Forecasting range shifts of a dioecious plant species under climate change

Jacob K. Moutouama 0000-0003-1599- $1671^{\rm a.c.1}$, Aldo Compagnoni 0000-0001-8302- $7492^{\rm b}$, and Tom E.X. Miller 0000-0003-3208- $6067^{\rm a}$

Global climate change has triggered an urgent need for predicting the reorganization of Earth's biodiversity. For dioecious species (in which female and male reproductive organs are not on the same individual), it is unclear how commonly unique climate sensitivities of females and males could influence projections for species-level responses to climate change. We developed demographic models of range limitation, parameterized from geographically distributed common garden experiments, with females and males of a dioecious grass species (Poa arachnifera) throughout and beyond its range in the south-central U.S. Female-dominant and two-sex model versions both predict that future climate change will alter population viability and will induce a poleward niche shift beyond current northern limits. However, the magnitude of niche shift was underestimated by the female-dominant model, because females have broader temperature tolerance than males and become mate-limited under female-biased sex ratios. Our result illustrate how explicit accounting for both sexes could enhance population viability forecasts and conservation planning for dioecious species in response to climate change.

13

11

19

22

demography | forecasting | global warming | matrix projection model| population dynamics | sex ratio | range limits |

Rising temperatures and extreme drought events associated with global climate change are leading to increased concern about how species will become redistributed across the globe under future climate conditions (1–3). Species' range limits, when not driven by dispersal limitation, should generally reflect the limits of the ecological niche (4). Niches and geographic ranges are often limited by climatic factors including temperature and precipitation (5). Therefore, any substantial changes in the magnitude of these climatic factors could impact population viability, with implications for range expansions or contractions based on which regions of a species' range become more or less suitable (6, 7).

Forecasting range shifts for dioecious species (most animals and ca. 7\% of plant species) is complicated by the potential for sexual niche differentiation, i.e. distinct responses of females and males to shared climate drivers (8–11). Accounting for sexual niche differentiation is a long-standing challenge in accurately predicting which sex will successfully track environmental change and how this will impact population viability and range shifts (12, 13). Populations in which males are rare under current climatic conditions could experience low reproductive success due to sperm or pollen limitation that may lead to population decline in response to climate change that disproportionately favors females (14). In contrast, climate change could expand male habitat suitability (e.g. upslope movement), which might increase seed set for mate-limited females and favor range expansion (15). Across dioecious plants, for example, studies suggest that future climate change toward hotter and drier conditions may favor male-biased sex ratios (10, 16). Although the response of species to climate warming is an urgent and active area of research, few studies have disentangled the interaction between sex and climate drivers to understand their combined effects on population dynamics and range shifts, despite calls for such an approach (10, 13).

31

32

33

34

35

37

39

40

41

42

43

44

45

46

47

49

54

55

56

57

58

59

The vast majority of theory and models in population biology, including those used to forecast biodiversity responses to climate change, ignore the complication of sex structure (but see 9, 17, 18). Traditional approaches instead focus exclusively on females, assuming that males are in sufficient supply as to never limit female fertility. In contrast, "two-sex" models are required to fully account for demographic differences between females and males and sex-specific responses to shared climate drivers (19, 20). Sex differences in maturation, reproduction, and mortality schedules can generate skew in the operational sex ratio (OSR; sex ratio of individuals available for mating) even if the birth sex ratio is 1:1 (14, 21). Climate and other environmental drivers can therefore influence the OSR via their influence on sex-specific demographic rates. In a two-sex framework, demographic rates both influence and respond to the OSR in a feedback loop that makes two-sex models inherently nonlinear and more data-hungry than corresponding female-dominant models. Given the additional complexity and data needs, forecasts of range dynamics for dioecious species under future climate change that explicitly account for females, males, and their inter-dependence are limited (15, 22).

Tracking the impact of climate change on population via bility (λ) and distributional limits of dioecious taxa depends on our ability to build mechanistic models that take into account the spatial and temporal context of sex specific response to

Significance Statement

The vast majority of models used to forecast population viability and range shifts in response to climate change overlook the complexity of sex structure, and thus the potential for females and males to differ in their sensitivity to climate drivers. Here, we combined common garden experiment with two-sex matrix population models to demonstrate that accounting for only one sex could lead to an underestimation of the impact of climate change on dioecious species, particularly in regions of their range that are biased toward one sex.

