

Original Articles

Climate Change Poses Risk of Local Extinction for Temperate Forest Species in Oaxaca, Mexico

Juan Carlos Guzmán-Santiago ^{ab} , Vicente Espinosa Hernández ^a , Rogelio Omar Corona-Núñez ^{cd*} , Juan Camilo Zapata-Calero ^e , Iván Felipe Benavides-Martínez ^f , Wenceslao Santiago-García ^g 

^a Colegio de Postgraduados, Posgrado en Ciencias Forestales, Campus Montecillos, Texcoco, Estado de México, México.

^b Centro de Investigación, Divulgación y Asesoría Técnica Forestal y Agropecuaria SC, Isabel la Católica, Tlaxiaco, Oaxaca, México.

^c Universidad Nacional Autónoma de México, Facultad de Ciencias, Investigación Científica, Coyoacán, Ciudad de México, México.

^d Universidad Autónoma Metropolitana Unidad Cuajimalpa, Contadero, Cuajimalpa de Morelos, Ciudad de México, México.

^e Universidad Nacional de Colombia, Sede Palmira, Palmira, Colombia.

^f Datambiente, Departamento de Analítica de Datos Ambientales, Bogotá, Colombia.

^g Universidad de la Sierra Juárez, División de Estudios de Postgrado-Instituto de Estudios Ambientales, Oaxaca, México.

Article Info

Received: 04-07-2025

Accepted: 21-11-2025

Published: 25-03-2026

Corresponding author: r.o.corona_nunez@ciencias.unam.mx

How to cite: Guzmán-Santiago J. C., Espinosa Hernández V., Corona-Núñez R. O., Zapata-Calero J. C., Benavides-Martínez I. F., Santiago-García W. (2026). Climate Change Poses Risk of Local Extinction for Temperate Forest Species in Oaxaca, Mexico. *Bosque*, 47, e4702, 2026.

<https://doi.org/10.4206/Bosque.e4702>

Abstract

Climate change is expected to drive significant shifts in species distributions, particularly in temperate forest ecosystems. Areas of long-term environmental stability, known as climatic refugia, may play a key role in preserving biodiversity under future climate scenarios. This study assessed the potential impact of climate change on the distribution of eleven temperate forest tree species in Oaxaca, Mexico. Species distribution models were developed for eleven tree-species of temperate forest using Random Forest, Generalized Additive Models, and Generalized Linear Models. Among these, Random Forest achieved the highest predictive performance. Future distributions were projected under two shared socioeconomic pathways (SSP126 and SSP585) for 2040 and 2100. Results indicate that under the SSP585 scenario, all species could lose over 90% of their current potential range by 2100. Although some are classified as “Least Concern” by the IUCN (2024), their local vulnerability is pronounced. Additionally, projections suggest declining representation within Natural Protected Areas, raising concerns about their long-term conservation capacity. These findings highlight the urgency of implementing adaptive conservation measures, including the identification and management of climatic refugia, promotion of assisted migration, and reduction of anthropogenic pressures to enhance species resilience.

Keywords: ecological niche, forest species distribution, *Pinus*, *Quercus*.

Introduction

Climate change (CC) has emerged as one of the main drivers of the transformation of terrestrial ecosystems globally (O'Neill et al., 2016). CC has modified the distribution patterns and resilience of forest ecosystems in several regions, including Mexico (Metcalf et al., 2010; Rodríguez-Ramírez et al., 2015). For example, CC is causing changes in species distribution, modifying its altitudinal and latitudinal ranges (Dyderski et al., 2018), which can promote the local extinction of species (Cruz-Cárdenas et al., 2016; Mendoza-Ponce et al., 2020). In addition, CC alters the typical functioning of species and ecosystems due to changes in the annual growth, affecting the viability of seeds, reducing the vigour of plants and affecting the survival of new growth (Murga-Orrillo et al., 2021), among other alterations. Therefore, CC represents a challenge for the adaptive capacity of forest species, acting as an environmental filter (Moeslund et al., 2013).

Global Circulation Models (GCMs) seek to simulate the behaviour of the atmosphere at the global level and its influence on future climate (Rivera & Amador, 2008). Different authors, through ecological niche modeling, have sought to understand the effect of future climate on the distribution of species (Guisan & Zimmermann, 2000; Manzanilla-Quijada et al., 2020). Ecological niche modeling, also known as species distribution modeling, has been instrumental in representing the potential distribution of species and changes over time. This distribution is based on recognizing species-specific characteristics such as climatic preference, tolerance to disturbance and resource requirements (Peterson et al., 2012). Currently there is a wide range of tools that help for this purpose. Among the most widely used models, the following stand out: GARP (Stockwell, 1999), Maxent (Phillips et al., 2004) and Random Forest (Evans et al., 2011).

Mexico is recognized as the center of origin and diversification for the genus *Pinus*, harboring a high number of endemic species. It contains 41% of all known *Pinus* species (Govaerts et al., 1998) and 76% of all *Quercus* species, while 51% of all species worldwide are considered endemic to Mexico (Valencia, 2004). The wide geographic distribution of *Pinus* across the country enables its coexistence with other conifers such as *Abies*, *Cupressus*, *Pseudotsuga*, *Juniperus*, and various hardwoods (Mendoza-Ponce et al., 2018). The evolutionary history of *Pinus* and *Quercus* reveals distinct phylogenetic patterns shaped by climatic changes and genetic adaptations, suggesting a shared evolutionary trajectory between the two genera (Hao et al., 2015; Li et al., 2021; Zeb et al., 2022). During the early Holocene, the genus *Pinus* expanded into various regions of Mexico in response to global warming (Lozano-García et al., 2007; Ortega-Rosas et al., 2008). These environmental conditions promoted the diversification of

numerous species and continue to play a critical role in the structure and function of temperate forest ecosystems (Gernandt & Pérez-de la Rosa, 2014). As a result, Mexico is considered a megadiverse country—not only due to its biological richness, but also its climatic heterogeneity.

At the subnational scale, the state of Oaxaca exhibits the highest biological diversity of temperate forest species in Mexico (Valencia, 2004; de Jesús Ordóñez & Rodríguez, 2008). However, this rich biodiversity is increasingly threatened by accelerated deforestation (Mendoza-Ponce et al., 2018), recurrent fires (Corona-Núñez et al., 2020), and the adverse effects of CC. Understanding the drivers and consequences of these threats is essential for developing effective conservation strategies (D'Amato & Palik, 2021), especially for species that are endangered and/or vulnerable (Trejo et al., 2011).

Although temperate forests in Oaxaca harbor high biodiversity and play a critical ecological role, few studies have quantitatively examined how CC may affect the distribution and persistence of key forest species in the region (Guzmán-Santiago, 2023). This knowledge gap limits the development of effective biodiversity management strategies under future CC scenarios. In response, this study evaluates the potential risk of range contractions and local extinctions among temperate forest tree species, with particular attention to their representation within Natural Protected Areas (NPAs) and the identification of climatic refugia as potential strongholds for long-term conservation.

Methods

Study area. The state of Oaxaca, located in southeastern Mexico (18°42'–15°39' N, 93°52'–98°32' W), reaches a maximum elevation of 3,750 m.a.s.l. and covers 95,354 km², ranking fifth nationally in area. It is divided into eight regions: *Cañada*, *Costa*, *Istmo*, *Mixteca*, *Papaloapan*, *Sierra Norte*, *Sierra Sur* and *Valles Centrales* (de Jesús Ordóñez, 2000). Each of these regions has distinct biodiversity shaped by unique biogeographic, bioclimatic, and biocultural conditions. Oaxaca encompasses both tropical and temperate biomes (Figure 1), with temperate forests accounting for over 50% of the state's forested area (Mendoza-Ponce et al., 2018).

