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RESEARCH ARTICLE

Automatic variable selection in ecological niche modeling: A case study using Cassin's Sparrow (*Peucaea cassinii*)

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Abstract

MERRA/Max provides a feature selection approach to dimensionality reduction that enables direct use of global climate model outputs in ecological niche modeling. The system accomplishes this reduction through a Monte Carlo optimization in which many independent Max-Ent runs, operating on a species occurrence file and a small set of randomly selected variables in a large collection of variables, converge on an estimate of the top contributing predictors in the larger collection. These top predictors can be viewed as potential candidates in the variable selection step of the ecological niche modeling process. MERRA/Max's Monte Carlo algorithm operates on files stored in the underlying filesystem, making it scalable to large data sets. Its software components can run as parallel processes in a high-performance cloud computing environment to yield near real-time performance. In tests using Cassin's Sparrow (Peucaea cassinii) as the target species, MERRA/Max selected a set of predictors from Worldclim's Bioclim collection of 19 environmental variables that have been shown to be important determinants of the species' bioclimatic niche. It also selected biologically and ecologically plausible predictors from a more diverse set of 86 environmental variables derived from NASA's Modern-Era Retrospective Analysis for Research and Applications Version 2 (MERRA-2) reanalysis, an output product of the Goddard Earth Observing System Version 5 (GEOS-5) modeling system. We believe these results point to a technological approach that could expand the use global climate model outputs in ecological niche modeling, foster exploratory experimentation with otherwise difficult-to-use climate data sets, streamline the modeling process, and, eventually, enable automated bioclimatic modeling as a practical, readily accessible, low-cost, commercial cloud service.

Introduction

Ecological niche modeling (ENM) consists of a set of techniques and tools that use species occurrence records and environmental data to predict the relative suitability of habitats [1]. It is used across a wide range of disciplines, including fields as diverse as biogeography and phylogeny [2], conservation biology and epidemiology [3, 4], invasion biology [5], and

archaeology [6]. In recent years, ecological niche models have become particularly important in understanding the influence of climate change on the geographic distribution of species [7]. This, in turn, has led to greater use of global climate model (GCM) outputs as environmental predictors [8]. GCMs combine observations from an array of satellite, airborne, and *in-situ* sensors to create global representations of the climate system, including historical simulations and future projections for hundreds of climate variables [9]. The largest and most sophisticated of these, however, produce complex, petabyte-scale data sets, which complicates variable selection and limits their direct use in ecological modeling [10–12].

Part of the problem lies in the fact that most ENM software tools require predictors and observations to be memory-resident in order for the programs to work [13, 14]. This results in run-times and space requirements that have linear or higher-order scaling properties with respect to the size of a model's inputs. This generally poses few difficulties. But when the number of predictors becomes large, compute times can become impractically long, models can become overly complex, and efforts to understand any particular variable's contribution to model formation, either as an aspect of model analysis or as a way of selecting subsets of variables for further model refinement, can become challenging [10, 15–19]. An effective way of dealing with large, externally-stored environmental data sets that preserves the advantages of conventional tools while overcoming this limitation would benefit the ENM community.

In previous work, we demonstrated the potential of a MaxEnt-based Monte Carlo method that addresses this issue by screening large data collections for viable predictors [14]. Based on a machine learning approach to maximum entropy modeling, MaxEnt is one of the most popular software packages in use today by the ENM community [20–22]. Among its many advantages, MaxEnt ranks the contribution of predictor variables in the formation of its models. Our Monte Carlo method exploits this feature in an ensemble strategy whereby many independent MaxEnt runs, each drawing on a small, random subset of variables stored in the filesystem, converge on a global estimate of the top contributing subset of variables in the larger collection. These top-contributing predictors can then be studied in more detailed ways, augmented with other variables, and further refined prior to final model construction. We believe a screening step, such as this, could help the ENM process, particularly when working with large, multidimensional data sets where selection through ecological reasoning or other means is not apparent.

In our earlier, proof-of-concept work, we implemented the Monte Carlo selection algorithm as a single-threaded program running on a MacBook Pro laptop computer [14]. In the current study, we have implemented a parallel version of the Monte Carlo method in a highperformance cloud computing environment. Our goal this time has been to characterize the run-time performance and scaling properties of a parallel implementation of the Monte Carlo algorithm and demonstrate its variable selection behavior with two example use cases. We call the prototype system MERRA/Max to reflect its reliance on MaxEnt and our interest in using the technology to screen for bioclimatic predictors in NASA's Modern-Era Retrospective Analysis for Research and Applications Version 2 (MERRA-2) dataset, what we view as an underutilized and potentially important GCM resource for the ecological modeling community [23, 24].

