

# Forecasting the flooding dynamics of flatwoods salamander breeding wetlands under future climate change scenarios

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# Abstract

Ephemeral wetlands are globally important systems that are regulated by regular cycles of wetting and drying, which are primarily controlled by responses to relatively short-term weather events (e.g., precipitation and evapotranspiration). Climate change is predicted to have significant effects on many ephemeral wetland systems and the organisms that depend on them through altered filling or drying dates that impact hydroperiod. To examine the potential effects of climate change on pine flatwoods wetlands in the southeastern United States, we created statistical models describing wetland hydrologic regime using an approximately 8-year history of water level monitoring and a variety of climate data inputs. We then assessed how hydrology may change in the future by projecting models forward (2025–2100) under six future climate scenarios (three climate models each with two emission scenarios). We used the model results to assess future breeding conditions for the imperiled Reticulated Flatwoods Salamander (*Ambystoma bishopi*), which breeds in many of the study wetlands. We found that models generally fit the data well and had good predictability across both training and testing data. Across all models and climate scenarios, there was substantial variation in the predicted suitability for flatwoods salamander reproduction. However, wetlands with longer hydroperiods tended to have fewer model iterations that predicted at least five consecutive years of reproductive failure (an important metric for population persistence). Understanding potential future risk to flatwoods salamander populations can be used to guide conservation and management actions for this imperiled species.

**Keywords:** *Ambystoma bishopi*, Amphibians, Conservation, Ephemeral wetlands, Hydrology, Management, Water level monitoring

# Introduction

Ephemeral wetlands are globally important ecosystems, contributing to landscape scale hydrological processes and nutrient cycles (McLaughlin, Kaplan & Cohen, 2014; Capps, Berven & Tiegs, 2015; Cohen et al., 2016) and providing critical habitat for diverse and productive communities (Galatowitsch & van der Valk, 1996; Kirkman et al., 1999; Jenkins, Grissom & Miller, 2001; Gibbons et al., 2006; Hunter & Lechner, 2017). Yet, ephemeral wetlands are imperiled, with an estimated global loss of approximately 50% and accelerated loss in tropical and subtropical areas and in some arid regions (Moser et al., 1996; OECD, 1996; Davidson, 2014). In the United States, wetland area has been reduced by approximately 50% since the 1780s, with losses along the coast of the Gulf of Mexico constituting 80% of all wetland losses (Dahl, 1990). While habitat change, especially conversion to agriculture, accounts for the majority of wetland loss (Frayer et al., 1983), climate change represents a future threat to ephemeral wetlands via increases in temperature and changes in precipitation patterns (Dahl, 2011, Zhu et al., 2017). Efforts to protect and restore ephemeral wetlands, therefore, will require an understanding of how climate change may affect wetland hydrologic regimes and associated functions (Erwin, 2009).

In the southeastern United States, climate projections suggest that there will be increased drought severity and frequency (Karl, Melillo & Peterson, 2009), increased rates of evapotranspiration through higher temperatures (Sun et al., 2002; Ingram et al., 2013), and a greater amount of total rainfall that arrives either less predictably or at different times of the year (Mulholland et al., 1997; Karl, Melillo & Peterson, 2009; Ingram et al., 2013). All of these climatic shifts, coupled with features of the local environment and broader landscape (Hayashi & Rosenberry, 2002; Winter & LaBaugh, 2003; Lu et al., 2009; Chandler et al., 2017; Jones et al.,

2018), have the potential to alter wetland hydrologic regimes in the region, with cascading effects on water quality, carbon storage, and wildlife populations (Zedler & Kercher, 2005; Davis et al., 2019).

The southeastern United States is a hotspot for amphibian diversity, with many species reliant on ephemeral wetlands to complete their life cycle (Blaustein et al., 2010; Walls et al., 2013, Greenberg et al., 2015). Amphibians often favor ephemeral wetlands for breeding habitat owing to increased productivity in the absence of predatory fish populations (Skelly, 1997). However, reproductive failure resulting from wetland drying is common, and altered hydrologic regimes may permanently erase the conditions necessary to support successful breeding cycles (Brooks, 2009; Yang & Rudolf, 2010; Benard, 2015). As such, amphibians are of particular concern when assessing the vulnerability of different taxa to climate-induced shifts in wetland hydrologic regime (Pounds, 2001; Carey & Alexander, 2003; Blaustein et al., 2010; Walls et al., 2013, Greenberg et al., 2015). In the context of managing imperiled species, failure to take such habitat alterations into account may lead to overly optimistic persistence probabilities derived from population viability analyses or present unexpected challenges to otherwise successful recovery efforts.

Here, we develop a hydrologic model for ephemeral wetlands used as breeding habitat by the federally endangered Reticulated Flatwoods Salamander (*Ambystoma bishopi*; USFWS, 2020). Flatwoods salamanders have declined throughout their range due to widespread loss of their pine flatwoods habitat to urbanization, agriculture, and plantation forestry and the degradation of remaining suitable breeding wetlands, mainly through fire suppression and exclusion (Bishop & Haas, 2005; Palis & Hammerson, 2008; O'Donnell et al., 2017). This degradation of wetland habitat can lead to a variety of conditions that are unsuitable for larval

growth and development (e.g., reduced hydroperiod and low dissolved oxygen levels; de Szalay & Resh, 1997; Skelly, Freidenburg & Kiesecker, 2002; Huxman et al., 2005; Gorman, Bishop & Haas, 2009; Sacerdote & King, 2009; Shulse et al., 2012; Jones et al., 2018). While vegetation management and prescribed fire application have improved some of these issues, there is substantial concern that climate change will continue to negatively impact flatwoods salamander populations, even on protected landscapes (Chandler et al., 2016; USFWS, 2020). Using water level data from 35 wetlands within the Gulf Coastal Plain of the southeastern U.S., we sought to predict hydrologic regimes across a series of climate projections to understand future suitability of study wetlands for larval flatwoods salamander survival and metamorphosis. Our research objectives were as follows: 1) model present-day dynamics of ephemeral wetlands, 2) project wetland hydrologic regimes onto future (2025–2100) climate space, and 3) evaluate the impact of future hydrologic regimes on flatwoods salamander breeding potential. Our findings shed light on the key drivers of hydrologic dynamics in the study region and carry implications for flatwoods salamander recovery efforts.

## Materials & Methods

### *Study Sites*

We conducted our work on Eglin Air Force Base (Eglin) in the Florida panhandle. Access to field sites was approved by the US Fish and Wildlife Service and Jackson Guard (Eglin’s Natural Resources Division; Cooperative Agreement Number F14AC00068). Eglin is a large military installation (>187,000 ha) located in Okaloosa, Walton, and Santa Rosa Counties and consists primarily of actively managed longleaf pine forests. The wetlands studied here have been the subject of ongoing research to understand the variation in hydrologic regime (e.g.,

Chandler et al., 2016; Chandler et al., 2017) across the landscape with the goal of managing and conserving valuable breeding habitat for flatwoods salamanders (see USFWS, 2009 for additional study site details). Extensive habitat management in both uplands and wetlands has made Eglin one of the few remaining strongholds for flatwoods salamanders (Gorman, Haas & Hines, 2013; USFWS, 2020).

# *Water level data*

As part of our long-term work on flatwoods salamanders, we previously installed water level monitoring wells at the approximate deepest point in 35 wetland basins on Eglin. All wetlands were located within 20 km of one another, and wetland areas ranged from 0.19–20.92 ha, although all but two wetlands were less than six hectares. Wells were installed at varying points in time (November 2014 through December 2017) such that we had between 1,061 and 2,338 daily measures of water level for each well (see Table S1 for more details). Each well consisted of a 3.8 diameter screened PVC pipe 1 m below ground with a HOBO U20 pressure transducer (Onset Computer Corporation, Bourne, MA) at the bottom of each well that recorded total pressure and temperature at 15-minute intervals. Total pressure data were corrected for barometric pressure variation using pressure sensors installed following methods in McLaughlin & Cohen (2011) at two locations within 9 km of any given wetland (Table S1). For each wetland, we used the resulting 15-min water level data to calculate mean daily water level (mm; positive indicating water levels above ground surface), along with the timing and duration of flooded conditions.