Species inventory. Species location records were obtained from the National Forest Inventory (CONAFOR, 2012), iNaturalist (iNaturalist, 2025), Global Biodiversity Information Facility (GBIF) and San Juan Quiahije Forest Inventory (IF-SJQ) (Table S1). To avoid pseudo-replication, records were clustered by merging observations within a 1-km², resulting in 1,383 independent records for 11 tree species (Figure 1 and Table S1). The most abundant genus is *Quercus* with

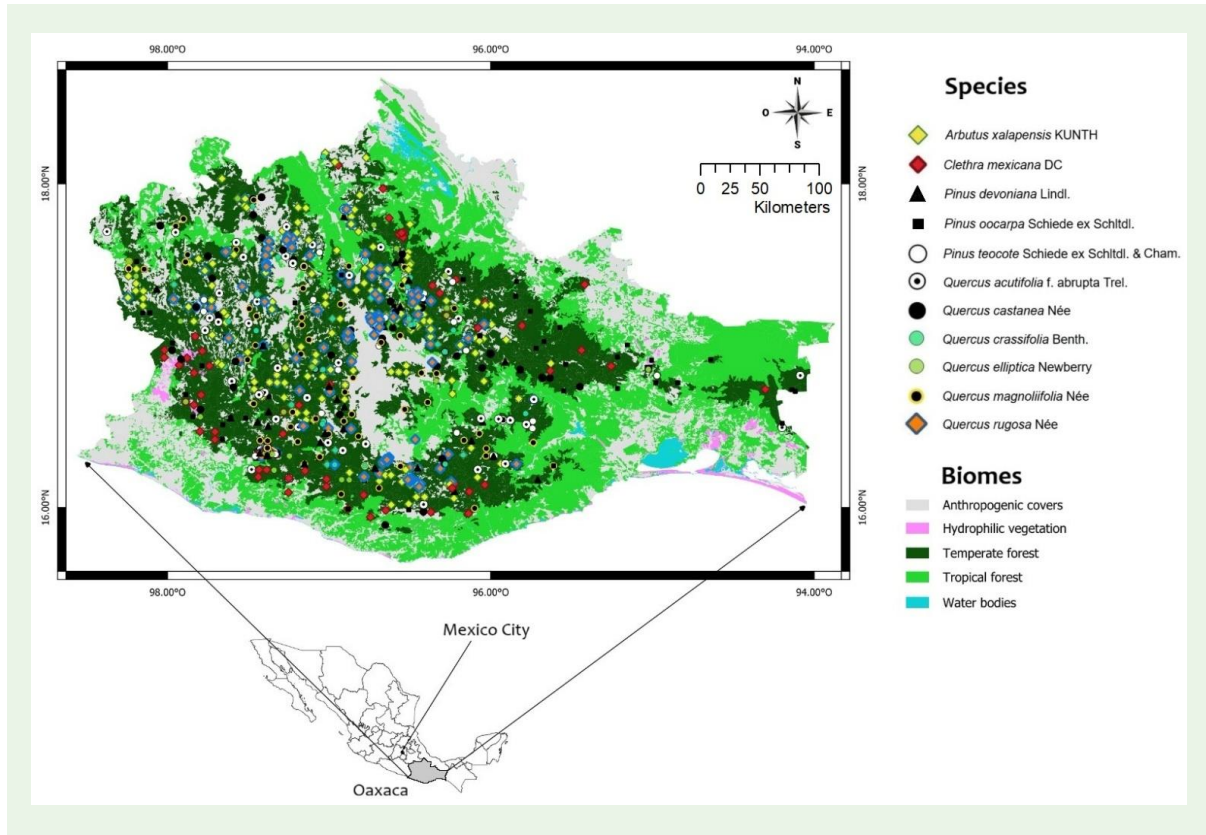


Figure 1. Study area showing the distribution of dominant biomes in the State of Oaxaca and the distribution of the evaluated species.

49.0% of the records, followed by *Arbutus* with 21.8% and *Pinus* with 21.4%. The species with the highest number of records was *Arbutus xalapensis* (21.8%) > *Pinus oocarpa* (12.4%) > *Quercus rugosa* (11.1%).

Bioclimatic variables. We used 19 bioclimatic variables from WorldClim v2.1 (Fick & Hijmans, 2017), representing both current (1970–2000) and future climate conditions (2021–2040 and 2081–2100), based on CMIP6 projections from the CNRM-CM6-1 global circulation model. CNRM-CM6-1 is a comprehensive Earth system model developed by CNRM/CERFACS for CMIP6, simulating climate dynamics and future scenarios using coupled atmosphere–ocean–land–ice components (Voldoire, 2023). These layers capture key climatic dimensions, including annual trends (e.g., mean annual temperature, annual precipitation), seasonality (e.g., temperature and precipitation ranges), and extreme or limiting environmental factors (e.g., temperature of the coldest and warmest month, precipitation of the wettest and driest quarters) (Table S2). All bioclimatic layers (Bio1–Bio19) were obtained from the WorldClim repository and have a spatial resolution of 30 arc-seconds (approximately 1 km² per pixel).

To complement climatic predictors, we incorporated topographic variables: altitude (derived from SRTM data) and slope (calculated from the SRTM). The SRTM Digital Elevation Model is a near-global dataset of Earth’s terrain elevation, captured by NASA’s Shuttle Radar Topography Mission in 2000 (NASA-JPL, 2013). These layers were processed to match the resolution of the bioclimatic variables, ensuring spatial consistency across all environmental predictors.

Species distribution models. Species distribution models were calibrated using 80% of the occurrence data, with the remaining 20% reserved for independent blind validation to assess predictive performance. For each species, pseudo-absences were generated randomly at distances >1 km to ensure model contrast and reduce spatial autocorrelation. Distributions were modeled using three algorithms: Random Forest (RF), Generalized Additive Models (GAM), and Generalized Linear Models (GLM), selected for their demonstrated accuracy and ecological interpretability in ecological and species distribution modelling applications (Elith & Leathwick, 2009; Corona-Núñez et al., 2017). The models were evaluated using Receiver Operating Characteristic curves (ROC), True Skill

Statistic (TSS), and the Kappa index (Swets, 1988; Allouche et al., 2006). The best-performing algorithm per species was selected for subsequent analysis.

Distributional ranges were defined by selecting pixels with suitability scores >0.5 , a threshold chosen to emphasize areas with a higher likelihood of species presence while reducing uncertainty in marginal zones. To evaluate model performance and identify optimal classification thresholds, we applied the Youden Index. The Youden Index is a widely used diagnostic metric in species distribution modeling that maximizes the sum of sensitivity and specificity, thereby balancing omission and commission errors. This index was selected for its robustness in threshold optimization, especially when presence–absence data are imbalanced or when both types of errors carry ecological consequences. We retained the >0.5 cutoff for spatial projections to ensure consistency across scenarios and facilitate ecological interpretation. This approach enhances model reliability and supports conservation-related decision-making by minimizing misclassification risks (Jiménez-Valverde & Lobo, 2007). This decision reflects a conservative approach that prioritizes ecological plausibility over statistical optimization, particularly in the context of future climate projections. Binary presence/absence maps were derived from these thresholds and used to estimate potential distributions under historical climate conditions (1970–2000) and two future periods: 2021–2040 (2040) and 2081–2100 (2100). Projections incorporated two contrasting Shared Socioeconomic Pathways (SSPs): SSP126 (optimistic) and SSP585 (pessimistic), representing divergent trajectories in global development and emissions.

Species at risk and changes in potential distribution. To assess the impact of CC on species distributions, we compared present and future projections across all scenarios and time periods. Distributional changes were classified as positive when species expanded their current distribution range, and negative when contractions occurred. To quantify these dynamics, we calculated three spatial metrics: loss (areas currently suitable but projected to become unsuitable), gain (areas unsuitable at present but projected to become suitable), and permanence (areas consistently suitable across time and scenarios, functioning as climate refugia). These metrics were derived from binary presence–absence maps generated using a standardized threshold, ensuring comparability across species and scenarios.

Climatic refugia were defined as zones of spatial overlap between present and future suitable habitats, indicative of climatic stability and ecological continuity. These areas are expected to remain within a species' environmental tolerance limits despite broader climatic shifts, and were identified using pixel-wise intersection across all

scenario outputs. We then aggregated refugia across species to delineate multi-species refugial regions, which may act as buffers against local extinction and support long-term persistence.

To evaluate the conservation relevance of these refugial zones, we assessed their spatial congruence with existing Natural Protected Areas (NPAs), following approaches similar to others (Keppel et al., 2012; Morelli et al., 2016). This comparison allowed us to identify priority areas where current protection aligns with future climatic stability. Finally, to quantify conservation potential within NPAs, we calculated the average species richness across all future scenarios, providing a comparative measure of species density retained under projected climatic conditions.

To contextualize the conservation relevance of modeled species, we identified the threat statuses using the IUCN Red List (2024). All analyses were conducted in the RStudio environment (R-Core-Team, 2018), integrating outputs from species distribution models and CC scenarios.