A second goal for this paper is to open a discussion about the potential merits of this technology and lay the groundwork for experiments to evaluate its scientific value more fully. Reanalyses, such as MERRA-2, simulate hundreds of low-level physical drivers of the Earth system at extraordinarily fine temporal scale, and they do so over the entire four-decade span of the satellite era. A technology like MERRA/Max makes this remarkable resource practically available to the ENM community. In this paper, we begin to make the case for that. Some of the most important conservation questions scientists hope to answer are hobbled by current predictor selection methods. Rare species, for example, are generally represented by sparce occurrence records. Effective ENM, in these cases, requires a small number of high-quality predictors to avoid overfitting. We show how MERRA/Max can help with that. As large climate data sets continue to grow in size, they become less accessible to the science community and less usable in today's suite of machine learning and statistical analysis tools. MERRA/Max shows how parallelizable, external-memory algorithms can address that problem. And, while the topic of variable selection in ENM is represented by a vast literature, most approaches in use today are difficult or impossible to automate, do not scale well to large data sets, or provide limited insight into the underlying biology or ecology of the organisms being studied. In the pages that follow, we show that a technology like MERRA/Max can potentially help overcome these limitations.

This project builds on a twenty-year history of technology research and development at NASA focusing on applications of high-performance computing to ecological modeling [25–29] and the big data challenges of Earth science [30-40]. It complements this body of work by looking at ways that machine learning and high-performance cloud computing can extend existing capabilities and open new opportunities for research. In a fully realized, operational implementation of the technologies described here, we see MERRA/Max as one element of a bioclimatic modeling service enabled by a suite of high-performance data subsetting and data analytic tools of the sort becoming increasingly available to the research community through commercial cloud services [41–46].

Materials and methods

System architecture and implementation

We implemented MERRA/Max in a 100-core testbed within the NASA Center for Climate Simulation's (NCCS's) Advanced Data Analytics Platform (ADAPT). ADAPT is a managed virtual machine (VM) environment most closely resembling a platform-as-a-service (PaaS) cloud [47]. It features over 300 physical hypervisors that host one or more VMs, each having access to multiple shared, centralized data repositories. The hypervisor hardware consists of 2.2 GHz 24-core Intel Xeon Broadwell E5-2650 v4 processors with 256 GB of memory. The MERRA/Max testbed consists of a dedicated set of ten 10-core Debian Linux 9 Stretch VMs. We used shell scripts, R Version 4.0.1 [48], ENMeval Version 0.3.1 [49, 50], and MaxEnt Version 3.4.1 [51] to develop MERRA/Max's software components, which collectively realize the Monte Carlo algorithm through the interactions shown in Fig 1.

Conceptually, MERRA/Max sits atop a collection of variables stored in the underlying filesystem; when provided a species occurrence file, the system screens the collection to find the most important predictors for the input provided (Fig 1, Steps 1–5). The Monte Carlo screening process is initiated by the *mcensemble.sh* script, which launches an *mcsprint.sh* script on each of the 10 MERRA/Max VMs (Fig 1, Step 2). The *mcsprint.sh* script, in turn, creates parallel sprint runs by launching an R run-time environment and an *MCSprint.R* program on each of the 10 VM's 10 processor cores (Fig 1, Steps 2.1). The *MCSprint.R* programs perform repeated MaxEnt runs on random pairs of variables read from the shared filesystem until a desired level of sampling is achieved (Fig 1, Steps 3.1). *MCSprint.R* maintains a tally table that tracks the number of times each variable is used along with its accumulating permutation importance, then writes the table to a shared directory (Fig 1, Steps 4.1). We used the operating system's *MCSprint.R* process identifiers to create unique *pid.tt* file names for the output tally tables. When all the sprints have completed their work, *MCSelect.R* concatenates the *pid.tt* files into a global tally table (Step 4), computes the average permutation importance for each variable, then sorts them to reveal the top contributing variables identified by the ensemble's runs





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(Fig_1, Step 5). While MERRA/Max relies on MaxEnt to perform selection, it is important to note that the resulting set of selected variables can be used in any ENM application or species distribution modeling approach.

Run-time performance and scaling properties

MERRA/Max's run-time performance and scaling properties are affected by several factors. The total amount of time needed for MERRA/Max to complete a screening run (T) is primarily determined by the number of variables in the collection being screened (N), the average number of random samples taken of each variable in the collection (S), the number of random variables used in each independent MaxEnt sampling run (V), and the number of processor cores available in the compute environment (C). To understand the interplay of these factors, we gathered timing metrics on a series of ensembles with varying values of N, S, and C.