# *Current climate data*

To determine precipitation inputs that each wetland received, we installed four rain gauges (HOBO Data Logging Rain Gauge RG3-M) such that the distance between any of the 35 wells and a rain gauge was no more than 4 km (Table S1). Three of the rain gauges were installed on 20 November 2014 and the fourth was installed on 19 June 2018. We used data from these gauges to calculate daily rainfall (mm), cumulative rainfall over the previous seven days (mm), and the number of days since the last rain event.

We obtained daily minimum and maximum temperature ( $^{\circ}\text{C}$ ) data from the Florida Automated Weather Network (FAWN) over the course of our study period using the Jay weather station (Station ID 110), which is located 32–58 km from each wetland. We calculated daily potential evapotranspiration (PET, mm) from minimum and maximum temperatures using the Hargreaves-Samani method (Hargreaves & Samani 1985). Lastly, we calculated the 12-month standardized precipitation-evapotranspiration index (SPEI) for each well from 1 January 1981 to 31 May 2022. The SPEI is useful for assessing drought severity (WMO, 2006) and quantifying and comparing water balances across locations (Stagge et al., 2014). To calculate monthly SPEI, we used daily minimum and maximum temperature data obtained from PRISM Climate group (Oregon State University, <http://prism.oregonstate.edu>, created 26 Oct 2022), which were necessary because the FAWN data do not have a long enough time series for the SPEI calculation. For the same time period, we used the Hargreaves method (Hargreaves, 1994) to calculate monthly PET, which was used for calculating SPEI.

### *Hydrologic Model*

We developed a statistical model to predict daily water levels at each of the 35 wells using a Bayesian first order autoregressive fixed effect model, which was based on visual



inspection of partial autocorrelation plots for each well. For each wetland basin, the water level of a given day was modeled as a function of the first order autoregressive term, the amount of rain received during the previous day, and their interaction. Models also included linear and quadratic terms for PET, the sum of the rain received in the previous seven days, and SPEI. We also included interactions between SPEI and the amount of rain received during the previous day, and the interaction between SPEI, the first order autoregressive term, and the amount of rain received during the previous day. Therefore, for each of the 35 well-specific models, we estimated 11 parameters, including an error term.

We selected these variables to model hydroregimes because of typically strong ephemeral wetland responses to regional climate forcing (Brooks, 2004; Greenberg et al., 2015; Lee et al., 2015). We defined water inputs through direct precipitation effects and parameters that included precipitation within an interaction. The interaction between precipitation and the autoregressive term is useful for defining stage-dependent inputs of rainfall. We included linear and quadratic terms for PET to define short-term (i.e., daily) drawdown of water caused by evapotranspiration because this relationship was nonlinear at low PET values. Lastly, we included SPEI and parameters that include SPEI within an interaction to define wetland response to long-term drought conditions.

For each of the 35 well-specific models, we assigned vague priors following a normal distribution (mean = 0; variance = 100) to all fixed effects, with the error term assumed to follow a normal distribution with mean = 0 and the prior of variance to follow a uniform distribution (min = 0, max = 100). We fit each model using Markov chain Monte Carlo (MCMC) simulations, generating three chains, each with 400,000 total iterations and a thinning rate of 100 (Kéry & Royle, 2016). We used an adaptation phase of 1,000 and discarded 300,000 burn-in

iterations, which retained 2,000 iterations for each chain (6,000 total samples) to estimate posterior distributions. We examined traceplots of parameters for adequate mixing among chains and the Gelman-Rubin's  $\hat{R}$  statistic to evaluate model convergence (Gelman, 2004). We assessed model predictive ability using a posterior predictive check based on the Bayesian P-value (Kéry, 2010; Link & Barker 2010). We evaluated parameter significance based on the overlap of 95% highest posterior density with zero.

### *Model Validation*

For each well-specific model, we used 75% of the available data to train the model and 25% for testing. Because models were autoregressive, rather than using random data points, the training dataset included consecutive data starting at a random position within the first 25% of the data and consisted of 75% of these data while the testing dataset consisted of the remaining 25% of the data. To determine how well each model predicted the measured water level of our testing and training datasets, we calculated the normalized root mean squared error (NRMSE) for each of the 6,000 iterations, which is the root mean squared error divided by the range of measured water levels within each basin to account for differences in scale among basins. To determine how well each model predicted flooded conditions, we calculated the proportion of days in our testing and training dataset, for each iteration, that were predicted correctly to either have standing water or not.

### *Predicting hydrologic regime from future climate data*

We obtained downscaled climate data for the years 2025–2100 using Localized Constructed Analogs (LOCA; Pierce, Cayan & Thrasher, 2014), which have a 6 x 6 km

resolution. For each well, we obtained daily estimates of precipitation (mm), minimum temperature (°C), and maximum temperature (°C) for each of three global circulation models (GCM): Hadley Centre Global Environment Model 2 Earth Systems (HadGEM2-ES), Hadley Centre Global Environment Model 2 Carbon Cycle (HadGEM2-CC), and the Community Climate System model version 4 (CCSM4). For each model, we used two different projections for representative concentration pathways (RCP): 4.5 assumes a peak in carbon emissions in 2040 (Thomson et al., 2011), and 8.5 assumes emissions will continue to increase throughout the 21st century (Riahi et al., 2011). Therefore, for each model, we were able to assess future wetland hydrologic regimes under six different climate scenarios. For each of the six scenarios, we used the daily temperature and precipitation variables to calculate the same climate metrics (i.e., PET, SPEI, sum of rain over the last seven days) as the current data and predicted daily water level for each well using model parameter estimates from their respective posterior distributions.

To predict future hydrologic regimes and associated habitat suitability for flatwoods salamanders, we combined the LOCA climate data and the parameter posterior distributions from our statistical models. For each breeding season from 2025–2100, we determined the length (number of days) of the maximum hydroperiod within a given season, as well as the day in which each wetland became flooded. We defined the flatwoods salamander breeding season as occurring between 1 November and 31 May of the following year (Palis, 1996). Additionally, within a given posterior draw, we determined the number of consecutive breeding seasons that did not contain a hydroperiod of at least 15 weeks. While metamorphosis in flatwoods salamanders has occurred in as little as 11 weeks (Palis, 1995), 15 weeks is a conservative estimate for the typical length of wetland filling required to allow larval development through

metamorphosis (Brooks et al., 2020; Haas, unpublished data). Lastly, for each of the maximum hydroperiod estimates, we determined the date the wetland filled as the number of days since 1 November for the respective breeding season.

All analyses were performed in program R (version 4.1.1; R Core Team, 2021). We used the *SPEI* (Beguería & Vicente-Serrano 2017) and *Evapotranspiration* (Guo, Westra & Peteron, 2019) packages for calculating SPEI and PET respectively, and the *jagsUI* package (Kellner, 2017) to call JAGS (Plummer, 2003), from Program R for MCMC analyses.

## Results

### *Model fitting*

Our dataset consisted of 72,266 daily observations of wetland water level across 35 monitoring wells (range: 795–1,753 training observations per well; Table S1). These data covered a range of hydrologic conditions, both among wetlands (e.g., mean across wetland hydroperiod during the flatwoods salamander breeding season ranged from 37–142 days) and across years (e.g., mean annual breeding season hydroperiod across all wetlands ranged from 31–168 days). All MCMC chains showed good mixing, and  $\hat{R}$  values were between 1.000 and 1.003, indicating model convergence. The posterior predictive check indicated that each model fit the data well (Bayesian P-value range: 0.49–0.51).

Our models generally showed good predictability of water level data, with the median training NRMSE ranging from 0.01 to 0.04 and the testing NRMSE ranging from 0.13 to 0.35 (Figure 1A, C). Variance in NRMSE was always  $< 0.01$  for training and testing datasets and generally showed a decreasing trend with the number of training or testing datapoints (Figure 1B, D). The median proportion of days correctly predicted to either have or not have surface

water ranged from 0.94–0.99 for training datasets and from 0.56–0.93 for testing datasets (Figure 1E, G). The variance in proportion of days correctly predicted to have surface water present ranged from  $< 0.001$ –0.01 and from 0.12–2.40 for training and testing datasets, respectively (Figure 1F, H).