Results

Model performance. For all species, the RF model showed the best performance, followed by GAM and finally GLM. On average RF resulted with a performance of 99.3%, GAM 93.1% and GLM 83.4%. The worst performance of GLM was with the Kappa $<$ ROC $<$ TSS metric. Specifically, for *Pinus teocote*, *Quercus elliptica* and *Q. castanea* species RF achieved a performance $>98%$, followed by GAM reaching values $>86%$ (Table 1). The species sensitivity ranges from 0.62 to 0.99 with an average of 0.93 ± 0.11 (Table S3).

Changes in species distribution. The Mixteca, Valles Centrales, Sierra Norte and Sierra Sur regions are where most of the species evaluated are concentrated (Figure 2). However, under CC scenarios all regions will show a reduction in their distribution, with low permanence in relation to their potential distribution. Gains in distribution are estimated to occur in the Sierra Sur and Costa (towards the south of the state), Mixteca (west), Valles Centrales (center) and Papaloapan (north).

Species exhibited diverse spatial responses to projected climate scenarios, reflecting varying degrees of resilience and sensitivity to environmental change (Table S4). Notably, *Pinus* species generally shifted upward in both altitude and latitude. In contrast, *Quercus* species displayed a divergent pattern, characterized by reductions in altitudinal range and increases in mean latitude. *Arbutus xalapensis* exhibited remarkable stability, with no detectable changes in altitude, latitude, or slope across all scenarios, suggesting strong resilience to climatic shifts. *Clethra mexicana* showed moderate latitudinal

Table 1. Validation metrics per species for each model. Kappa: Cohen’s kappa statistic, ROC: area under the curve, TSS: true skill statistic, GAM: Generalized Additive Models, GLM: Generalized Linear Models, RF: Random Forest.

Species	Kappa			ROC			TSS		
	GAM	GLM	RF	GAM	GLM	RF	GAM	GLM	RF
<i>Arbustus xalapensis</i>	86.4	92.0	99.3	93.7	92.0	99.3	94.7	92.0	99.3
<i>Clethra mexicana</i>	90.7	82.4	99.1	94.4	82.4	99.1	94.4	82.4	99.1
<i>Pinus devoniana</i>	96.5	89.5	99.9	96.5	89.5	99.9	96.5	89.5	99.9
<i>Pinus oocarpa</i>	94.7	74.9	98.8	94.7	84.8	99.9	95.9	83.6	99.9
<i>Pinus teocote</i>	95.5	79.1	98.5	94.0	98.5	98.5	95.5	88.1	98.5
<i>Quercus acutifolia</i>	94.9	49.2	99.2	94.9	71.2	99.9	94.9	84.7	99.2
<i>Quercus castanea</i>	90.4	69.1	98.9	90.4	80.9	98.9	90.4	86.2	98.9
<i>Quercus crassifolia</i>	86.5	89.2	99.6	93.7	95.6	99.9	93.6	95.6	99.9
<i>Quercus elliptica</i>	96.3	74.1	98.8	96.3	75.3	98.8	96.3	77.8	98.8
<i>Quercus magnoliifolia</i>	86.0	79.1	99.2	89.9	85.3	99.2	90.7	85.3	99.2
<i>Quercus rugosa</i>	88.9	83.0	99.3	91.5	85.6	99.3	92.2	85.6	99.3

displacement and a notable decrease in altitude under SSP585-2100, indicating potential vulnerability to warming trends and habitat compression. Among *Pinus* species, *P. devoniana* exhibited gradual increases in altitude and latitude, with slight slope variability, indicating a shift toward higher terrain. In contrast, *P. oocarpa* and *P. teocote* both displayed a retreat to lower, flatter landscapes, particularly under SSP585-2100. Oak species responded more uniformly: *Quercus acutifolia*, *Q. castanea*, and *Q. crassifolia* demonstrated consistent declines in elevation and slope.

In the SSP126-2040 an average loss of 31.4±7.7% among all species is estimated, where *Pinus* would lose 39.2±8.3% and *Quercus* 29.9±5.3% (Figure 3 and Table S5), while for the year 2100 the average loss of the species amounts to 44.0±9.4%, *Pinus* would lose 53.9±9.8% and *Quercus* 42.0±6.7%. The species of greatest losses are *Pinus oocarpa*, *P. devoniana* and *Quercus rugosa* with 62.8%, 55.5% and 51.2%, respectively. The species with the lowest losses were *Q. castanea* with 32.9%, *Arbutus xalapensis* with 33.8% and *Q. magnoliifolia* with 36.4%.

These same analyses for the SSP858-2040 scenario indicate that on average the species would lose 32.6±7.8% of their potential distribution, while by the year 2100 the losses are estimated to increase to 88.1±6.1% of their potential distribution (Figure 3 and Table S5). The *Pinus* genera will show a loss of 92.2±3.9 and *Quercus* of 88.0± 6.2%. The species *Pinus devoniana*, *P. oocarpa*, *Quercus*

acutifolia, *Q. crassifolia* and *Q. elliptica* will have losses > 90%. While the species with the lowest losses are *Arbutus xalapensis* with 78.4% and *Quercus castanea* with 78.6%.

Although most of the anticipated changes in distribution are expected to result in a reduction of the species’ current extent, these losses will be partly compensated by minor expansions into new regions where the species are currently absent (Figure 3 and Table S5). For example, when analysing the expansions the SSP126-2100 scenario, on average the species will increase their surface area by 12.7± 8.8%, where *Pinus* will be 23.3± 9.1% and *Quercus* 8.8±5.1. The species with the greatest expansion is *Pinus teocote* (33.7%), and those with the least gains are *Quercus acutifolia* (3.1%) and *Q. castanea* (5.0%). For the SSP585-2100 scenario the average expansion of all species is lower (8.5± 7.3%), where *Pinus* is expected to have a gain of 17.0± 3.6% and *Quercus* 6.7±5.9%. *Pinus oocarpa* will have the highest expansion of all *Pinus* species (20.5%). While those showing the lowest expansion capacity are *Quercus rugosa* (6.1%) > *Q. magnoliifolia* (3.5%) > *Q. castanea* (2.3%) > *Q. acutifolia* (0.3%).

Climatic refugia and the importance of natural protected areas.

Under CC scenarios, projected shifts in species distributions are not expected to align with their current spatial extent, including areas within NPAs. Our results indicate that in the SSP126-2100 scenario,