J.K.M., A.C. and T.E.X.M. designed the study.
A.C. and T.E.X.M. collected the data.
All authors conducted the statistical analyses and modeling.
J.K.M. drafted the manuscript, and T.E.X.M contributed to revisions

The authors declare no conflict of interest

^a Program in Ecology and Evolutionary Biology, Department of BioSciences, Rice University, Houston, Texas, USA; ^b Institute of Biology, Martin Luther University Halle-Wittenberg, Halle, Germany; and German Centre for Integrative Biodiversity Research (iDiv), Leipzig, Germany

²To whom correspondence should be addressed. E-mail: jmoutouama@gmail.com

climate change, while accounting for sources of uncertainty (6, 23). Structured population models built from demographic data collected from geographically distributed observations or common garden experiments provide several advantages for studying the impact of climate change on species' range shifts (24–26). First, demographic models link individual-level life history events (mortality, development, and regeneration) to population demography, allowing the investigation of factors explaining vital rate responses to environmental drivers (27–29). Second, demographic models have a natural interface with statistical estimation of individual-level vital rates that provide quantitative measures of uncertainty and isolate different sources of variation, features that can be propagated to population-level predictions (30, 31). Finally, structured demographic models can be used to identify which aspects of climate are the most important drivers of population dynamics. For example, Life Table Response Experiments (LTRE) built from structured models have become widely used to understand the relative importance of covariates in explaining variation in population growth rate (32-34).

61

62

63

64

67

68

69

70

71

72

73

74

75

76

77

78

79

81

82

83

84

85

86

87

88

89

90

91

93

94

95

96

97

100

101

102

103

104

106

107

108

109

110

111

112

113

114

In this study, we combined geographically-distributed common garden experiments, hierarchical Bayesian statistical modeling, two-sex population projection modeling, and climate back-casting and forecasting to understand demographic responses to climate change and their implications for past, present, and future range dynamics. Our work focused on the dioecious plant Texas bluegrass (Poa arachnifera), which is distributed along environmental gradients in the south-central U.S. corresponding to variation in temperature across latitude and precipitation across longitude (Fig. 1A)*. Our previous study showed that, despite evidence for differentiation of climatic niche between sexes, the female niche mattered the most in driving longitudinal range limits of Texas bluegrass (35). However, that study used a single proxy variable (longitude) to represent environmental variation related to aridity and did not consider variation in temperature, which is the Developing a rigorous forecast for the implications of future climate change requires that we transition from implicit to explicit treatment of multiple climate drivers, as we do here. Leveraging the power of Bayesian inference, we take a probabilistic view of past, present, and future range limits by quantifying the probability of population viability $(Pr(\lambda \geq 1))$ in relation to climate drivers of demography, an approach that fully accounts for uncertainty arising from multiple sources of estimation and process error. Specifically, we asked:

- 1. What are the sex-specific vital rate responses to variation in temperature and precipitation across the species' range?
- 2. How do sex-specific vital rates combine to determine the influence of climate variation on population growth rate (λ)?
- 3. What is the impact of climate change on operational sex ratio throughout the range?
- 4. What are the likely historical and projected dynamics of the Texas bluegrass geographic niche and how does accounting for sex structure modify these predictions?

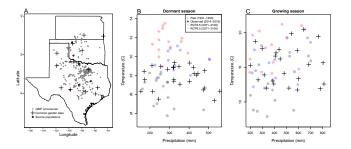


Fig. 1. Experimental gardens and climate of the study region. **A**: Map of 14 experimental garden sites (crosses) in Texas, Oklahoma, and Kansas relative to GBIF occurrences of *Poa arachnifera* (gray points). Red points indicate source populations for plants used in the common garden experiment. **B,C**: Past, future, and observed climate space for growing and dormant seasons. Crosses show observed conditions for the sites and years of the common garden experiment. Gray points show historical (1901-1930) climate normals, and blue and red points show end-of-century (2071-2100) climate normals for RCP4.5 and RCP8.5 projections, respectively, form MIROC5.

ACKNOWLEDGMENTS. This research was supported by National Science Foundation Division of Environmental Biology awards 2208857 and 2225027. We thank the institutions who hosted us at their field station facilities, including

- Bertrand R, et al. (2011) Changes in plant community composition lag behind climate warming in lowland forests. Nature 479(7374):517–520.
- Gamelon M, et al. (2017) Interactions between demography and environmental effects are important determinants of population dynamics. Science Advances 3(2):e1602298.
- Smith MD, et al. (2024) Extreme drought impacts have been underestimated in grasslands and shrublands globally. Proceedings of the National Academy of Sciences 121(4):e2309881120.
- Lee-Yaw JA, et al. (2016) A synthesis of transplant experiments and ecological niche models suggests that range limits are often niche limits. Ecology letters 19(6):710–722.
- Sexton JP, McIntyre PJ, Angert AL, Rice KJ (2009) Evolution and ecology of species range limits. Annu. Rev. Ecol. Evol. Syst. 40:415

 –436.
- Davis MB, Shaw RG (2001) Range shifts and adaptive responses to quaternary climate change. Science 292(5517):673