In a first ensemble with N = 2 and C = 10, we used 10 *parallel* sprints (one sprint per core) in which each sprint performed five *sequential* two-variable sampling runs to achieve an average sample size for each collection variable of S = 50. We then completed the S = 50 series with ensembles in which N and C were proportionally increased to N = 18 and C = 90. This process was repeated using 10- and 15-run sprints respectively to create a timing series for S = 100 and S = 150. These measurements allowed us to quantify MERRA/Max's run-time performance, estimated optimal performance, and its scaling properties within the constraints of a 100-core testbed.

We did not quantify non-algorithmic influences on run time, such as predictor resolution, occurrence file size, competing system processes, filesystem performance, processor failures, process failures (abends), or MaxEnt parameter settings, as these tend to be intrinsic properties of the science question being studied, the compute environment, or the MaxEnt software itself. These non-algorithmic factors either have an idiosyncratic impact on overall run time that is constant for any particular application of MERRA/Max, or they are beyond user control.

bio01	Annual mean temperature
bio02	Mean diurnal range (mean of monthly (max temp—min temp))
bio03	Isothermality (bio2/bio7) (×100)
bio04	Temperature seasonality (standard deviation ×100)
bio05	Maximum temperature of warmest month
bio06	Minimum temperature of coldest month
bio07	Temperature annual range (bio5-bio6)
bio08	Mean temperature of wettest quarter
bio09	Mean temperature of driest quarter
bio10	Mean temperature of warmest quarter
bio11	Mean temperature of coldest quarter
bio12	Annual precipitation
bio13	Precipitation of wettest month
bio14	Precipitation of driest month
bio15	Precipitation seasonality (coefficient of variation)
bio16	Precipitation of wettest quarter
bio17	Precipitation of driest quarter
bio18	Precipitation of warmest quarter
bio19	Precipitation of coldest quarter

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For development testing, we used Cassin's Sparrow (*Peucaea cassinii* Woodhouse, 1852) as the target species [52] and Worldclim Version 2.1's 19 Bioclim variables at a resolution of 5.0 arc-minutes (Table 1) as environmental predictors [53, 54]. We obtained Cassin's Sparrow observational records for the year 2016 from the Global Biodiversity Information Facility (GBIF) [55]. Because of their secretive nature, Cassin's Sparrows are generally detected in the field by the presence of singing males that define and defend breeding territories that range in size from 0.6 to 12.9 acres [52, 56–58]. After removing replicates, we thinned the records to non-overlapping observations within a 16 km buffer around each point to avoid double counting the same individuals. This resulted in a total of 609 observations, which were used for testing throughout the study. The predictor layers were clipped to the coverage area of our observational data, reprojected, and formatted for use by MaxEnt using rgdal Version 1.5–18 [59] following the guidelines of Hijmans et al. [60].

We adopted a standard MERRA/Max screening configuration that we used as the default in all our timing trials and use cases. This included a MaxEnt feature class (FC) setting of LQHP (linear, quadratic, hinge, and product), a regularization multiplier (RM) setting of 1.0, 10 replicate cross-validation, and ten thousand background points from across the study area [14]. We used V = 2 random variables in all the independent MaxEnt sampling runs. Additional detail about MERRA/Max's default screening parameters and the rationale for their choice are provided in S1 Appendix.

Use case scenarios and selection behavior

To demonstrate MERRA/Max's selection behavior and show how the system might be used in actual practice, we developed two use case scenarios in which we modeled the bioclimatic niche of Cassin's Sparrow, a species known to be sensitive to many of the variables used in the study [52, 56, 58, 61]. Each use case involved three steps. The first was a *Variable Screening* step, in which MERRA/Max selected the top six contributing predictors from a collection of variables using an average sampling rate of S = 50. In previous work, we demonstrated that

sampling at this rate converges quickly on a stable set of top predictors [14]. Here, we confirmed this behavior by first performing three ensemble runs. We then used the averaged results from these three screenings to settle on the top contributors. This was followed by a *Predictor Refinement* step, where we used variance inflation factor (VIF) analysis to reduce collinearities in the selected predictors [62]. VIF shows the degree to which standard errors are inflated due to the levels of multicollinearities. Using ENMtools Version 1.4.4 [63], we first calculated Pearson correlation coefficient (r), coefficient of determination (r²), and VIF [1÷ (1– r²)] values for the selected predictors, then eliminated the least contributing variable in any pair of variables having r > 0.8, r² > 0.8, and VIF > 10.0 [62]. In a final *Model Calibration / Final Model Run* step, we used the ENMeval R package [49, 50] to identify optimal settings for the remaining, non-collinear predictors by performing a series of MaxEnt runs across all possible combinations of five feature classes (L, LQ, H, LQH, and LQHP) and regularization multiplier values ranging from 0.5 to 4.0 in increments of 0.5. The combination of settings resulting in the lowest value for Akaike's information criterion corrected for small sample size (AICc) [64] was taken to be an optimal tuning configuration.