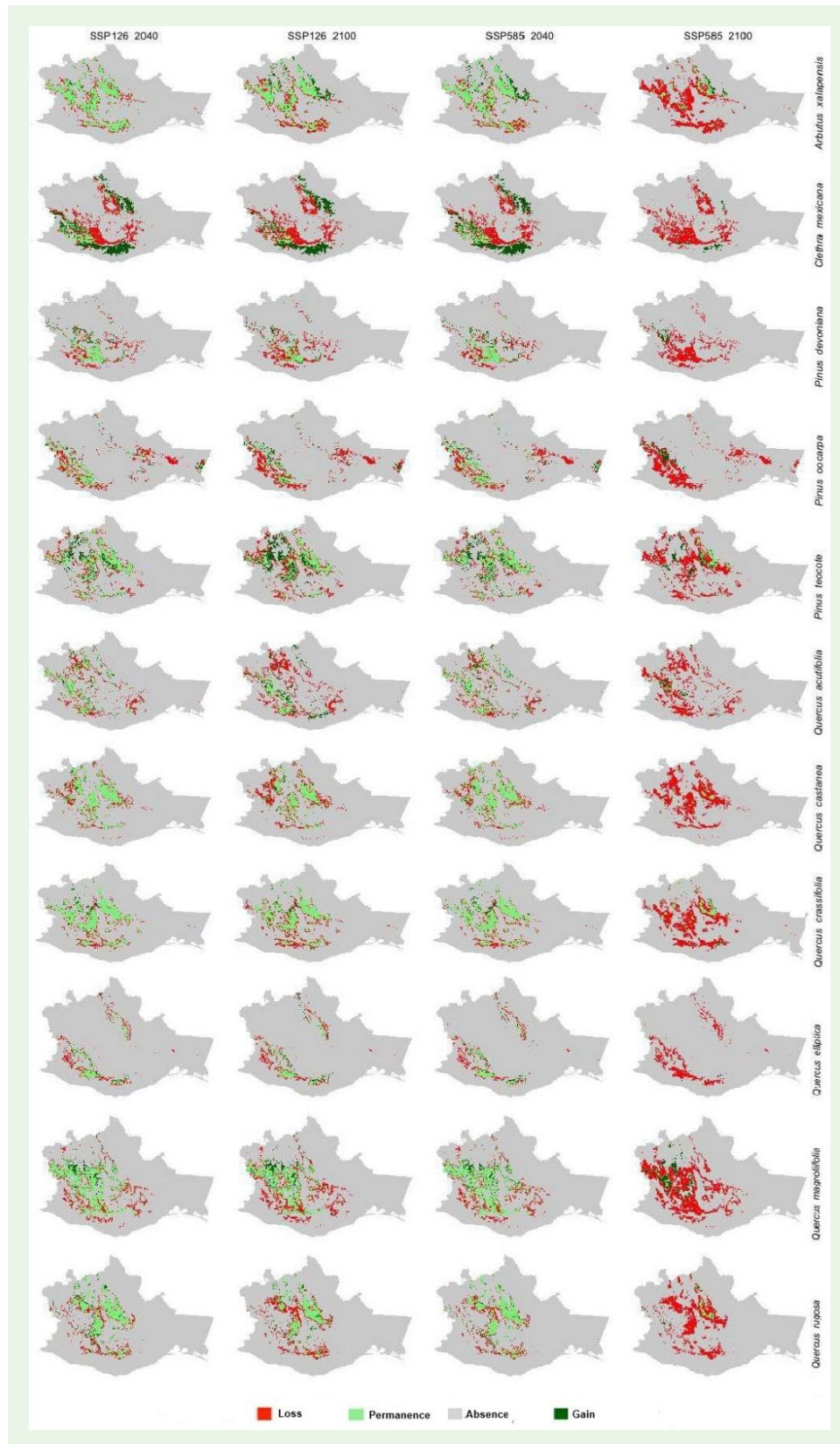


Figure 2. Changes in the distribution of species evaluated by period and climate change scenario.

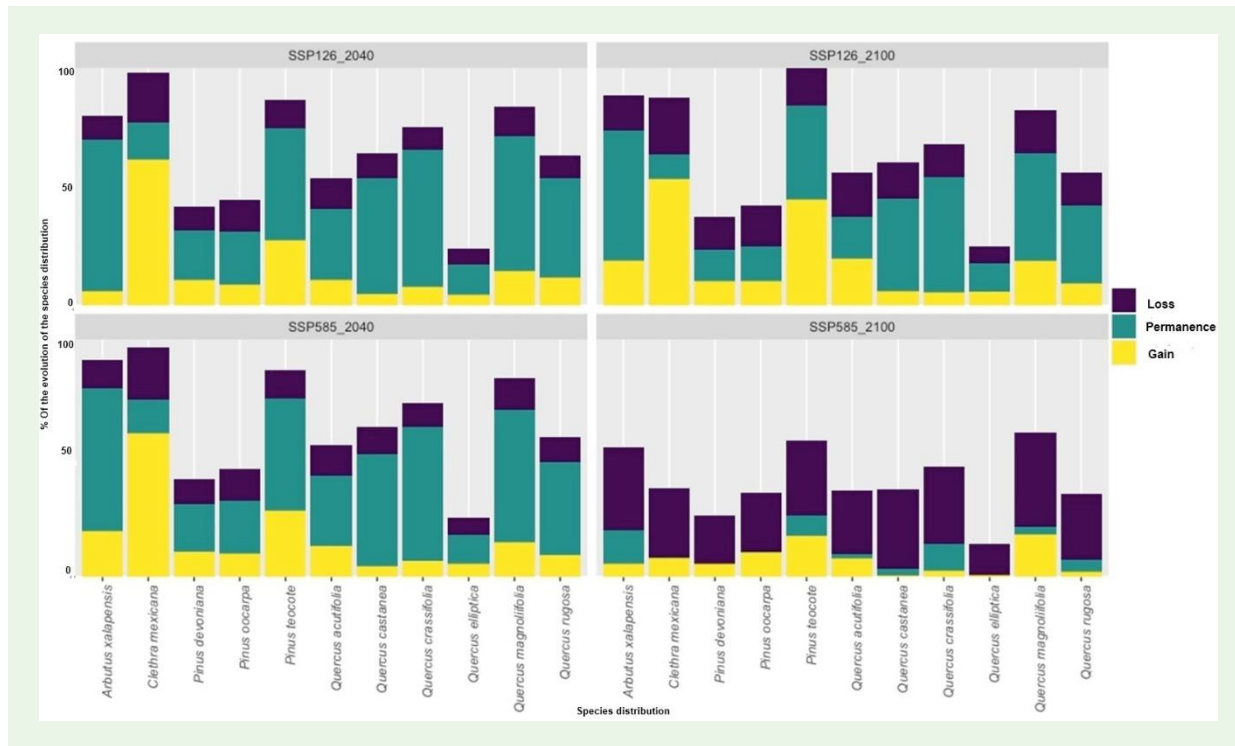


Figure 3. Bar graph of the distribution of species in different climatic scenarios.

the total extent of suitable habitat is projected to decline by 23.1%, while under the more severe SSP585-2100 scenario, this reduction could reach up to 81.0% (Figure 4). This contraction is accompanied by a decrease in species richness. Under SSP126-2100, up to seven species may be lost from areas where complete extinction is not recorded, while under SSP585-2100, this number increases to nine species (Figure 5). Consequently, NPAs will also experience a reduction in the extent of temperate forest they encompass, along with a diminished capacity to conserve forest species under future climate conditions.

Our study shows that NPAs conserve a maximum of 9 forest species of the 11 species evaluated for Oaxaca (Figure 4). However, under CC scenarios this number is reduced to a maximum of 8 and 6 for the SSP126 scenario for the years 2040 and 2100, respectively. For SSP585 the reduction is even more drastic, with a maximum of 7 and 5 for the same years. In addition, NPAs currently protect an average richness of 2.4 ± 1.6 species per km^2 . Under CC scenarios, this richness will also be affected. In the SSP126 scenario by the year 2100 this richness decreases to 1.7 ± 1.8 species, and in SSP585 it is 0.4 ± 1.0 species per km^2 .

Discussion

Models performance. Among the modeling algorithms tested, RF demonstrated the highest performance, followed by GAM and GLM, consistent with previous findings (Evans et al., 2011; Barrio-Anta et al., 2020; Alegria et al., 2023). It is well established that both model choice and the quantity of calibration data significantly influence model accuracy (Corona-Núñez et al., 2017). Additionally, our results indicate that species-specific biological traits may impact model performance. Species with broad environmental tolerances tend to exhibit greater spatial heterogeneity, complicating accurate modeling. Conversely, species with narrow bioclimatic ranges are typically modeled with higher precision, corroborating earlier studies (Wisiz et al., 2008; Guzmán-Santiago, 2023).

Changes in species distribution. Environmental change in temperate regions is driving *Pinus* species to higher elevations and more extreme latitudes, where conditions may offset precipitation deficits and offer more favourable temperatures (Castellanos-Acuña et al., 2014; Rehfeldt et al., 2014; Safaei et al., 2021; Mirhashemi et al., 2023). Interestingly, some *Quercus* species may reduce their altitudinal range while increasing their mean latitude, a pattern also observed by

Barradas et al. (2011). However, in addition to altitude, topographic slope has also been shown to constrain species distributions and development, consistent with the findings of this study (Moeslund et al., 2013; Cueva et al., 2019; Antúnez, 2021).

Pinus and *Quercus* species are particularly vulnerable to shifts in lower elevational limits, which may reduce available habitat and hinder growth rates (Castellanos-Acuña et al., 2015). Areas where species distributions remain stable under CC scenarios may function