### *Model Parameters*

The mean value for the first order autoregressive coefficient was positive and slightly less than one for all wetland basins. We found that daily precipitation had the largest magnitude effect, when compared to all other parameters, on wetland water level (Figure 2; Table S2). The amount of rain over the previous seven days had a smaller positive effect than daily rainfall on water level. The model intercept ( $\alpha$ ), the SPEI, the quadratic term for PET, and the interactions between SPEI and daily rainfall and daily rainfall, autoregressive term, and SPEI had both negative and positive effects depending on the model. Finally, the effects of PET and the interaction between precipitation and the autoregressive term tended to have negative effects on daily water level (Figure 2; Table S2).

### *Future predictions*

Future projections revealed suitable hydroperiods for flatwoods salamander larval development (i.e., at least 15 consecutive weeks of surface water during the breeding season) in at least some years across all climate scenarios (Figure 3 and 4). When comparing the various climate scenarios, the CCSM4 model predictions typically had fewer years with suitable hydroperiods compared to the HadGEM2-CC or HadGEM2-ES model (Figures 3 and 4). For date of wetland filling, wetland basins generally showed a tendency towards earlier fill dates

(Figure 5). For both hydroperiod and date of wetland filling, there was considerable variation among years and iterations for all wetland basins (Figures 3–5). Lastly, across all wetlands and climate scenarios examined in this study, we found substantial variability in their general suitability for flatwoods salamander reproduction (Figure 6). We found that wetlands experiencing longer hydroperiods, on average, were less likely to have model iterations of at least five consecutive years of reproductive failure (a metric thought to be important for population persistence; Palis, Aresco & Kilpatrick., 2006). Some wetlands did have a high probability ( $> 75\%$ ) of extended recruitment failures or had a low number of years with a suitable hydroperiod, indicating they are likely unsuitable for flatwoods salamanders. Overall, most wetland basins had suitable hydroperiods for flatwoods salamanders in at least 50% of future years (Figures 5 and 6).

## Discussion

Here, we present wetland-specific models describing hydrologic regime in ephemeral wetlands embedded within the pine flatwoods ecosystem of the southeastern United States. We show that, by harnessing information contained within multiple years (2014–2022) of daily water level data, models accurately predicted the dynamics of 35 flatwoods salamander breeding wetlands. Forecasting across six climate change scenarios, we demonstrate that wetlands vary in their response to changing climate conditions, with many wetlands exhibiting earlier fill dates in future years. Despite these predicted alterations in hydrologic regime, most sites appear to remain hydrologically suitable for flatwoods salamanders. For the handful of wetlands that may become unsuitable, our results can be used to direct appropriate management practices to mitigate climate-induced changes.

Hydrologic regimes were most strongly determined by rainfall patterns and PET rates. Specifically, daily rainfall had the largest positive influence on wetland water level, and linear relationships with PET had the largest negative effect on wetland water level. It is unsurprising that daily rainfall had a large positive effect on water levels as these wetlands are largely rain-fed, with minimal connections to other water sources (i.e., often referred to as geographically isolated wetlands; Tiner, 2003). Similarly, the effects of vegetation (and thus ET rates), both within and surrounding wetlands, are known to influence groundwater inputs and resulting water levels (Jones et al., 2018). A logical conservation application, therefore, is to devise management strategies designed to artificially manipulate these important determinants of hydrologic regime. Future research should seek to test the effectiveness of shallow groundwater wells or different vegetation management practices (including the restoration of natural fire regimes) in extending wetland hydroperiods (Seigel et al., 2006; Jones et al., 2018).

Our long-term water level data allowed us to accurately model wetland hydrologic regime under a range of both current and future climate conditions. However, accurate predictions were contingent on having multiple years of data. The amount of data necessary to construct robust models likely represents a need to capture different water levels under a wide range of climate conditions. Unsurprisingly, annual climate variation is reflected in hydroperiod as these ephemeral wetlands display markedly different hydroperiods from one year to the next (Chandler et al., 2016). Therefore, a single breeding season or year will likely not capture this variation, especially as precipitation events vary in number and magnitude. Additionally, the magnitude of increases in wetland water level to a given amount of precipitation will change under long-term drought (e.g., low values of SPEI) or wet conditions (high values of SPEI), highlighting the need for extended time series to quantify long-term trends.

Although predictive accuracy varied considerably across the 35 instrumented wetlands, this variability was unrelated to the number of training data points used in model fitting. This suggests two things, 1) even the most recently instrumented sites yielded enough data points to accurately predict their dynamics ( $> 1,000$  total data points), and 2) residual error in model predictions was due to unmeasured variables, as opposed to insufficient data. Characteristics such as area, shape, hydraulic conductivity, and vegetation structure are all expected to affect wetland hydrologic regime and its response to climate forcing (Brooks, 2005; Jones et al., 2018; Cianciolo et al., 2021). For example, the large variation in the degree to which daily rainfall increases water levels at our study wetlands likely results from differences in canopy interception, wetland bathymetry, and storm event surface runoff (Brooks, 2005). Further, larger wetlands have been shown to exhibit lower recession rates and longer hydroperiods compared to smaller wetlands (e.g., Vanschoenwinkel et al., 2009; Chandler et al., 2016; Chandler et al., 2017). It is likely that other factors, such as wetland bathymetry (Haag, Lee & Herndon, 2005) and landscape position, may also affect wetland hydrologic response to climate variables, and as such, incorporating other wetland characteristics could improve the predictive accuracy of these models and provide additional insights into an individual wetland's sensitivity to climate change.

Changes in the timing or duration of wetland filling can negatively impact several aspects of amphibian reproduction (Parmesan, 2006; Li, Cohen & Rohr, 2013). During the fall and early winter, adult flatwoods salamanders migrate to wetlands to lay their eggs in dry wetland basins (Anderson & Williamson 1976; Palis, 1996). Embryos begin to develop terrestrially but do not hatch until inundated by rising water levels. Although eggs can persist in dry basins for up to two months, they risk mortality from desiccation or freezing if exposed for too long (Anderson & Williamson, 1976). After inundation, larvae can take between 11 to 18 weeks to metamorphose



into terrestrial adults (Palis, 1995). Therefore, reproductive success is dependent upon both the timing of when a wetland fills at the beginning of the breeding season as well as the length of time a wetland is flooded. Reassuringly, future projections of hydrological suitability suggest that most of the currently monitored wetlands will remain suitable for flatwoods salamander larval development and metamorphosis. Additionally, the strong influence of PET on water level uncovers a potential management strategy involving the intensive removal of woody vegetation (e.g., Liu et al., 2020). Indeed, many of these wetlands, as they are either currently occupied or potentially occupied by flatwoods salamanders, have been the focus of habitat management in the form of physical removal and chemical treatment (i.e., herbicide and fire) of wetland shrubs (Gorman, Haas & Himes, 2013), presenting an opportunity to quantify the additional benefits of vegetation removal on habitat quality.

## Conclusions

There is a growing body of literature highlighting observed and potential impacts of climate change on amphibian species in the southeastern United States (e.g., Todd et al., 2011; Greenberg et al., 2015; Walls et al., 2019). Our results do not suggest an immediately worrisome scenario for flatwoods salamanders when considering breeding wetland hydrology in the coming decades. However, it is important to acknowledge that population viability is contingent on the timing of breeding migrations in relation to environmental conditions and the survival of adults in upland habitats surrounding breeding wetlands (Brooks, 2020). Only by integrating the wetland models presented here with additional phenological and demographic information can we explicitly model flatwoods salamander persistence under future climate change and guide ongoing recovery efforts. More broadly, our approach can be used to discern the relative

363 vulnerability of ephemeral wetlands in the southeastern United States to climate change and  
364 devise strategies to safeguard the species that rely on them.

365

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# References

- Anderson, J. D., and G. K. Williamson. 1976. Terrestrial mode of reproduction in *Ambystoma cingulatum*. *Herpetologica* 32:214–221.
- Baguería, S., and S. M. Vicente-Serrano. 2017. SPEI: Calculation of the standardized precipitation-evapotranspiration index. R package version 1.7.
- Benard, M. F. 2015. Warmer winters reduce frog fecundity and shift breeding phenology, which consequently alters larval development and metamorphic timing. *Global Change Biology* 21:1058–1065.
- Bishop, D. C., and C. A. Haas. 2005. Burning trends and potential negative impacts on flatwoods salamanders. *Natural Areas Journal* 25:290–294.
- Blaustein, A. R., S. C. Walls, B. A. Bancroft, J. J. Lawler, C. L. Searle, and S. S. Gervasi. 2010. Direct and indirect effects of climate change on amphibian populations. *Diversity* 2:281–313.
- Brooks, R. T. 2004. Weather-related effects on woodland vernal pool hydrology and hydroperiod. *Wetlands* 24:104–114.
- Brooks, R. T. 2005. A review of basin morphology and pool hydrology of isolated ponded wetlands: Implications for seasonal forest pools of the northeastern United States. *Wetlands Ecology and Management* 13:335–348.
- Brooks, R. T. 2009. Potential impacts of global climate change on hydrology and ecology of ephemeral freshwater systems of the forests of the northeastern United States. *Climatic Change* 95:469–483.