We used the same 2016 Cassin's Sparrow occurrence data in each scenario that we used for development testing. However, the two use cases operated on different sets of environmental predictors. In the first, we again used WorldClim's 19 Bioclim variables. In the second use case, to gain experience with an even larger collection and demonstrate the system's application to a novel set of Intergovernmental Panel on Climate Change (IPCC)-class GCM outputs [10, 65], we used variables obtained directly from the Modern-Era Retrospective Analysis for Research and Applications Version 2 (MERRA-2) reanalysis. In contrast to Worldclim's Bioclim predictors, which are derived from 30-year averages of spatially interpolated weather station temperature and precipitation data [53, 54], the MERRA-2 reanalysis is produced by NASA's Goddard Earth Observing System Version 5 (GEOS-5) [23, 66, 67]. The system integrates observational data with numerical models to produce a global temporally and spatially consistent synthesis of over 600 climate-related variables. MERRA-2's spatial resolution is $1/2^\circ$ latitude $\times 5/8^\circ$ longitude (i.e., 55.5×69.4 km at the equator) $\times 72$ vertical levels extending through the stratosphere. Its temporal resolution is hourly and extends from 1979 to the present, nearly the entire satellite era. The complete MERRA-2 collection is about one petabyte in size.

For the current study, we created a test collection of 86 MERRA-2 variables of potential ENM interest. These were drawn from four MERRA-2 collections and included modeled, two-dimensional values for atmospheric attributes and heat, wind, radiation, and land surface attributes (Table 2). The test collection contains weekly and monthly maximum, minimum, and average values (or sums as appropriate) for each variable for the 40 years spanning 1980 to 2020. Importantly, the collection contains modeled values for the temperature and precipitation variables that form the basis for Bioclim's 19 predictors, which highlight climate conditions generally understood to relate to a species' physiology, *plus* an extended array of environmental attributes of potentially more direct biological significance, such as soil moisture and evaporation, wind direction and speed, and various solar radiation fluxes (Table 2) [53, 68-72]. For our use case, we used xarray [73] to create annual averages for the 86 variables for the year 2016, the year corresponding to the observation year of our Cassin's Sparrow occurrence data. We used modeled values at 850 hPa, where appropriate, to reflect surface conditions. The hPa (hectopascal) atmospheric pressure unit is an expression of altitude. Generally, 850 hPa lies immediately above the atmospheric boundary layer (about 1.5 km), where daily surface variations in temperature, humidity, wind speed, etc. have little if any effect on measured or modeled values [9]. These layers were then prepared for use with MaxEnt as described above.

To evaluate MERRA/Max's selection behavior, we created initial MaxEnt models using the top six predictors selected by the three screenings in the *Variable Screening* step. Then, using

	M2T1NXSLV	2D Atmospheric single-level diagnostics			
M01	PS	Time averaged surface pressure			
M02	U850	Eastward wind at 850 hPa			
M03	V850	Northward wind at 850 hPa			
M04	T850	Temperature at 850 hPa			
M05	Q850	Specific humidity at 850 hPa			
M06	H1000	Height at 1000 hPa			
M07	TS	Surface skin temperature			
M08	QV2M	2-meter specific humidity			
M09	QV10M	10-meter specific humidity			
M10	T2M	2-meter air temperature			
M11	T10M	10-meter air temperature			
M12	U2M	2-meter eastward wind			
M13	U10M	10-meter eastward wind			
M14	U50M	Eastward wind at 50 meters			
M15	V2M	2-meter northward wind			
M16	V10M	10-meter northward wind			
M17	V50M	Northward wind at 50 meters			
	M2T1NXFLX	2D Surface turbulent flux diagnostics			
M18	EFLUX	Latent heat flux (positive upward)			
M19	HFLUX	Sensible heat flux (positive upward)			
M20	TAUX	Eastward surface wind stress			
M21	TAUY	Northward surface wind stress			
M22	RHOA	Surface air density			
M23	TSH	Effective turbulence skin temperature			
M24	QSH	Effective turbulence skin humidity			
M25	PGENTOT	Total generation of precipitation			
M26	PREVTOT	Total re-evaporation of precipitation			
	M2T1NXRAD	2D Surface and TOA radiation fluxes			
M27	EMIS	Surface emissivity			
M28	ALBEDO	Surface albedo			
M29	LWGEM	Emitted longwave at the surface			
M30	LWGAB	Surface absorbed longwave			
M31	LWGABCLR	Surface absorbed longwave assuming clear sky			
M32	LWGABCLRCLN	Surface absorbed longwave assuming clear clean sky			
M33	LWGNT	Surface net downward longwave flux			
M34	LWGNTCLR	Surface net downward longwave flux assuming clear day			
M35	LWGNTCLRCLN	Surface net downward longwave flux assuming clear clean day			
M36	SWGDN	Surface incident shortwave flux			
M37	SWGDNCLR	Surface incident shortwave flux assuming clear sky			
M38	SWGNT	Surface net downward shortwave flux			
M39	SWGNTCLR	Surface net downward shortwave flux assuming clear sky			
M40	SWGNTCLN	Surface net downward shortwave flux assuming clean sky			
M41	SWGNTCLRCLN	Surface net downward shortwave flux assuming clear clean sky			
M42	TAUTOT	Optical thickness of all clouds			
M43	CLDTOT	Total cloud fraction			
	M2T1NXLND	2D Land surface diagnostics			
M44	GRN	Vegetation greenness fraction (LAI-weighted)			