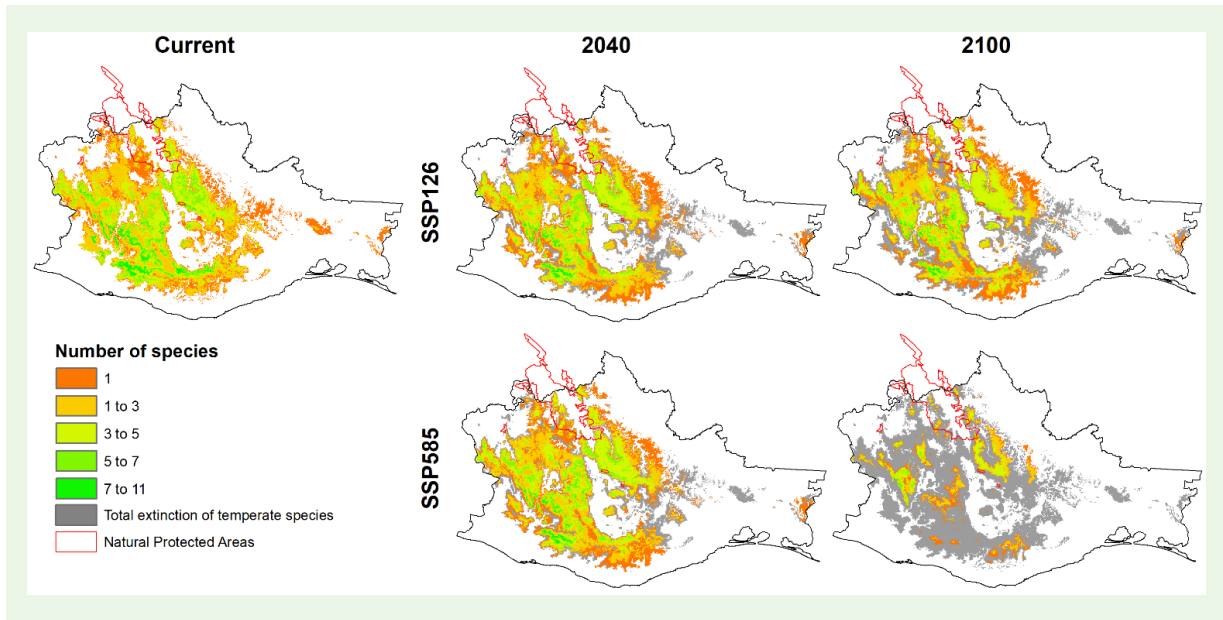


Figure 4. Species richness in temperate forests. Natural Protected Area polygons in the state of Oaxaca are highlighted in red.

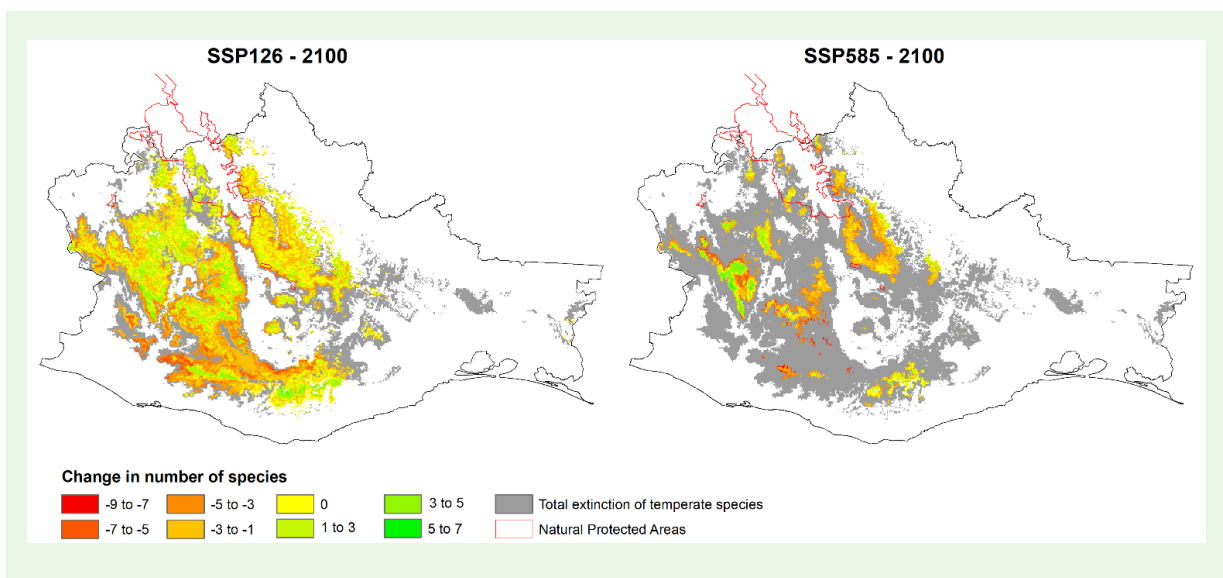


Figure 5. Changes in temperate forest species richness by the year 2100. Natural Protected Area polygons in the state of Oaxaca are highlighted in red.

as climatic refugia. These zones are essential for preserving genetic diversity and maintaining biotic interactions under reduced levels of disturbance. Similar patterns have been observed in studies of terrestrial vertebrates (Mendoza-Ponce et al., 2020). These refugia thus warrant targeted conservation efforts to mitigate pressures from deforestation and ecosystem degradation.

Species vulnerability. Our findings indicate that temperate species are projected to experience reductions in their potential distribution ranges. Although some may exhibit greater capacity to migrate into newly suitable areas, such shifts are insufficient to offset overall habitat loss. Consequently, localized extinctions are expected across various regions of Oaxaca. This vulnerability is further underscored by the fact that most of these species are currently classified as “Least Concern” by the IUCN (2024), with *Quercus acutifolia* listed as “Vulnerable”. Global conservation statuses may overlook region-specific threats, especially in fragile temperate ecosystems where climate-induced range reductions intensify conservation challenges.

To reduce the vulnerability of these ecosystems, it is crucial to implement CC adaptation measures, including the establishment of NPAs within identified climate refugia. Additionally, landscape management should prioritize the reduction of anthropogenic pressures, such as forest land conversion (Mendoza-Ponce et al., 2020), and the prevention of fires (Corona-Núñez et al., 2020). Assisted migration strategies (Gray et al., 2011; Dumroese et al., 2015) may also play a complementary role by supporting species persistence within future suitable habitats. Collectively, these actions are essential for enhancing the resilience and long-term viability of temperate forest species under CC scenarios.

Importance of natural protected areas under climate change scenarios. NPAs were established globally to safeguard biodiversity. However, their initial designation did not account for their function as climate refugia, prompting recent concerns over their capacity to conserve species under changing climatic conditions (Dumroese et al., 2015). Our findings suggest that the representation of temperate forest species within NPAs will decline. This reduction affects not only total area but also species richness and the number of species each NPA conserves. Such shifts will compromise ecosystem functionality at multiple levels. Alterations in forest species composition and structure may disrupt trophic interactions by diminishing feeding, nesting, and sheltering sites (Gray et al., 2011; Millar & Stephenson, 2015). Additionally, changes in microclimates and biogeochemical cycles are expected, potentially undermining essential ecosystem services. To align with global goals such as the Kunming-Montreal Global Biodiversity Framework and the 2030 Agenda, it is critical

to integrate climate refugia into conservation planning. Protecting these stable zones can enhance the resilience of NPAs and ensure the persistence of species in a changing climate (Keppel et al., 2024).

Conclusions

The results show that regardless of the algorithm used, and regardless of future CC scenarios, in all cases temperate forest species will show significant changes in their potential distribution, mainly with high losses of their potential extent, being SSP585-2100 where these changes will be more noticeable. Changes in species distributions expose the fragility of temperate forest ecosystems, through the modification of their species composition, and most likely of abundances. It is essential to establish adaptation strategies to CC that seek to conserve climatic refuges to conserve germplasm. Finally, it is important to evaluate the efficiency of NPAs as climatic refuges for biodiversity conservation.

Author contributions

ROCN, JCGS, and JCZC carried out data analysis and document writing. VEH, IFBM and WSG reviewed the wording.

Funding

The first author thanks the Secretaría de Ciencia, Humanidades, Tecnología e Innovación (SECIHTI) for the scholarship awarded for the postdoctoral stay, with project ID 2524710 (managed by the same beneficiary). Also, ROCN want to grate for its postdoctoral scholarship received from SECIHTI.

Acknowledgements

The agrarian authorities of the community of San Juan Quiahije, Juquila, and the forestry technicians are thanked, Ing. Gregorio Morales and Biol. María del Sol for her followers in the various field and cabinet activities. The research is carried out at the College of Postgraduates, Postgraduate in Forest Sciences, Montecillo Campus, Texcoco, Mexico.