 –679.
- Pease CM, Lande R, Bull J (1989) A model of population growth, dispersal and evolution in a changing environment. Ecology 70(6):1657–1664.
- Tognetti R (2012) Adaptation to climate change of dioecious plants: does gender balance matter? Tree Physiology 32(11):1321–1324.
- Pottier P, Burke S, Drobniak SM, Lagisz M, Nakagawa S (2021) Sexual (in) equality? a metaanalysis of sex differences in thermal acclimation capacity across ectotherms. Functional Ecology 35(12):2663–2678.
- Hultine KR, et al. (2016) Climate change perils for dioecious plant species. Nature Plants 2(8):1–8.
- Morrison CA, Robinson RA, Clark JA, Gill JA (2016) Causes and consequences of spatial variation in sex ratios in a declining bird species. *Journal of Animal Ecology* 85(5):1298–1306.
- Jones MH, Macdonald SE, Henry GH (1999) Sex-and habitat-specific responses of a high arctic willow, salix arctica, to experimental climate change. Oikos pp. 129–138.
- Gissi E, et al. (2023) Exploring climate-induced sex-based differences in aquatic and terrestrial ecosystems to mitigate biodiversity loss. nature communications 14(1):4787.
- Eberhart-Phillips LJ, et al. (2017) Sex-specific early survival drives adult sex ratio bias in snowy plovers and impacts mating system and population growth. Proceedings of the National Academy of Sciences 114(27):E5474–E5481.
- Petry WK, et al. (2016) Sex-specific responses to climate change in plants alter population sex ratio and performance. Science 353(6294):69–71.
- Field DL, Pickup M, Barrett SC (2013) Comparative analyses of sex-ratio variation in dioecious flowering plants. Evolution 67(3):661–672.
- Ellis RP, et al. (2017) Does sex really matter? explaining intraspecies variation in ocean acidification responses. Biology letters 13(2):20160761.
- Gissi E, Bowyer RT, Bleich VC (2024) Sex-based differences affect conservation. Science 384(6702):1309–1310.
- Gerber LR, White ER (2014) Two-sex matrix models in assessing population viability: when do male dynamics matter? Journal of Applied Ecology 51(1):270–278.
- Miller TE, Shaw AK, Inouye BD, Neubert MG (2011) Sex-biased dispersal and the speed of two-sex invasions. The American Naturalist 177(5):549–561.
- Shelton AO (2010) The ecological and evolutionary drivers of female-biased sex ratios: two-sex models of perennial seagrasses. *The American Naturalist* 175(3):302–315.
 Lynch HJ, et al. (2014) How climate extremes—not means—define a species' geographic
- range boundary via a demographic tipping point. *Ecological Monographs* 84(1):131–132.
- Evans ME, Merow C, Record S, McMahon SM, Enquist BJ (2016) Towards process-based range modeling of many species. Trends in Ecology & Evolution 31(11):860–871.
- 24. Merow C, Bois ST, Allen JM, Xie Y, Silander Jr JA (2017) Climate change both facilitates

116

117

118

119

120

121

122

123

124

125

126

127

128

129

130

131

132

133

134

135

136

137

138

139

140

141

142

143

144

145

146

147

148

149

150

151

152

153

154

155

156

157

158

159

160

161

162

163

164

165

166

167

168

^{*} Fig. A does not show what we are saying here. Maybe I should add the Figure with the raster

- and inhibits invasive plant ranges in new england. Proceedings of the National Academy of
 Sciences 114(16):E3276–E3284.
- 25. Schwinning S, Lortie CJ, Esque TC, DeFalco LA (2022) What common-garden experiments
 tell us about climate responses in plants.
- 26. Schultz EL, et al. (2022) Climate-driven, but dynamic and complex? a reconciliation of competing hypotheses for species' distributions. *Ecology letters* 25(1):38–51.

175

176

- Ehrlén J, Morris WF (2015) Predicting changes in the distribution and abundance of species under environmental change. Ecology letters 18(3):303–314.
- 28. Louthan AM, et al. (2022) Climate change weakens the impact of disturbance interval on the growth rate of natural populations of venus flytrap. *Ecological Monographs* 92(4):e1528.
- Dahlgren JP, Bengtsson K, Ehrlén J (2016) The demography of climate-driven and density regulated population dynamics in a perennial plant. *Ecology* 97(4):899–907.
- Elderd BD, Miller TE (2016) Quantifying demographic uncertainty: Bayesian methods for integral projection models. *Ecological Monographs* 86(1):125–144.
- 183
 Ellner SP, et al. (2022) A critical comparison of integral projection and matrix projection models
 for demographic analysis: Comment. *Ecology*.
- 32. Ellner SP, Childs DZ, Rees M, , et al. (2016) Data-driven modelling of structured populations.
 A practical guide to the Integral Projection Model. Cham: Springer.
- 33. Hernández CM, Ellner SP, Adler PB, Hooker G, Snyder RE (2023) An exact version of life
 table response experiment analysis, and the r package exactltre. Methods in Ecology and
 Evolution 14(3):939–951.
- 34. Czachura K, Miller TE (2020) Demographic back-casting reveals that subtle dimensions of climate change have strong effects on population viability. *Journal of Ecology* 108(6):2557– 2570.
- Miller TE, Compagnoni A (2022) Two-sex demography, sexual niche differentiation, and
 the geographic range limits of texas bluegrass (poa arachnifera). The American Naturalist

Moutouama et al. PNAS | September 23, 2024 | vol. XXX | no. XX | 3