Table 2. MERRA-2 variables.

(Continued)

M45	LAI	Leaf area index		
M46	GWETPROF	Total profile soil wetness		
M47	GWETROOT	Root zone soil wetness		
M48	GWETTOP	Top soil layer wetness		
M49	TSURF	Mean land surface temperature (incl. snow)		
M50	TPSNOW	Top snow layer temperature		
M51	TUNST	Surface temperature of unsaturated (but non-wilting) zone		
M52	TSA T	Surface temperature of saturated zone		
M53	TWLT	Surface temperature of wilting zone		
M54	SNODP	Snow depth		
M55	RUNOFF	Overland runoff		
M56	BASEFLOW	Baseflow		
M57	QINFIL	Soil water infiltration rate		
M58	FRUNST	Fractional unsaturated (but non-wilting) area		
M59	FRSAT	Fractional saturated area		
M60	FRSNO	Fractional snow-covered area		
M61	FRWLT	Fractional wilting area		
M62	PARDFLAND	Surface downward photosynthetically active radiation diffuse flux		
M63	PARDR LAND	Surface downward photosynthetically active radiation beam flux		
M64	SHLAND	Sensible heat flux from land		
M65	LHLAND	Latent heat flux from land		
M66	LWLAND	Net downward longwave flux over land		
M67	SWLAND	Net downward shortwave flux over land reservoirs		
M68	GHLAND	Downward heat flux into top soil layer		
M69	TWLAND	Total water stored in land reservoirs		
M70	TELAND	Energy stored in all land		
M71	WCHANGE	Total land water change per unit time		
M72	ECHANGE	Total land energy change per unit time		
M73	SPLAND	Spurious land energy source		
M74	SPWATR	Spurious land water source		
M75	SPSNOW	Spurious snow energy source		
M76	PRMC	Total profile soil moisture content		
M77	RZMC	Root zone soil moisture content		
M78	SFMC	Top soil layer soil moisture content		
M79	PRECTOT	Total surface precipitation		
M80	SNOMAS	Snow mass		
M81	EVPSOIL	Bare soil evaporation		
M82	EVPTRNS	Transpiration		
M83	EVPINTR	Interception loss		
M84	EVPSBLN	Sublimation		
M85	SMLAND	Snowmelt over land		
M86	EVLAND	Evaporation from land		

Table 2. (Continued)

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the overall top six variables found in the *Variable Screening* step, we created a final MaxEnt model in the *Model Calibration / Final Model Run* step that reflected any improvements gained in the *Predictor Refinement* step or by *Model Calibration*. The potential distribution maps produced by the final models were judged for reasonableness based on first-hand knowledge of the species, its habitat preferences, what is known about Cassin's Sparrow's range from the

published literature [52, 56–58, 61], and observational records from Cornell Lab's eBird citizen-scientist database [74].

We further compared these final model predictions to results obtained by replicating, in part, the work of Salas et al. [75], in which traditional MaxEnt variable-selection techniques were used to model the bioclimatic niche of Cassin's Sparrow. Here, we used our 2016 Cassin's Sparrow occurrence data in combination with the seven Worldclim Bioclim variables used by the Salas team: bio03, bio06, bio08, bio09, bio12, bio14, and bio18. The Salas team chose these predictors by first removing one of each pair of highly correlated variables to avoid collinearity among the variables. The team then chose between highly correlated variables by selecting those that were identified in one or more species-specific studies as influencing the species' range or population dynamics. In cases where the literature search could not differentiate between two highly correlated variables, the team used a qualitative assessment of the distribution of values of the variable at all presence points and the relationship between the variable and species presence or pseudo-absence [75]. We used ENMeval, as described above, to identify optimal tuning parameters for the Salas-derived model.

