, A. Gütlein, M. Helbig-Bonitz, C. Hemp, W. J. Kindeketa, A. Kühnel, A. V. Mayr, E. Mwangomo, C. Ngereza, H. K. Njovu, I. Otte, H. Pabst, M. Renner, J. Röder, G. Rutten, D. Schellenberger Costa, N. Sierra-Cornejo, M. G. R. Vollstädt, H. I. Dulle, C. D. Eardley, K. M. Howell, A. Keller, R. S. Peters, A. Ssymank, V. Kakengi, J. Zhang, C. Bogner, K. Böhning-Gaese, R. Brandl, D. Hertel, B. Huwe, R. Kiese, M. Kleyer, Y. Kuzyakov, T. Nauss, M. Schleuning, M. Tschapka, M. Fischer, I. Steffan-Dewenter, Climate–land-use interactions shape tropical mountain biodiversity and ecosystem functions. *Nature.* **568**, 88–92 (2019).

25. C. L. Outhwaite, P. McCann, T. Newbold, Agriculture and climate change reshape insect biodiversity worldwide. *Nature* (In press).

26. J. G. Rodger, J. M. Bennett, M. Razanajatovo, T. M. Knight, M. van Kleunen, T.-L. Ashman, J. A. Steets, C. Hui, G. Arceo-Gómez, M. Burd, L. A. Burkle, J. H. Burns, W. Durka, L. Freitas, J. E. Kemp, J. Li, A. Pauw, J. C. Vamosi, M. Wolowski, J. Xia, A. G. Ellis, Widespread vulnerability of flowering plant seed production to pollinator declines. *Science Advances.* **7**, eabd3524 (2021).

27. A. Klein, I. Steffan–Dewenter, T. Tscharntke, Fruit set of highland coffee increases with the diversity of pollinating bees. *Proceedings of the Royal Society of London. Series B: Biological Sciences.* **270**, 955–961 (2003).

28. T. H. Ricketts, G. C. Daily, P. R. Ehrlich, C. D. Michener, Economic value of tropical forest to coffee production. *Proceedings of the National Academy of Sciences.* **101**, 12579–12582 (2004).

29. A. Holzschuh, C. F. Dormann, T. Tscharntke, I. Steffan-Dewenter, Expansion of mass-flowering crops leads to transient pollinator dilution and reduced wild plant pollination. *Proceedings of the Royal Society B: Biological Sciences.* **278**, 3444–3451 (2011).

30. B. A. Woodcock, M. P. D. Garratt, G. D. Powney, R. F. Shaw, J. L. Osborne, J. Soroka, S. A. M. Lindström, D. Stanley, P. Ouvrard, M. E. Edwards, F. Jauker, M. E. McCracken, Y. Zou, S. G. Potts, M. Rundlöf, J. A. Noriega, A. Greenop, H. G. Smith, R. Bommarco, W. van der Werf, J. C. Stout, I. Steffan-Dewenter, L. Morandin, J. M. Bullock, R. F. Pywell, Meta-analysis reveals that pollinator functional diversity and abundance enhance crop pollination and yield. *Nature Communications.* **10**, 1481 (2019).

31. M. A. Aizen, L. A. Garibaldi, S. A. Cunningham, A. M. Klein, Long-term global trends in crop yield and production reveal no current pollination shortage but increasing pollinator dependency. *Current Biology.* **18**, 1572–1575 (2008).

32. A.-M. Klein, B. E. Vaissière, J. H. Cane, I. Steffan-Dewenter, S. A. Cunningham, C. Kremen, T. Tscharntke, Importance of pollinators in changing landscapes for world crops. *Proceedings of the Royal Society B: Biological Sciences.* **274**, 303–313 (2007).

33. F. D. S. Silva, L. G. Carvalheiro, J. Aguirre-Gutiérrez, M. Lucotte, K. Guidoni-Martins, F. Mertens, Virtual pollination trade uncovers global dependence on biodiversity of developing countries. *Science Advances.* **7**, eabe6636 (2021).

34. M. Albrecht, B. Schmid, Y. Hautier, C. B. Müller, Diverse pollinator communities enhance plant reproductive success. *Proceedings of the Royal Society B: Biological Sciences.* **279**, 4845–4852 (2012).