394 Brooks, G. C. 2020. On the use of demographic models to inform amphibian conservation and  
395 management: A case study of the Reticulated Flatwoods Salamander. Virginia Tech,  
396 Blacksburg, Virginia.

397 Brooks, G.C., T. A. Gorman, Y. Jiao, and C.A. Haas. 2020. Reconciling larval and adult  
398 sampling methods to model growth across life-stages. PLoS ONE 15:e0237737.

399 Capps, K. A., K. A. Berven, and S. D. Tiegs. 2015. Modelling nutrient transport and  
400 transformation by pool-breeding amphibians in forested landscapes using a 21-year  
401 dataset. *Freshwater Biology* 60:500–511.

402 Carey, C., and M. A. Alexander. 2003. Climate change and amphibian declines: Is there a link?  
403 *Diversity and Distributions* 9:111–121.

404 Chandler, H. C., D. L. McLaughlin, T. A. Gorman, K. J. McGuire, J. B. Feaga, and C. A. Haas.  
405 2017. Drying rates of ephemeral wetlands: implications for breeding amphibians.  
406 *Wetlands* 37:545–557.

407 Chandler, H. C., A. L. Rypel, Y. Jiao, C. A. Haas, and T. A. Gorman. 2016. Hindcasting  
408 historical breeding conditions for an endangered salamander in ephemeral wetlands of the  
409 southeastern USA: implications for climate change. PLoS ONE 11:e0150169.

410 Cianciolo, T. R., J. S. Diamond, D. L. McLaughlin, R. A. Slesak, A. W. D'Amato, and B. J.  
411 Palik. 2021. Hydrologic variability in black ash wetlands: Implications for vulnerability  
412 to emerald ash borer. *Hydrological Processes* 35:e14014.

413 Cohen, M.J., I. F. Creed, L. Alexander, N. B. Basu, A. J. K. Calhoun, C. Craft, E. D'Amico, E.  
414 DeKeyser, L. Fowler, H. E. Golden, J. W. Jawitz, P. Kalla, L. K. Kirkman, C. R. Lane,  
415 M. Lang, S. G. Leibowitz, D. B. Lewis, J. Marton, D. L. McLaughlin, D. M. Mushet, H.

416 Raanan-Kiperwas, M. C. Rains, L. Smith, and S. C. Walls. 2016. Do geographically  
417 isolated wetlands influence landscape functions? *PNAS* 113:1978–1986.

418 Dahl, T. E. 1990. Wetland losses in the United States: 1780s to 1980s. US Fish and Wildlife  
419 Service, Washington, D.C.

420 Dahl, T. E. 2011. Status and trends of wetlands and deepwater habitats in the conterminous  
421 United States 2004 to 2009. US Fish and Wildlife Service, Washington, D.C.

422 Davidson, N. C. 2014. How much wetland has the world lost? Long-term and recent trends in  
423 global wetland area. *Marine and Freshwater Research* 65:934–941.

424 Davis, C. L., D. A. W. Miller, E. H. C. Grant, B. J. Halstead, P. M. Kleeman, S. C. Walls, and  
425 W. J. Barichivich. 2019. Linking variability in climate to wetland habitat suitability: Is it  
426 possible to forecast regional responses from simple climate measures? *Wetlands Ecology*  
427 *and Management* 27:39–53.

428 de Szalay, F. A., and V. H. Resh. 1997. Responses of wetland invertebrates and plants important  
429 in waterfowl diets to burning and mowing of emergent vegetation. *Wetlands* 17:149–156.

430 Erwin, K. L. 2009. Wetlands and global climate change: The role of wetland restoration in a  
431 changing world. *Wetlands Ecology and Management* 17:71–84.

432 Frayer, W. E., T. J. Monahan, D. C. Bowden, and F. A. Graybill. 1983. Status and trends of  
433 wetlands and deepwater habitats in the conterminous United States, 1950's to 1970's. US  
434 Fish and Wildlife Service, Washington, D.C.

435 Galatowitsch, S. M., and A. G. van der Valk. 1996. Characteristics of recently restored wetlands  
436 in the prairie pothole region. *Wetlands* 16:75–83.

437 Gelman, A. 2004. Parameterization and Bayesian modeling. *Journal of the American Statistical*  
438 *Association* 99:537–545.

Gibbons, J. W., C. T. Winne, D.E. Scott, J. D. Wilson, X. Glaudas, K. M. Andrews, B. D. Todd,  
L. A. Fedewa, L. Wilkinson, R. N. Tsaliagos, S. J. Harper, J. L. Greene, T. D. Tuberville,  
B. S. Metts, M. E. Dorcas, J. P. Nestor, C. A., Young, T. Akre, R. N. Reed, K. A.  
Buhlmann, J. Norman, D. A. Croshaw, C. Hagen, and B. B. Rothermel. 2006.  
Remarkable amphibian biomass and abundance in an isolated wetland: Implications for  
wetland conservation. *Conservation Biology* 20:1457–1465.

Gorman, T. A., C. A. Haas, and J. G. Himes. 2013. Evaluating methods to restore amphibian  
habitat in fire-suppressed pine flatwoods wetlands. *Fire Ecology* 9:96–109.

Gorman, T. A., D. C. Bishop, and C. A. Haas. 2009. Spatial interactions between two species of  
frogs: *Rana okaloosae* and *R. clamitans clamitans*. *Copeia* 2009: 138–141.

Greenberg, C. H., S. Goodrick, J. D. Austin, and B. R. Parresol. 2015. Hydroregime prediction  
models for ephemeral groundwater-driven sinkhole wetlands: A planning tool for climate  
change and amphibian conservation. *Wetlands* 35:899–911

Guo, D., S. Westra, and T. Peterson. 2020. Evapotranspiration: Modelling actual, potential and  
reference crop evapotranspiration. R package version 1.15.

Haag, K. H., T. M. Lee, and D. C. Herndon. 2005. Bathymetry and vegetation in isolated marsh  
and cypress wetlands in the northern Tampa Bay Area, 2000–2004. U.S. Geological  
Survey Scientific Investigations Report 2005–5109, U.S. Geological Survey, Reston, VA.

Hargreaves, G. H. 1994. Defining and using reference evapotranspiration. *Journal of Irrigation  
and Drainage Engineering* 120:1132–1139.

Hargreaves, G. H., and Z. A Samani. 1985. Reference crop evapotranspiration from temperature.  
*Applied Engineering in Agriculture*. 1:96–99.

461 Hayashi, M., and D. O. Rosenberry. 2002. Effects of ground water exchange on the hydrology  
462 and ecology of surface water. *Ground Water* 40:309–316.

463 Hunter, J. T., and A. M. Lechner. 2017. A multiscale, hierarchical, ecoregional and floristic  
464 classification of arid and semi-arid ephemeral wetlands in New South Wales, Australia.  
465 *Marine and Freshwater Research* 69:418–431.

466 Huxman, T. E., B. P. Wilcox, D. D. Breshears, R. L. Scott, K. A. Snyder, E. E. Small, K.  
467 Hultine, W. T. Pockman, and R. B. Jackson. 2005. Ecohydrological implications of  
468 woody plant encroachment. *Ecology* 86:308–319.

469 Ingram, K. T., K. Dow, L. Carter, and J. Anderson. 2013. Climate of the southeast United States:  
470 Variability, change, impacts, and vulnerability. Inland Press, Washington, D. C.

471 Jenkins, D.G., S. Grissom, and K. Miller. 2001. Consequences of prairie wetland drainage for  
472 crustacean biodiversity and metapopulations. *Conservation Biology* 17:158–167.