To gain a quantitative perspective on performance, we used AICc [64] as a measure of a model's relative explanatory power (lower values indicating less information loss) and area under the receiver operating characteristic curve (AUC) [76], percent correctly classified (PCC) [77], and the True Skill Statistic (TSS) [78, 79] as measures of model accuracy (higher values in all cases indicating greater accuracy). Similarities between our first use case's final Bioclim model and the Salas-derived Bioclim model were examined using Warren's *I* statistic [80], Schoener's *D* statistic [81], and Pearson's *r* statistic [82]. All input data used in this study, along with a set of example scripts are provided in S1 File.

Results

Run-time performance and scaling properties



The first ensemble of the 50-sample timing series (S = 50) required a total run time of T = 7.9 minutes to screen a two-variable collection (N = 2) using 10 processor cores (C = 10) (Fig 2A).

Fig 2. MERRA/Max run-time performance and scaling properties. Figure shows the relationship between the amount of time it takes MERRA/Max to complete a screening run (T) (shown by the left Y axis and the colored lines labeled A, B, and C), the number of variables in the collection being scanned (N), the average number of random samples taken of each variable in the collection during the screening process (S), and the number of processor cores available in the compute environment (C) (shown by the colored vertical bars and right Y axis). MERRA/Max's parallel implementation scales linearly with respect to S, and, for any given collection of size N and sample size S, the estimated minimum possible run time (T*min*) (shown in parentheses) can be achieved when enough cores are available for a completely parallel screening of the collection.

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At one sprint per core, and with each MaxEnt sampling run operating on two (V = 2) randomly selected variables at a time, this first ensemble needed 50 MaxEnt runs to do its work. The shortest achievable screening time (T*min*) is possible only when the number of cores needed for perfect parallelism (*Cmax*) are actually available, in this case, 50:

$$Cmax = (N \times S) \div V = (2 \times 50) \div 2 = 50.$$

Because only 10 cores were available, each of the parallel sprints had to perform five sequential MaxEnt sampling runs to achieve the S = 50 sampling goal, a repeat factor (R) of 5:

$$R = [Cmax \div C] = [50 \div 10] = 5.$$

By accounting for this performance cost, we estimate that MERRA/Max's minimum possible run time, in a completely parallel screening of this first data set, would have been about 1.6 minutes:

$$Tmin = T \div R = 7.9 \div 5 = 1.6$$
 minutes.

In each subsequent ensemble of the S = 50 series, we added two variables to the scanned collection and 10 cores to the pool of available processors. With this proportional scaling of variables and processors, average run times remained constant across the series at T = 7.9 ± 0.3 minutes (Tmin = 1.6 ± 0.1 minutes) (Fig 2A). In the S = 100 timing series, R = 10 sequential MaxEnt runs were used in each sprint to achieve the desired sampling level (Fig 2B), and in the S = 150 series, R = 15 runs were used (Fig 2C). In both cases, run times scaled linearly with sample size and remained relatively constant across the series, with T = 14.8 ± 2.5 minutes (Tmin = 1.6 ± 0.1 minutes) for the S = 100 series and T = 22.6 ± 0.5 minutes (Tmin = $1.5. \pm 0.1$ minutes) for the S = 150 series.

Use case scenarios and selection behavior

The Bioclim collection consists of N = 19 variables. To achieve an average per-variable sampling goal of S = 50 with C = 100 cores, each sprint in the Bioclim use case (Fig 3A) performed R = 5 sequential MaxEnt runs in the *Variable Screening* step, resulting in ensembles comprising a total of 500 runs. In an average of three such ensembles, MERRA/Max took T = 6.4 ± 0.5 minutes (Tmin = 1.3 ± 0.1 minutes) to identify bio18 (precipitation of the warmest quarter), bio03 (isothermality), bio05 (maximum temperature of the warmest month), bio08 (mean temperature of the wettest quarter), bio13 (precipitation of the wettest month), and bio16 (precipitation of the wettest quarter) as the top six contributing variables of the collection. In the subsequent *Variable Refinement* step, predictor pairs bio13-bio16 and bio16-bio18 were shown to be correlated, which led us to discard bio16 from the selection set. In the *Model Calibration / Final Model Run* step, the remaining five non-correlated variables were used to create a final model in which the top four contributing variables (bio18, bio03, bio05, and bio13) accounted for approximately 98% of overall permutation importance, and the performance metrics were AICc 12,232, AUC 0.83, PCC 0.75, and TSS 0.49.