References

- Alegria, C., Almeida, A. M., Roque, N., Fernandez, P., & Ribeiro, M. M. (2023). Species distribution modelling under climate change scenarios for maritime pine (*Pinus pinaster* Aiton) in Portugal. *Forests*, *14*(3), 591.
- Allouche, O., Tsoar, A., & Kadmon, R. (2006). Assessing the accuracy of species distribution models: prevalence, kappa and the true skill statistic (TSS). *Journal of applied ecology*, *43*(6), 1223-1232.
- Antúnez, P. (2021). Influence of physiography, soil and climate on *Taxus globosa*. *Nordic Journal of Botany*, *39*(3). DOI: <https://doi.org/10.1111/njb.03058>
- Barradas, V. L., Tapia Vargas, L. M., & Cervantes Pérez, J. (2011). Consecuencias del cambio climático en la ecofisiología vegetal de un bosque templado en Veracruz. *Revista mexicana de ciencias agrícolas*, *2*, 183-194.
- Barrio-Anta, M., Castedo-Dorado, F., Cámara-Obregón, A., & López-Sánchez, C. A. (2020). Predicting current and future suitable habitat and productivity for Atlantic populations of maritime pine (*Pinus pinaster* Aiton) in Spain. *Annals of Forest Science*, *77*, 1-19.
- Castellanos-Acuña, D., Lindig-Cisneros, R., & Sáenz-Romero, C. (2015). Altitudinal assisted migration of Mexican pines as an adaptation to climate change. *Ecosphere*, *6*(1). DOI: <https://doi.org/10.1890/ES14-00375.1>
- Castellanos-Acuña, D., Lindig-Cisneros, R. A., Silva-Farias, M. Á., & Sáenz-Romero, C. (2014). Zonificación altitudinal provisional de *Abies religiosa* en un área cercana a la Reserva de la Biósfera de la Mariposa Monarca, Michoacán. *Revista Chapingo serie ciencias forestales y del ambiente*, *20*(2), 215-225.
- CONAFOR. (2012). *National Forest inventory and soils 2004-2009. Protocolo de muestreo de campo*.
- Corona-Núñez, R. O., Li, F., & Campo, J. E. (2020). Fires Represent an Important Source of Carbon Emissions in Mexico. *Global Biogeochemical Cycles*, *34*(12), e2020GB006815. DOI: <https://doi.org/10.1029/2020GB006815>
- Corona-Núñez, R. O., Mendoza-Ponce, A., & López-Martínez, R. (2017). Model selection changes the spatial heterogeneity and total potential carbon in a tropical dry forest. *Forest Ecology and Management*, *405*, 69-80. DOI: <https://doi.org/10.1016/j.foreco.2017.09.018>
- Cruz-Cárdenas, G., López-Mata, L., Silva, J. T., Bernal-Santana, N., Estrada-Godoy, F., & López-Sandoval, J. A. (2016). Modelado de la distribución potencial de especies de Pinaceae bajo escenarios de cambio climático en Michoacán. *Revista Chapingo serie ciencias forestales y del ambiente*, *22*(2), 135-148.
- Cueva, E., Lozano, D., & Yaguana, C. (2019). Efecto de la gradiente altitudinal sobre la composición florística, estructura y biomasa arbórea del bosque seco andino, Loja, Ecuador. *Bosque (Valdivia)*, *40*, 365-378. DOI: <https://doi.org/10.4067/S0717-92002019000300365>
- D'Amato, A. W., & Palik, B. J. (2021). Building on the last "new" thing: exploring the compatibility of ecological and adaptation silviculture. *Canadian Journal of Forest Research*, *51*(2), 172-180.
- de Jesús Ordóñez, M. (2000). El territorio del estado de Oaxaca: una revisión. *Investigaciones geográficas*, *42*, 67-86.
- de Jesús Ordóñez, M., & Rodríguez, P. (2008). Oaxaca, el estado con mayor diversidad biológica y cultural en México, y sus productores rurales. *Ciencias*(091).
- Dumroese, R. K., Williams, M. I., Stanturf, J. A., & Clair, J. B. S. (2015). Considerations for restoring temperate forests of tomorrow: forest restoration, assisted migration, and bioengineering. *New Forests*, *46*(5), 947-964. DOI: <https://doi.org/10.1007/s11056-015-9504-6>
- Dyderski, M. K., Paź, S., Frelich, L. E., & Jagodziński, A. M. (2018). How much does climate change threaten European forest tree species distributions? *Global Change Biology*, *24*(3), 1150-1163.
- Elith, J., & Leathwick, J. R. (2009). Species Distribution Models: Ecological Explanation and Prediction Across Space and Time. *Annual Review of Ecology, Evolution, and Systematics*, *40*(Volume 40, 2009), 677-697. DOI: <https://doi.org/10.1146/annurev.ecolsys.110308.120159>
- Evans, J. S., Murphy, M. A., Holden, Z. A., & Cushman, S. A. (2011). Modeling Species Distribution and Change Using Random Forest. In C. A. Drew, Y. F. Wiersma, & F. Huettmann (Eds.), *Predictive Species and Habitat Modeling in Landscape Ecology: Concepts and Applications* (pp. 139-159). Springer New York. DOI: https://doi.org/10.1007/978-1-4419-7390-0_8
- Fick, S. E., & Hijmans, R. J. (2017). WorldClim 2: new 1-km spatial resolution climate surfaces for global land areas. *International journal of climatology*, *37*(12), 4302-4315.

- Gernandt, D. S., & Pérez-de la Rosa, J. A. (2014). Biodiversidad de Pinophyta (coníferas) en México. *Revista mexicana de biodiversidad*, 85, 126-133.
- Govaerts, R., Frodin, D. G., & Govaerts, R. (1998). *World checklist and bibliography of fagales (Betulaceae, Corylaceae, Fagaceae and Ticodendraceae)* (Vol. 2). Royal Botanic Gardens Kew.
- Gray, L. K., Gylander, T., Mbogga, M. S., Chen, P.-y., & Hamann, A. (2011). Assisted migration to address climate change: recommendations for aspen reforestation in western Canada. *Ecological Applications*, 21(5), 1591-1603. DOI: <https://doi.org/10.1890/10-1054.1>
- Guisan, A., & Zimmermann, N. E. (2000). Predictive habitat distribution models in ecology. *Ecological modelling*, 135(2-3), 147-186.
- Guzmán-Santiago, J. C., Santos-Posadas, Héctor Manuel De los, Ángeles-Pérez, Gregorio, Vargas-Larreta, Benedicto, Gómez-Cárdenas, Martín, Rodríguez-Ortiz, Gerardo, & Corona-Núñez, Rogelio Omar. (2023). Effect of climate change on the distribution of temperate climate species in Oaxaca, Mexico. *Botanical Sciences*, 102(1), 39-53. DOI: <https://doi.org/10.17129/botsci.3355>
- Hao, Z.-Z., Liu, Y.-Y., Nazaire, M., Wei, X.-X., & Wang, X.-Q. (2015). Molecular phylogenetics and evolutionary history of sect. *Quinquefoliae* (Pinus): Implications for Northern Hemisphere biogeography. *Molecular Phylogenetics and Evolution*, 87, 65-79.
- iNaturalist (2025). <https://www.inaturalist.org>
- Jiménez-Valverde, A., & Lobo, J. M. (2007). Threshold criteria for conversion of probability of species presence to either–or presence–absence. *Acta Oecologica*, 31(3), 361-369. DOI: <https://doi.org/10.1016/j.actao.2007.02.001>
- Keppel, G., Stralberg, D., Morelli, T. L., & Bátori, Z. (2024). Managing climate-change refugia to prevent extinctions. *Trends in Ecology & Evolution*, 39(9), 800-808. DOI: <https://doi.org/10.1016/j.tree.2024.05.002>
- Keppel, G., Van Niel, K. P., Wardell-Johnson, G. W., Yates, C. J., Byrne, M., Mucina, L., Schut, A. G. T., Hopper, S. D., & Franklin, S. E. (2012). Refugia: identifying and understanding safe havens for biodiversity under climate change. *Global Ecology and Biogeography*, 21(4), 393-404. DOI: <https://doi.org/10.1111/j.1466-8238.2011.00686.x>
- Li, X., Li, Y., Sylvester, S. P., Zang, M., El-Kassaby, Y. A., & Fang, Y. (2021). Evolutionary patterns of nucleotide substitution rates in plastid genomes of *Quercus*. *Ecology and Evolution*, 11(19), 13401-13414.
- Lozano-García, M. d. S., Caballero, M., Ortega, B., Rodríguez, A., & Sosa, S. (2007). Tracing the effects of the Little Ice Age in the tropical lowlands of eastern Mesoamerica. *Proceedings of the National Academy of Sciences*, 104(41), 16200-16203.
- Manzanilla-Quijada, G. E., Treviño-Garza, E. J., Aguirre-Calderón, O. A., Yerena-Yamallel, J. I., & Manzanilla-Quiñones, U. (2020). Current and future potential distribution and identification of suitable areas for the conservation of *Cedrela odorata* L. in the Yucatan Peninsula. *Revista Chapingo serie ciencias forestales y del ambiente*, 26(3), 391-408.
- Mendoza-Ponce, A., Corona-Núñez, R., Kraxner, F., Leduc, S., & Patrizio, P. (2018). Identifying effects of land use cover changes and climate change on terrestrial ecosystems and carbon stocks in Mexico. *Global Environmental Change*, 53, 12-23.
- Mendoza-Ponce, A. V., Corona-Núñez, R. O., Kraxner, F., & Estrada, F. (2020). Spatial prioritization for biodiversity conservation in a megadiverse country. *Anthropocene*, 32, 100267.
- Metcalfe, S. E., Jones, M. D., Davies, S. J., Noren, A., & MacKenzie, A. (2010). Climate variability over the last two millennia in the North American Monsoon region, recorded in laminated lake sediments from Laguna de Juanacatlán, Mexico. *The Holocene*, 20(8), 1195-1206.
- Millar, C. I., & Stephenson, N. L. (2015). Temperate forest health in an era of emerging megadisturbance. *Science*, 349(6250), 823-826. DOI: <https://doi.org/10.1126/science.aaa9933>
- Mirhashemi, H., Heydari, M., Ahmadi, K., Karami, O., Kavgaci, A., Matsui, T., & Heung, B. (2023). Species distribution models of Brant's oak (*Quercus brantii* Lindl.): The impact of spatial database on predicting the impacts of climate change. *Ecological Engineering*, 194, 107038.
- Moeslund, J. E., Arge, L., Bøcher, P. K., Dalgaard, T., & Svenning, J. C. (2013). Topography as a driver of local terrestrial vascular plant diversity patterns. *Nordic Journal of Botany*, 31(2), 129-144.
- Morelli, T. L., Daly, C., Dobrowski, S. Z., Dulen, D. M., Ebersole, J. L., Jackson, S. T., Lundquist, J. D., Millar, C. I., Maher, S. P., Monahan, W. B., Nydick, K. R., Redmond, K. T., Sawyer, S. C., Stock, S., & Beissinger, S. R. (2016). Managing Climate Change Refugia for Climate Adaptation. *PLOS ONE*, 11(8), e0159909. DOI: <https://doi.org/10.1371/journal.pone.0159909>
- Murga-Orrillo, H., Coronado Jorge, M. F., Abanto-Rodríguez, C., & Almeida Lobo, F. D. (2021). Altitudinal gradient and its influence on the edofoclimatic characteristics of tropical forests. *Madera y bosques*, 27(3).