473 Jones, C. N., D. L. McLaughlin, K. Henson, C. A. Haas, and D. A. Kaplan. 2018. From  
474 salamanders to greenhouse gases: Does upland management affect wetland functions?  
475 *Frontiers in Ecology and the Environment* 16:14–19.

476 Karl, T. R., J. M. Melillo, and T. C. Peterson. 2009. Global climate change impacts in the United  
477 States. Cambridge University Press, Cambridge, UK.

478 Kellner, K. 2021. jagsUI: A wrapper around ‘rjags’ to streamline ‘JAGS’ analyses. R package  
479 version 1.5.2.

480 Kéry, M. 2010. Introduction to WinBUGS for Ecologists. Academic Press, Bulington, MA.

481 Kéry, M., and J. A. Royle. 2016. Applied Hierarchical modeling in ecology: Analysis of  
482 distribution, abundance and species richness in R and BUGS. Academic Press, London,  
483 UK.

484 Kirkman, L. K., S. W. Golladay, L. Laclaire, and R. D. Sutter. 1999. Biodiversity in  
485 southeastern, seasonally ponded, isolated wetlands: Management and policy perspectives  
486 for research and conservation. *Journal of the North American Benthological Society*  
487 18:553–562.

488 Lee, S.-Y., M. E. Ryan, A. F. Hamlet, W. J. Palen, J. J. Lawler, and M. Halabisky. 2015.  
489 Projecting the hydrologic impacts of climate change on montane wetlands. *PLoS ONE*  
490 10:e0142960.

491 Li, Y., J. M. Cohen, and J. R. Rohr. 2013. Review and synthesis of the effects of climate change  
492 on amphibians. *Integrative Zoology* 8:145–161.

493 Link, W. A., and R. J. Barker. 2010. Bayesian inference with ecological applications. Academic  
494 Press, London, UK.

495 Liu, C., N. Cui, D. Gong, X. Hu, and Y. Feng. 2020. Evaluation of seasonal evapotranspiration  
496 of winter wheat in humid region of East China using large-weighted lysimeter and three  
497 models. *Journal of Hydrology* 590:125388.

498 Lu, J., G. Sun, S. G. McNulty, and N. B. Comerford. 2009. Sensitivity of pine flatwoods  
499 hydrology to climate change and forest management in Florida, USA. *Wetlands* 29:826–  
500 836.

501 McLaughlin, D. L., and M. J. Cohen. 2011. Thermal artifacts in measurements of fine-scale  
502 water level variation. *Water Resources Research* 47:W09601.

503 McLaughlin, D.L., D. A. Kaplan, and M. J. Cohen. 2014. A significant nexus: Geographically  
504 isolated wetlands influence landscape hydrology. *Water Resources Research* 50:7153–  
505 7166.



- Moser, S. C. 1996. A partial instructional module on global and regional land use/cover change: Assessing the data and searching for general relationships. *GeoJournal* 39:241–283.
- Mulholland, P. J., G. R. Best, C. C. Coutant, G. M. Hornberger, J. L. Meyer, P. J. Robinson, J. R. Stenberg, R. E. Turner, F. Vera-Herrera, and R. G. Wetzel. 1997. Effects of climate change on freshwater ecosystems of the south-eastern United States and the gulf coast of Mexico. *Hydrological Processes* 11:949–970.
- O'Donnell, K. M., A. F. Messerman, W. J. Barichivich, R. D. Semlitsch, T. A. Gorman, H. G. Mitchell, N. Allan, D. Fenolio, A. Green, F. A. Johnson, A. Keever, M. Mandica, J. Martin, J. Mott, T. Peacock, J. Reinman, S. S. Romanach, G. Titus, C. P. McGowan, and S. C. Walls. 2017. Structured decision making as a conservation tool for recovery planning of two endangered salamanders. *Journal of Nature Conservation* 37:66–72.
- Organization for Economic Cooperation and Development (OECD). 1996. Guidelines for aid agencies for improved conservation and sustainable use of tropical and sub-tropical wetlands. OECD, Paris.
- Palis, J. G. 1995. Larval growth, development, and metamorphosis of *Ambystoma cingulatum* on the Gulf Coastal Plain of Florida. *The Florida Scientist* 58:352–358.
- Palis, J. G. 1996. Flatwoods salamander (*Ambystoma cingulatum* Cope). *Natural Areas Journal* 16:49–54.
- Palis, J., and G. Hammerson. 2008. *Ambystoma bishopi*. The IUCN Red List of Threatened Species: <https://www.iucnredlist.org/species/136128/4245386>. Downloaded on 27 October 2021.

527 Palis, J. G., M. J. Aresco, and S. Kilpatrick. 2006. Breeding biology of a Florida population of  
528 *Ambystoma cingulatum* (Flatwoods salamander) during a drought. Southeastern  
529 Naturalist 5:1–8.

530 Parmesan, C. 2006. Ecological and evolutionary responses to recent climate change. Annual  
531 Review of Ecology, Evolution, and Systematics 37:637–669.

532 Pierce, D. W., D. R. Cayan, and B. L. Thrasher. 2014. Statistical downscaling using localized  
533 constructed analogs (LOCA). Journal of Hydrometeorology 15:2558–2585.

534 Plummer, M. 2003. JAGS: A program for analyses of Bayesian graphical models using Gibbs  
535 sampling. Proceedings of the 3rd International Workshop on Distributed Statistical  
536 Computing.

537 Pounds, J. A. 2001. Climate and amphibian declines. Nature 410:639–640.

538 R Core Team. 2021. R: A language and environment for statistical computing. R Foundation for  
539 Statistical Computing, Vienna, Austria.

540 Riahi, K., S. Rao, V. Krey, C. Cho, V. Chirkov, G. Fischer, G. Kindermann, N. Nakicenovic, and  
541 P. Rafaj. 2011. RCP 8.5—A scenario of comparatively high greenhouse gas emissions.  
542 Climatic Change 109:33–57.

543 Sacerdote, A. B., and R. B. King. 2009. Dissolved oxygen requirements for hatching success of  
544 two ambystomatid salamanders in restored ephemeral ponds. Wetlands 29:1202–1213.

545 Seigel, R. A., A. Dinsmore, and S. C. Richter. 2006. Using well water to increase hydroperiod as  
546 a management option for pond-breeding amphibians. Wildlife Society Bulletin 34:1022–  
547 1027.

548 Shulze, C. D., R. D. Semlitsch, K. M. Trauth, and J. E. Gardner. 2012. Testing wetland features  
549 to increase amphibian reproductive success and species richness for mitigation and  
550 restoration. *Ecological Applications* 22:1675–1688.

551 Skelly, D. K. 1997. Tadpole communities: Pond permanence and predation are powerful forces  
552 shaping the structure of tadpole communities. *American Scientist* 85:36–45.

553 Skelly, D. K., L. K. Freidenburg, and J. M. Kiesecker. 2002. Forest canopy and the performance  
554 of larval amphibians. *Ecology* 83:983–992.

555 Stagge, J. H., L. M. Tallaksen, L. C.-Y. Xu, and H. A. J. Van Lanen. 2014. Standardized  
556 Precipitation-Evapotranspiration Index (SPEI): Sensitivity to potential evapotranspiration  
557 model and parameters. *Hydrology in a Changing World: Environmental and Human*  
558 *Dimensions*. In *Proceedings of FRIEND-Water 2014*, Daniell, T. M. (ed). Montpellier,  
559 France, 7–10 October 2014. IAHS Publ. No. 363, IAHS Press, Centre for Ecology and  
560 Hydrology: Wallingford, UK, 367–373.

561 Sun, G., S. G. McNulty, D. M. Amatya, R. W. Skaggs, L. W. Swift Jr., J. P. Shepard, and H.  
562 Riekerk. 2002. A comparison of the watershed hydrology of coastal forested wetlands  
563 and the mountainous uplands in the Southern US. *Journal of Hydrology* 263:92–104.

564 Thomson, A. M., K. V. Calvin, S. J. Smith, G. P. Kyle, A. Volke, P. Patel, S. Delgado-Arias, B.  
565 Bond-Lamberty, M. A. Wise, L. E. Clarke, J. A. Edmonds. 2011. RCP 4.5: A pathway  
566 for stabilization of radiative forcing by 2100. *Climatic Change* 109:77–94.