In the MERRA-2 use case (Fig 3B), MERRA/Max screened a collection of N = 86 variables of coarser resolution (approximately 50 km for MERRA-2 vs. 8 km for Bioclim). To achieve the S = 50 sampling goal, each sprint performed R = 22 MaxEnt sampling runs, which resulted in 2200-run ensembles. The average run time across three such ensembles in the *Variable Screening* step increased to T = 18.0 \pm 2.2 minutes; however, because of the coarser predictor resolution, times for the MaxEnt sampling runs decreased, which resulted in an estimated theoretical lower bound of only T*min* = 0.8 \pm 0.1 minutes. The six top contributing variables



Fig 3. MERRA/Max use case scenarios. Figure shows the results of two use cases involving Cassin's Sparrow observational data and predictor data sets of contrasting size and complexity: the Bioclim collection with N = 19 variables (A) and a MERRA-2 reanalysis test collection comprising N = 86 variables (B). A Variable Screening step was used in each scenario to select the top six contributing variables in the underlying collection. Correlated variables (indicated with red text and yellow highlight) were identified in a Predictor Refinement step and thinned to reduce collinearities. In a third step, Model Calibration and a Final Model Run were performed with the remaining non-correlated variables (green highlight). AICc is Akaike's information criterion corrected for small sample size, AUC is area under the receiver operating characteristic curve, PCC is percent correctly classified, TSS is True Skill Statistic, Parameters is MaxEnt's measure of model complexity, r is Pearson's correlation coefficient, r² is the coefficient of determination, and VIF is variable inflation factor. The estimated minimum run time (Tmin) for a completely parallel screening is shown in parentheses. Maps created by the authors show MaxEnt logistic output, which can be interpreted as an estimate of habitat suitability between 0 and 1 with warmer colors indicating better predicted conditions for the species.

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identified in the Variable Screening step included M05 (specific humidity), M39 (surface net downward shortwave flux assuming a clear day), M38 (surface net downward shortwave flux), M81 (bare soil evaporation), M03 (northward wind), and M04 (temperature). In the Predictor *Refinement* step, the M38-M39 pair showed strong correlation, which led us to discard M38. In the Model Calibration / Final Model Run step, the remaining five non-correlated variables were used to create a final model in which the top four contributing variables (M39, M05, M04, and M81) accounted for approximately 97% of over overall permutation importance, and the performance metrics were AICc 7,023, AUC 0.83, PCC 0.72, and TSS 0.44.

In the Salas-derived Cassin's Sparrow model (Fig 4A), where a traditional approach to variable selection was used to identify the seven predictors used in MaxEnt, the top four contributing variables (bio18, bio06, bio14, and bio09) accounted for approximately 83% of overall permutation importance, and the model's performance metrics were AICc 12,169, AUC 0.83, PCC 0.76, and TSS 0.50.

Discussion

Climate change research is giving rise to new technology requirements at the intersection of big data, machine learning, and high-performance computing [84]. There are few places where this is more clearly seen than with studies focusing on the climate's impact on species distribution and abundance [85]. For nearly twenty years, the ecological modeling community's tool-of-choice for this work has been MaxEnt [22]. Few, if any, machine learning programs have been more widely used or more carefully studied [16, 18, 84, 86-92]. Today,



(A) Salas-derived Cassin's Sparrow MaxEnt Model

(B) Cassin's Sparrow Range Map

Fig 4. Cassin's Sparrow baseline model and maps. Figure shows results from a MaxEnt run that builds on the Cassin's Sparrow bioclimatic modeling work of Salas et al. [75] and reflects a more traditional approach to ENM (A) and Cassin's Sparrow's range map based on observational data (B). Highlighted variables indicate those that were also selected by MERRA/Max in the Bioclim use case. Range map provided by eBird (www.ebird.org), created 28 July 2020, and reprinted from [83] under a CC BY license, with permission from the Cornell Lab of Ornithology.

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however, there is increasing interest in using GCM outputs as predictors in ENM [10–12], which has brought into focus one of the more challenging problems with existing machine learning systems: how to make them work with large, complex, feature-rich, high-dimensional data sets [93–99].

This study reflects our efforts to address this problem. Our approach to dimensionality reduction involves a parallel, out-of-core Monte Carlo selection method implemented in a high-performance, cloud computing setting. Monte Carlo optimizations are a way of finding approximate answers to problems that are solvable in principle but lack a practical means of solution [100]. Out-of-core (or external memory) algorithms process data sets that are too large to fit into a computer's main memory [101, 102]. They are currently a major focus of research in the machine learning community [102–104]. With MERRA/Max, we bring these concepts together to find a useful subset of predictors in a large collection of environmental variables in a reasonable amount of time. Early results are encouraging and suggest that the approach holds promise from both a technological and scientific perspective.