- NASA-JPL. (2013). *NASA Shuttle Radar Topography Mission Global 1 arc second*. DOI: <https://doi.org/10.5067/MEASURES/SRTM/SRTMGL1.003>
- O'Neill, B. C., Tebaldi, C., van Vuuren, D. P., Eyring, V., Friedlingstein, P., Hurtt, G., Knutti, R., Kriegler, E., Lamarque, J. F., Lowe, J., Meehl, G. A., Moss, R., Riahi, K., & Sanderson, B. M. (2016). The Scenario Model Intercomparison Project (ScenarioMIP) for CMIP6. *Geosci. Model Dev.*, *9*(9), 3461-3482. DOI: <https://doi.org/10.5194/gmd-9-3461-2016>
- Ortega-Rosas, C., Peñalba, M., & Guiot, J. (2008). Holocene altitudinal shifts in vegetation belts and environmental changes in the Sierra Madre Occidental, Northwestern Mexico, based on modern and fossil pollen data. *Review of Palaeobotany and Palynology*, *151*(1-2), 1-20.
- Peterson, A. T., Soberón, J., Pearson, R. G., Anderson, R. P., Martínez-Meyer, E., Nakamura, M., & Araújo, M. B. (2012). Princeton University Press. DOI: <https://doi.org/10.1515/9781400840670>
- Phillips, S. J., Dudík, M., & Schapire, R. E. (2004). A maximum entropy approach to species distribution modeling. Proceedings of the twenty-first international conference on Machine learning,
- R-Core-Team. (2018). *R: A language and environment for statistical Computing*. In R Foundation for Statistical Computing.
- Rehfeldt, G. E., Leites, L. P., Bradley St Clair, J., Jaquish, B. C., Sáenz-Romero, C., López-Upton, J., & Joyce, D. G. (2014). Comparative genetic responses to climate in the varieties of *Pinus ponderosa* and *Pseudotsuga menziesii*: Clines in growth potential. *Forest Ecology and Management*, *324*, 138-146. DOI: <https://doi.org/10.1016/j.foreco.2014.02.041>
- Rivera, E. R., & Amador, J. A. (2008). Predicción estacional del clima en Centroamérica mediante la reducción de escala dinámica. Parte I: Evaluación de los modelos de circulación general CCM3. 6 y ECHAM4. 5. *Revista de matemática: teoría y aplicaciones*, *15*(2), 131-173.
- Rodríguez-Ramírez, A., Caballero, M., Roy, P., Ortega, B., Vázquez-Castro, G., & Lozano-García, S. (2015). Climatic variability and human impact during the last 2000 years in western Mesoamerica: evidence of late Classic (AD 600–900) and Little Ice Age drought events. *Climate of the Past*, *11*(9), 1239-1248.
- Safaei, M., Rezayan, H., Zeaiean Firouzabadi, P., & Sadidi, J. (2021). Optimization of species distribution models using a genetic algorithm for simulating climate change effects on Zagros forests in Iran. *Ecological Informatics*, *63*, 101288. DOI: <https://doi.org/10.1016/j.ecoinf.2021.101288>
- Stockwell, D. (1999). The GARP modelling system: problems and solutions to automated spatial prediction. *International journal of geographical information science*, *13*(2), 143-158.
- Swets, J. A. (1988). Measuring the accuracy of diagnostic systems. *Science*, *240*(4857), 1285-1293.
- Trejo, I., Martínez-Meyer, E., Calixto-Pérez, E., Sánchez-Colón, S., Vázquez De La Torre, R., & Villers-Ruiz, L. (2011). Analysis of the effects of climate change on plant communities and mammals in México. *Atmósfera*, *24*(1), 1-14.
- Valencia, S. (2004). Diversidad del género *Quercus* (Fagaceae) en México. *Boletín de la sociedad Botánica de México*(75), 33-53.
- Voldoire, A. (2023). IPCC DDC: CNRM-CERFACS CNRM-CM6-1 model output prepared for CMIP6 ScenarioMIP. *World Data Center for Climate (WDCC) at DKRZ*. DOI: <https://doi.org/10.26050/WDCC/AR6.C6SPCECC1>
- Wisz, M. S., Hijmans, R., Li, J., Peterson, A. T., Graham, C., Guisan, A., & Group, N. P. S. D. W. (2008). Effects of sample size on the performance of species distribution models. *Diversity and distributions*, *14*(5), 763-773.
- Zeb, U., Wang, X., AzizUllah, A., Fiaz, S., Khan, H., Ullah, S., Ali, H., & Shahzad, K. (2022). Comparative genome sequence and phylogenetic analysis of chloroplast for evolutionary relationship among *Pinus* species. *Saudi Journal of Biological Sciences*, *29*(3), 1618-1627.

Supporting information

Table S1. Records of presence of the evaluated species.

Species	IF_SJQ	INFyS Oaxaca	GBIF	iNaturalist	Records
<i>Arbutus xalapensis</i> KUNTH	14	144	57	87	302
<i>Clethra mexicana</i> DC	41	57	6	4	108
<i>Pinus devoniana</i> Lindl.	11	31	8	7	57
<i>Pinus oocarpa</i> Schiede ex Schltdl.	63	78	11	20	172
<i>Pinus teocote</i> Schiede ex Schltdl. & Cham.	24	23	0	20	67
<i>Quercus acutifolia</i> f. <i>abrupta</i> Trel.	47	51	11	9	118
<i>Quercus castanea</i> Née	13	32	25	24	94
<i>Quercus crassifolia</i> Benth.	17	61	0	24	102
<i>Quercus elliptica</i> Newberry	39	37	0	5	81
<i>Quercus magnoliifolia</i> Née	14	86	14	15	129
<i>Quercus rugosa</i> Née	13	54	24	62	153
Total	296	654	156	277	1,383

Table S2. Environmental variables used for modelling the potential distribution of species.