567 Tiner, R. W. 2003. Geographically isolated wetlands of the United States. *Wetlands* 23:494–516.

568 Todd, B. D., D. E. Scott, J. H. K. Pechmann, and J. W. Gibbons. 2011. Climate change correlates  
569 with rapid delays and advancements in reproductive timing in an amphibian community.  
570 *Proceedings of the Royal Society B* 278:2191–2197.

571 U.S. Fish and Wildlife Service. 2009. Endangered and threatened wildlife and plants;  
572 determination of endangered status for reticulated flatwoods salamander; designation of  
573 critical habitat for frosted flatwoods salamander and reticulated flatwoods salamander.  
574 Federal Register. 74 FR 6700.

575 U.S. Fish and Wildlife Service. 2020. Species status assessment for the Reticulated Flatwoods  
576 Salamander (*Ambystoma bishopi*), Version 1.0. Panama City Field Office Panama City,  
577 Florida.

578 Vanschoenwinkel, B., A. Hulsmans, E. de Roeck, C. de Vries, M. Seaman, and L. Brendonck.  
579 2009. Community structure in temporary freshwater pools: Disentangling the effects of  
580 habitat size and hydroregime. *Freshwater Biology* 54:1487–1500.

581 Walls, S. C., W. J. Barichivich, M. E. Brown, D. E. Scott, and B. R. Hossack. 2013. Influence of  
582 drought on salamander occupancy of isolated wetlands on the southeastern coastal plain  
583 of the United States. *Wetlands* 33:345–354.

584 Walls, S. C., W. J. Barichivich, M. Milinichik, K. M. O'Donnell, M. E. Owens, T. Peacock, J.  
585 Reinman, R. C. Watling, and O. E. Wetsch. 2019. Seeking shelter from the storm:  
586 Conservation and management of imperiled species in a changing climate. *Ecology and*  
587 *Evolution* 9:7122–7133.

588 Winter, T. C., and J. W. LaBaugh. 2003. Hydrologic considerations in defining isolated  
589 wetlands. *Wetlands* 23:532–540.

590 WMO. 2006. Drought monitoring and early warning: Concepts, progress, and future challenges.  
591 WMO-No. 1006, World Meteorological Organization, Geneva, Switzerland.

592 Yang, L. H., and V. H. W. Rudolf. 2010. Phenology, ontogeny, and the effects of climate change  
593 on the timing of species interactions. *Ecology Letters* 13:1–10.

594 Zedler, J. B., and S. Kercher. 2005. Wetland resources: Status, trends, ecosystem services, and  
 595 restorability. *Annual Review of Environment and Resources* 30:39–74.

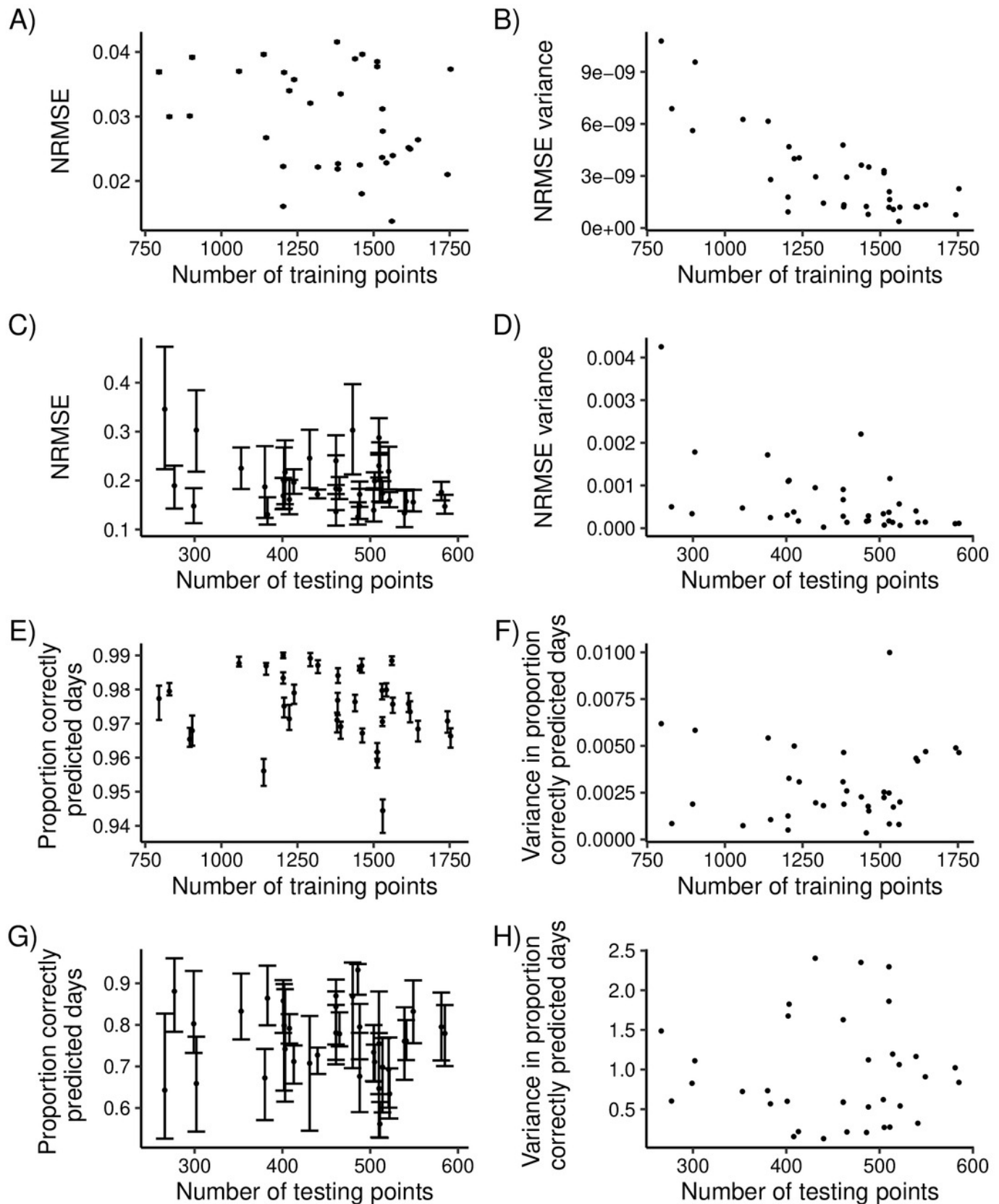
596 Zhu, J., G. Sun, W. Li, Y. Zhang, G. Miao, A. Noormets, S. G. McNulty, J. S. King, M. Kumar,  
 597 and X. Wang. 2017. Modeling the potential impacts of climate change on the water table  
 598 level of selected forested wetlands in the southeastern United States. *Hydrology and*  
 599 *Earth System Sciences* 21:6289–6305.

600

# Figure 1

Plots describing performance of models predicting water levels in ephemeral wetlands on Eglin Air Force Base, Florida