35. C. Brittain, C. Kremen, A.-M. Klein, Biodiversity buffers pollination from changes in environmental conditions. *Global Change Biology.* **19**, 540–547 (2013).

36. M. A. Genung, J. Fox, N. M. Williams, C. Kremen, J. Ascher, J. Gibbs, R. Winfree, The relative importance of pollinator abundance and species richness for the temporal variance of pollination services. *Ecology.* **98**, 1807–1816 (2017).

37. R. Winfree, J. R. Reilly, I. Bartomeus, D. P. Cariveau, N. M. Williams, J. Gibbs, Species turnover promotes the importance of bee diversity for crop pollination at regional scales. *Science.* **359**, 791–793 (2018).

38. M. Dainese, E. A. Martin, M. A. Aizen, M. Albrecht, I. Bartomeus, R. Bommarco, L. G. Carvalheiro, R. Chaplin-Kramer, V. Gagic, L. A. Garibaldi, J. Ghazoul, H. Grab, M. Jonsson, D. S. Karp, C. M. Kennedy, D. Kleijn, C. Kremen, D. A. Landis, D. K. Letourneau, L. Marini, K. Poveda, R. Rader, H. G. Smith, T. Tscharntke, G. K. S. Andersson, I. Badenhausser, S. Baensch, A. D. M. Bezerra, F. J. J. A. Bianchi, V. Boreux, V. Bretagnolle, B. Caballero-Lopez, P. Cavigliasso, A. Ćetković, N. P. Chacoff, A. Classen, S. Cusser, F. D. da Silva e Silva, G. A. de Groot, J. H. Dudenhöffer, J. Ekroos, T. Fijen, P. Franck, B. M. Freitas, M. P. D. Garratt, C. Gratton, J. Hipólito, A. Holzschuh, L. Hunt, A. L. Iverson, S. Jha, T. Keasar, T. N. Kim, M. Kishinevsky, B. K. Klatt, A.-M. Klein, K. M. Krewenka, S. Krishnan, A. E. Larsen, C. Lavigne, H. Liere, B. Maas, R. E. Mallinger, E. M. Pachon, A. Martínez-Salinas, T. D. Meehan, M. G. E. Mitchell, G. A. R. Molina, M. Nesper, L. Nilsson, M. E. O'Rourke, M. K. Peters, M. Plećaš, S. G. Potts, D. de L. Ramos, J. A. Rosenheim, M. Rundlöf, A. Rusch, A. Sáez, J. Scheper, M. Schleuning, J. M. Schmack, A. R. Sciligo, C. Seymour, D. A. Stanley, R. Stewart, J. C. Stout, L. Sutter, M. B. Takada, H. Taki, G. Tamburini, M. Tschumi, B. F. Viana, C. Westphal, B. K. Willcox, S. D. Wratten, A. Yoshioka, C. Zaragoza-Trello, W. Zhang, Y. Zou, I. Steffan-Dewenter, A global synthesis reveals biodiversity-mediated benefits for crop production. *Science Advances.* **5**, eaax0121 (2019).

39. E. Lonsdorf, C. Kremen, T. Ricketts, R. Winfree, N. Williams, S. Greenleaf, Modelling pollination services across agricultural landscapes. *Annals of Botany.* **103**, 1589–1600 (2009).

40. L. N. Hudson, T. Newbold, S. Contu, S. L. L. Hill, I. Lysenko, A. De Palma, B. Collen, R. M. Ewers, G. M. Mace, D. W. Purves, J. P. W. Scharlemann, A. Purvis, The database of the PREDICTS (Projecting Responses of Ecological Diversity In Changing Terrestrial Systems) project. *Ecology and Evolution.* **7**, 145–188 (2017).

41. C. Monfreda, N. Ramankutty, J. A. Foley, Farming the planet: 2. Geographic distribution of crop areas, yields, physiological types, and net primary production in the year 2000. *Global Biogeochemical Cycles.* **22**, GB1022 (2008).

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Competing interests

All authors declare that they have no competing interests.

Data and materials availability

Supplementary Materials

Materials and Methods

Figs. S1-S13

Tables

References