Key	Description (Unit)
Bio-1	Annual Mean Temperature (°C)
Bio-2	Mean Diurnal Range (Mean of monthly (max temp - min temp)) (°C)
Bio-3	Isothermality (BIO2/BIO7) (×100) (°C)
Bio-4	Temperature seasonality (standard deviation × 100) (°C)
Bio-5	Max Temperature of Warmest Month (°C)
Bio-6	Min Temperature of Coldest Month (°C)
Bio-7	Temperature Annual Range (BIO5-BIO6) (°C)
Bio-8	Mean Temperature of Wettest Quarter (°C)
Bio-9	Mean Temperature of Driest Quarter (°C)
Bio-10	Mean Temperature of Warmest Quarter (°C)
Bio-11	Mean Temperature of Coldest Quarter (°C)
Bio-12	Annual precipitation (mm)
Bio-13	Precipitation of Wettest Month (mm)
Bio-14	Precipitation of Driest Month (mm)
Bio-15	Precipitation Seasonality (Coefficient of Variation) (%)
Bio-16	Precipitation of Wettest Quarter (mm)
Bio-17	Precipitation of Driest Quarter (mm)
Bio-18	Precipitation of Warmest Quarter (mm)
Bio-19	Precipitation of Coldest Quarter (mm)

Table S3. Species adaptation threshold.

Species	Threshold	Accuracy	Sensitivity
<i>Arbutus xalapensis</i>	0.53	0.86	0.97
<i>Clethra mexicana</i>	0.42	0.81	0.62
<i>Pinus devoniana</i>	0.46	0.88	0.98
<i>Pinus oocarpa</i>	0.50	0.91	0.98
<i>Pinus teocote</i>	0.56	0.91	0.97
<i>Quercus acutifolia</i>	0.54	0.92	0.97
<i>Quercus castanea</i>	0.53	0.87	0.95
<i>Quercus crassifolia</i>	0.46	0.86	0.99
<i>Quercus elliptica</i>	0.54	0.96	0.95
<i>Quercus magnoliifolia</i>	0.64	0.92	0.89
<i>Quercus rugosa</i>	0.58	0.92	0.95

Table S4. Spatial attributes of studied species across climate change scenarios.

Species	Scenario	Altitude (m a.s.l.)	Latitude (°)	Slope (°)
<i>Arbutus xalapensis</i>	Current	1,107±824	16.967±0.637	5.8±4.7
	ssp126-2040	1,108±824	16.967±0.637	5.8±4.7
	ssp585-2040	1,108±824	16.967±0.637	5.8±4.7
	ssp126-2100	1,108±824	16.967±0.637	5.8±4.7
	ssp585-2100	1,108±824	16.967±0.637	5.8±4.7
<i>Clethra mexicana</i>	Current	939±685	17.342±1.030	7.2±5.3
	ssp126-2040	963±697	17.349±1.037	7.4±5.5
	ssp585-2040	982±694	17.346±1.036	7.2±5.4
	ssp126-2100	978±716	17.358±1.036	7.0±5.4
	ssp585-2100	979±885	17.393±1.044	7.0±4.6
<i>Pinus devoniana</i>	Current	897±770	17.346±1.047	6.4±5.1
	ssp126-2040	934±816	17.359±1.045	6.7 ±5.3
	ssp585-2040	924±808	17.359±1.046	6.8±5.3
	ssp126-2100	950±844	17.373±1.047	6.5±5.6
	ssp585-2100	989±827	17.398±1.045	6.0±5.1
<i>Pinus oocarpa</i>	Current	1,034±856	17.314±0.988	5.4±5.1
	ssp126-2040	977±921	17.321±1.000	5.0±5.1
	ssp585-2040	960±915	17.317±1.001	5.0±5.0
	ssp126-2100	870±922	17.321±1.003	4.6±4.9
	ssp585-2100	985±735	17.308±1.007	4.0±4.1
<i>Pinus teocote</i>	Current	1,123±952	17.364±1.030	5.7±4.9
	ssp126-2040	1,105±998	17.375±1.032	5.3±4.9
	ssp585-2040	1,079±985	17.374±1.033	5.3±4.9
	ssp126-2100	1,026±1012	17.380±1.035	4.7±4.8
	ssp585-2100	1,086±1057	17.400±1.045	4.0±3.8
<i>Quercus acutifolia</i>	Current	1,420±878	17.344±1.023	7.6±5.3
	ssp126-2040	1,350±924	17.359±1.034	8.0±5.8
	ssp585-2040	1,343±930	17.357±1.034	7.9±5.7
	ssp126-2100	1,303±969	17.367±1.037	7.8±5.7
	ssp585-2100	1,341±1,178	17.393±1.039	6.3±5.9
<i>Quercus castanea</i>	Current	1,444±1,002	17.347±1.011	6.0±4.7
	ssp126-2040	1,306±1,084	17.340±1.019	5.33±4.7
	ssp585-2040	1,199±1,085	17.335±1.023	5.0±4.6
	ssp126-2100	1,220±1,117	17.343±1.026	4.9±4.6
	ssp585-2100	1,187±1,061	17.374±1.041	3.0±3.4

Continue

Table S4. Continued

<i>Quercus crassifolia</i>	Current	1,642±947	17.352±1.003	7.5±4.9
	ssp126-2040	1,543±1,034	17.364±1.008	6.9±4.9
	ssp585-2040	1,493±1,054	17.368±1.006	6.8±4.9
	ssp126-2100	1,398±1,094	17.371±1.006	6.4±5.0
	ssp585-2100	1,496±1,119	17.391±1.016	5.1±5.0
<i>Quercus elliptica</i>	Current	1,056±749	17.364±1.032	8.3±5.5
	ssp126-2040	1,003±801	17.375±1.037	8.1±5.8
	ssp585-2040	994±783	17.373±1.036	7.9±5.7
	ssp126-2100	970±803	17.383±1.037	7.8±5.9
	ssp585-2100	908±835	17.405±1.038	6.5±6.8
<i>Quercus magnoliifolia</i>	Current	1,305±778	17.307±1.009	6.6±4.4
	ssp126-2040	1,221±871	17.310±1.013	6.1±4.6
	ssp585-2040	1,236±858	17.312±1.015	6.1±4.6
	ssp126-2100	1,183±904	17.319±1.017	5.7±4.6
	ssp585-2100	1,291±866	17.283±1.045	4.7±4.2
<i>Quercus rugosa</i>	Current	1,731±856	17.331±0.974	6.8±4.5
	ssp126-2040	1,615±976	17.354±0.980	6.6±4.6
	ssp585-2040	1,579±996	17.360±0.982	6.7±4.7
	ssp126-2100	1,417±1,067	17.372±0.983	6.4±4.8
	ssp585-2100	1,655±760	17.382±1.007	5.5±5.0

Table S5. Distribution percentages presented by forest species in Gain, Loss and Permanence (Per). Gain refers to the percentage increase in area with respect to the current potential distribution. The sum of Loss and Per corresponds to 100% in relation to the current potential distribution.

Species	SSP126-2040			SSP126-2100			SSP585-2040			SSP585-2100		
	Gain	Loss	Per	Gain	Loss	Per	Gain	Loss	Per	Gain	Loss	Per
<i>Arbutus xalapensis</i>	9.0	25.5	74.5	11.1	33.8	66.2	9.2	25.9	74.1	1.4	78.4	21.6
<i>Clethra mexicana</i>	6.2	22.2	77.8	5.4	36.7	63.3	5.8	24.2	75.8	1.6	85.9	14.1
<i>Pinus devoniana</i>	16.9	37.0	63.0	18.5	55.5	44.5	16.5	38.6	61.4	17.1	95.2	4.8
<i>Pinus oocarpa</i>	16.6	48.3	51.7	17.6	62.8	37.2	17.4	50.7	49.3	20.5	93.5	6.5
<i>Pinus teocote</i>	22.0	32.2	67.8	33.7	43.3	56.7	22.1	32.6	67.4	13.4	87.8	12.2
<i>Quercus acutifolia</i>	3.1	28.2	71.8	3.1	40.8	59.2	2.8	30.6	69.4	0.3	91.8	8.2
<i>Quercus castanea</i>	5.4	22.9	77.1	5.0	32.9	67.1	4.5	23.7	76.3	2.3	78.6	21.4
<i>Quercus crassifolia</i>	12.8	36.5	63.5	15.7	46.8	53.2	13.9	36.6	63.4	14.3	94.6	5.4
<i>Quercus elliptica</i>	10.0	30.5	69.5	13.3	43.6	56.4	10.0	32.1	67.9	13.5	93.2	6.8
<i>Quercus magnoliifolia</i>	11.0	26.0	74.0	10.6	36.4	63.6	9.2	28.0	72.0	3.5	84.1	15.9
<i>Quercus rugosa</i>	6.3	35.5	64.5	5.5	51.2	48.8	7.2	36.1	63.9	6.1	85.6	14.4