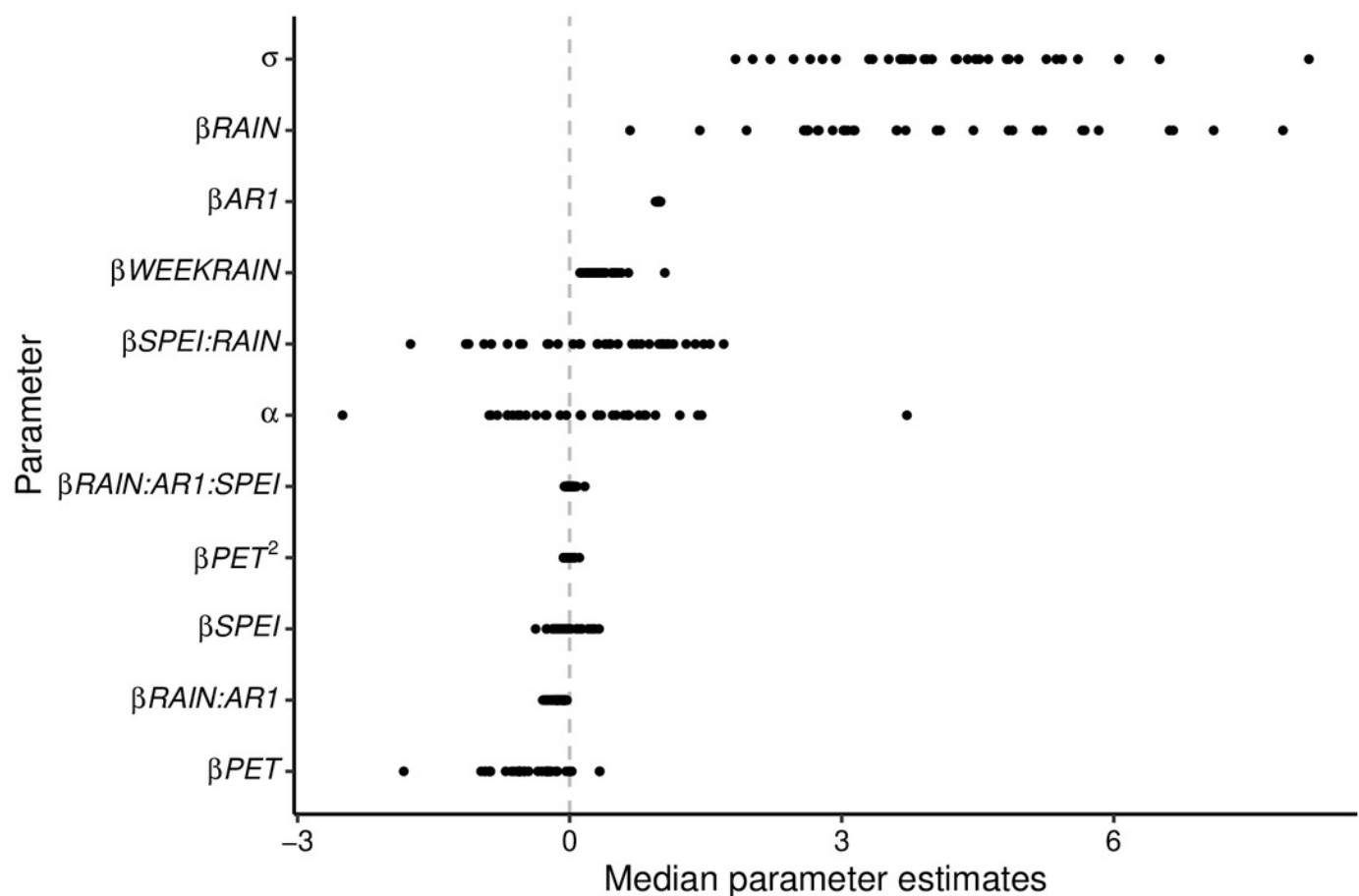
(A, B, C, D) Model accuracy for predicted water levels was estimated by normalized root mean squared error (NRMSE) and (E, F, G, H) for the prediction of length of presence of surface water by the proportion of days that correctly predicted to either have or not have water. Results are divided by the number of training (A, B, E, F) or testing (C, D, G, H) data points (daily observations) used in the model. Error bars in A, C, E, and G represent the upper and lower 95% highest density posterior and points represent median values, while points in B, D, F, and H are variance estimates. Note that the y-axis scale is different for all eight panels.



# Figure 2

Median parameter estimates for Bayesian first order autoregressive fixed effect models that predicted daily water levels for wetlands on Eglin Air Force Base, Florida.

Model parameters included an error term ( $\sigma$ ), precipitation ( $\beta_{RAIN}$ ), an autoregressive term ( $\beta_{AR1}$ ), an intercept ( $\alpha$ ), total precipitation over the previous seven days ( $\beta_{WEEKRAIN}$ ), the 12-month standardized precipitation-evapotranspiration index ( $\beta_{SPEI}$ ), potential evapotranspiration ( $\beta_{PET}$ ), and associated interactions and quadratic effects. Points represent the median value for each of the 35 monitoring wells included in this study.

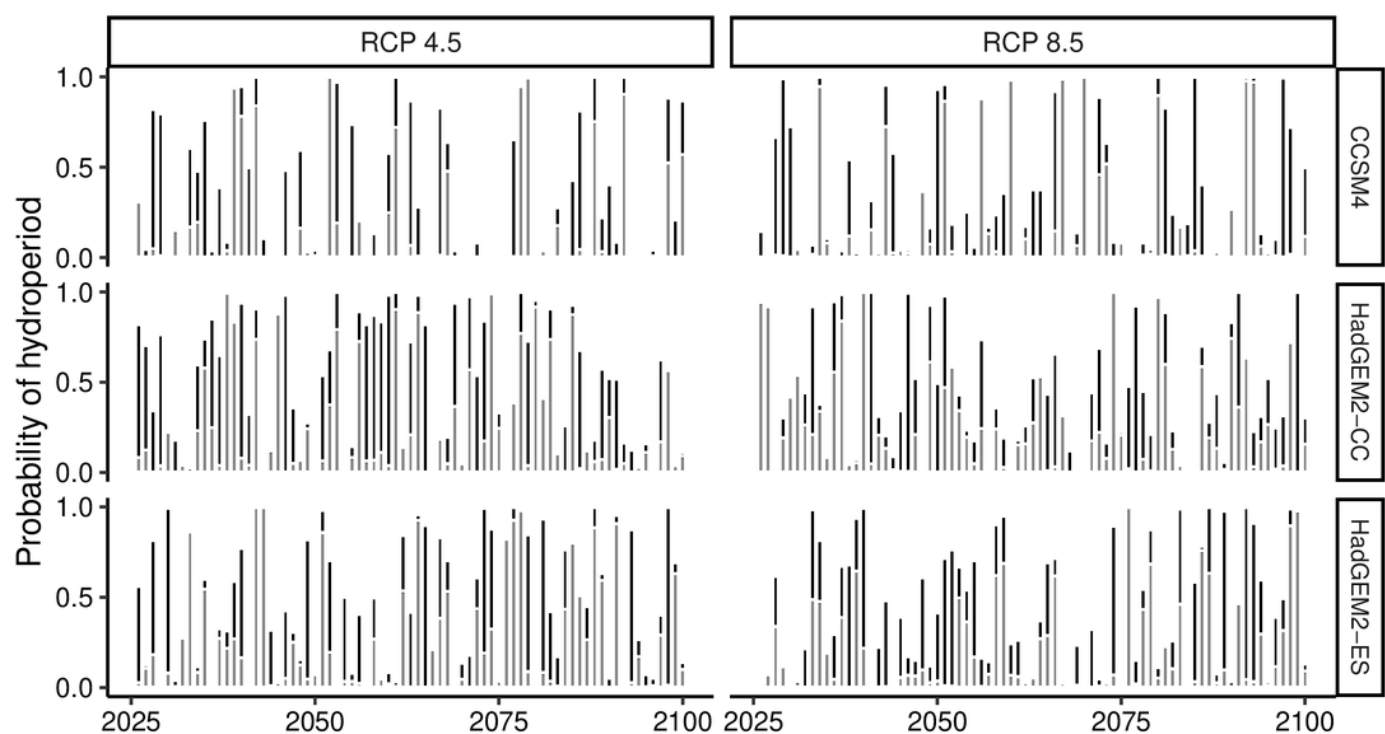




# Figure 3

Predicted hydroperiods for a single ephemeral wetland on Eglin Air Force Base, Florida.

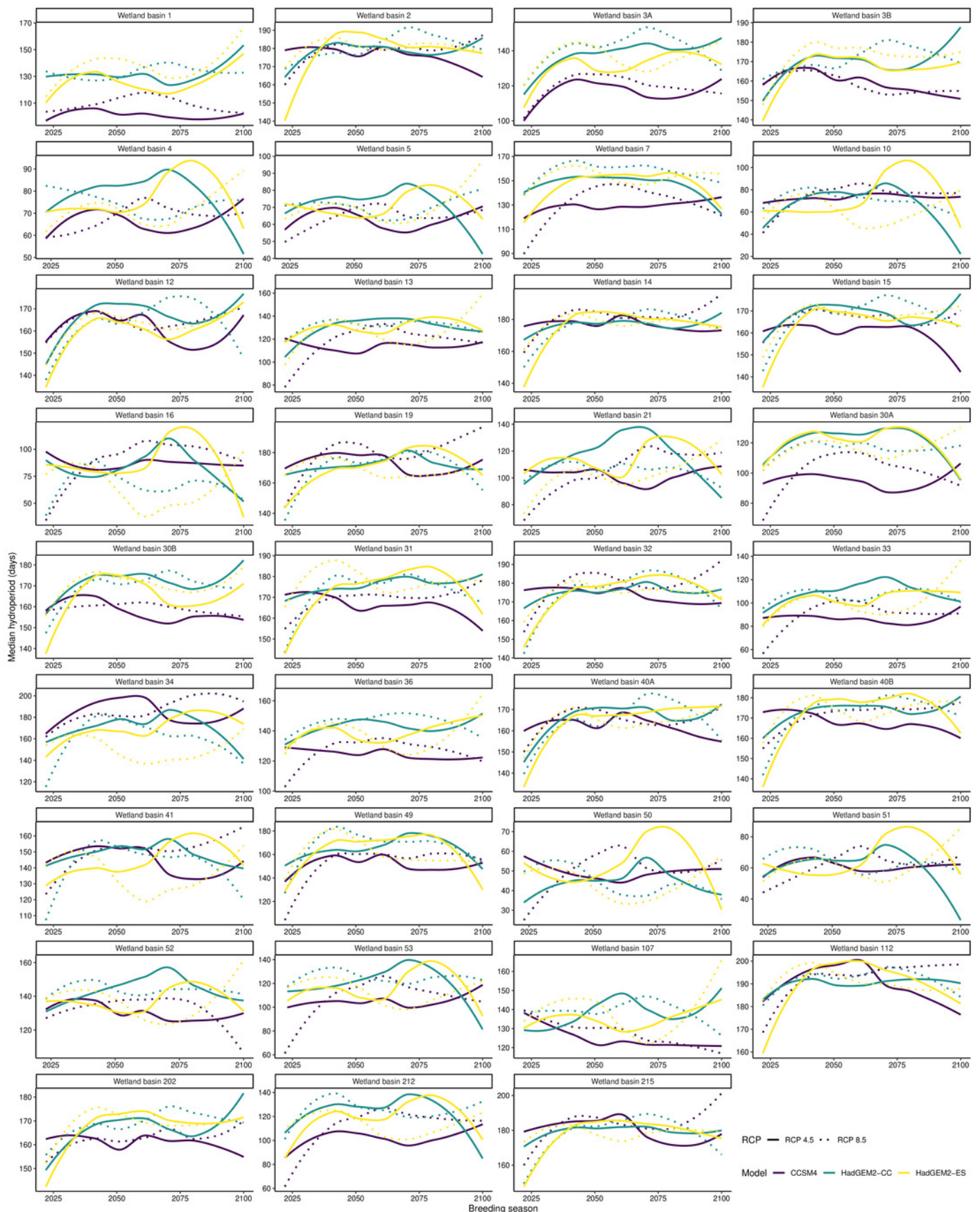
Probability of an 11-week (black) or 15-week (gray) hydroperiod in a single Reticulated Flatwoods Salamander (*Ambystoma bishopi*) breeding wetland. There have been reports of successful metamorphosis of flatwoods salamanders after an 11-week hydroperiod, but 15-weeks is a more conservative estimate of suitable hydroperiod. Predictions were made across three global climate models and two emission scenarios (RCPs). Hydroperiods were calculated from 1 November to 31 May.



# Figure 4

Median predicted hydroperiod from 2025 to 2100 for 35 wetlands on Eglin Air Force Base, Florida.

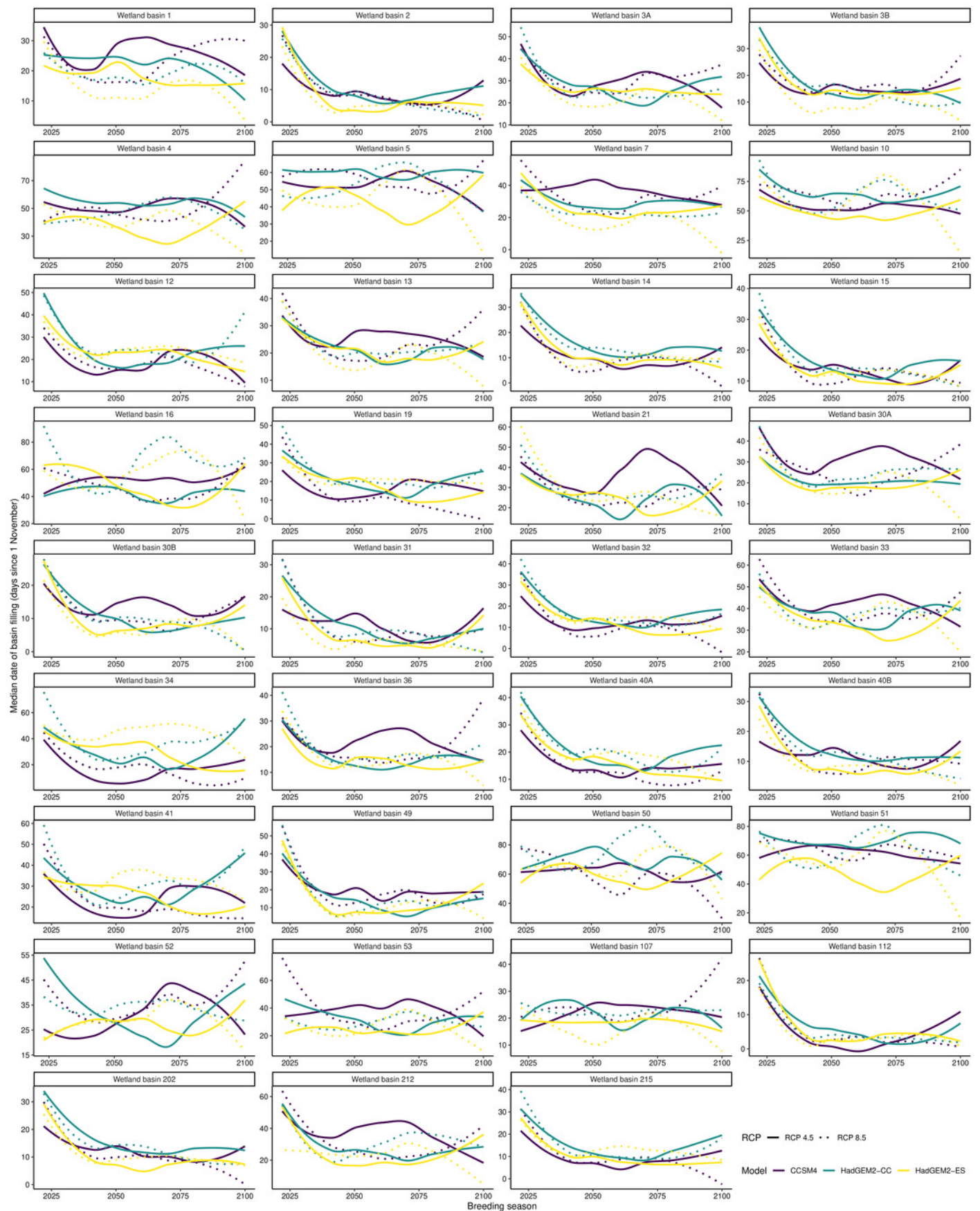
Lines represent smoothed (using the *loess* function in R) predictions for three global climate model and two emission scenario (RCPs) combinations (six total scenarios). Hydroperiods were calculated across the Reticulated Flatwoods Salamander (*Ambystoma bishopi*) breeding season (1 November to 31 May).



# Figure 5

Median predicted date of basin filling from 2025 to 2100 for 35 wetlands on Eglin Air Force Base, Florida.

Lines represent smoothed (using the *loess* function in R) predictions for three global climate model and two emission scenario (RCPs) combinations (six total scenarios). Fill dates were calculated relative to the Reticulated Flatwoods Salamander (*Ambystoma bishopi*) breeding season (1 November to 31 May).



# Figure 6

Relationship between wetland hydroperiod and probability of consecutive short-hydroperiod years for wetlands on Eglin Air Force Base, Florida.

Relationship between the median number of Reticulated Flatwoods Salamander (*Ambystoma bishopi*) breeding seasons (1 November to 31 May) with an at least 15-week hydroperiod from 2025–2100 versus the proportion of iterations for which a given wetland had at least five consecutive years without a hydroperiod of 15 weeks. The vertical (0.75) and horizontal (15) dotted lines represent cutoffs indicating a high probability (i.e., above 75% of proportions or less than 20% of breeding seasons) of extirpation of the salamander population. Predictions were made across three global climate models and two emission scenarios (RCPs).