Run-time performance and scaling properties

To begin, we have shown that MERRA/Max's parallel implementation of the Monte Carlo selection algorithm scales linearly with respect to the average number of samples taken of each variable in the collection being screened. It can recruit additional processor cores to maintain a constant run time regardless of the size of the collection. In a best case, where sufficient processors are available for complete parallelism, MERRA/Max can screen a predictor data set of any size in the time it takes for a single MaxEnt run using only two predictors in the target collection. Collectively, these results confirm that near real-time performance and, in the vernacular of high-performance cloud computing, "infinite scalability" are achievable [105–107].

Use case scenarios and selection behavior

MERRA/Max's run-time performance and scaling properties are consistent with what one would expect of an "embarrassingly parallel" workload, where subtasks are completely independent and able to run concurrently. The more important question is: Does this approach actually benefit science? We try to assess MERRA/Max's potential value to science by addressing three interrelated questions:

(1) Is MERRA/Max making useful, ecologically plausible selections?. It is difficult to know how best to evaluate selection success in this work, given that we are proposing the Monte Carlo method as a preliminary screening step, which, presumably, would be followed by further refinements to the set of selected predictors based on the biology and ecology of the species, collinearity, or the other considerations that have traditionally guided ENM variable selection. Depending on the circumstances, post-selection refinement might mean additional winnowing, substitution or augmentation of the screened predictors with other variables, or that the selected variables are discarded altogether. That being said, the Bioclim use case seems to confirm that MERRA/Max's selections are both valid and useful.

In the most general, qualitative sense, the habitat suitability map produced by the final model in the Bioclim use case is consistent with what is known about Cassin's Sparrow's range from observational records [83] (Figs <u>3A</u> and <u>4B</u>). Likewise, the set of selected variables are consistent with what is known of the species' natural history. Cassin's Sparrow is a desert-adapted, ground-dwelling (and, notably, ground-nesting) species, whose breeding biology is exquisitely linked to conditions of temperature and precipitation and their consequent influence on vegetation availability, insect abundance, and terrestrial microclimates [52, 56, 58, 61, <u>108–110</u>]. In fact, field studies over the past century suggest that Cassin's Sparrow is an itinerant breeder, so responsive to temperature and precipitation that they make seasonal, interclutch moves within their range to find optimal conditions for breeding [52, 109, 111].

The Bioclim scenario's ordered selection of bio18 (precipitation of the warmest quarter), bio03 (isothermality, i.e., temperature evenness, or how large the daily temperature variation is compared to its annual variation), bio05 (maximum temperature of the warmest month), bio13 (precipitation of the wettest month), and bio08 (mean temperature of the wettest quarter) is entirely consistent with this picture. It is also largely consistent with the variables assembled by the Salas team using a more traditional approach to variable selection [75]. Both sets have bio18 and bio03 in common, both of these variables are highly influential, and both are known to be important determinants of range in arid-adapted birds, especially in desert and grassland species of conservation concern [75, 85, 112–118]. Where the two predictor data sets differ, for example, bio05 (maximum temperature of the warmest month) in the Bioclim use case vs. bio06 (minimum temperature of the coldest month) in the Salas-derived model, bio13 (precipitation of the wettest month) vs. bio12 (annual precipitation) and bio 14 (precipitation of the driest month), an argument could be made, in light of Cassin's Sparrows distinctive seasonal breeding dynamics, which is likely influenced by the North American Monsoon, that MERRA/Max found the more relevant predictors [119–121].

From a quantitative perspective, the final model in MERRA/Max's Bioclim scenario (Fig 3A) demonstrated strong evaluation metrics (AUC, PCC, TSS of 0.83, 0.75, 0.49 respectively) that were almost identical to those obtained in the Salas-derived model (0.83, 0.76, 0.50) (Fig 4A) [122]. A high degree of similarity between the Bioclim use case and Salas-derived model is further confirmed by their AICc values (12,232 and 12,169 respectively) and the results we obtained for the *D*, *I*, and *r* statistics (0.974, 0.999, and 0.997 respectively) [80]. The top four predictors in the Bioclim scenario accounted for 98% of overall permutation importance in the final model; the top four predictors in the Salas-derived model accounted for 83%.

While these results reflect an admittedly limited trial at this point, taken together, they suggest that MERRA/Max's use in the Bioclim scenario produced a bioclimatic niche model for Cassin's Sparrow that, within its training range, is ecologically reasonable, statistically robust, and at least as good (if not better) than what might be obtained in a traditional application of MaxEnt. This gives us confidence that the Monte Carlo method is, in fact, finding a useful subset of predictors in a larger pool of possible predictors. (2) Does MERRA/Max create new research opportunities?. Another measure of MERRA/ Max's potential value to science is to consider whether new avenues of research are opened up with this technology. We believe they are, and the MERRA-2 use case helps explain why. Here we have a situation where the size and complexity of the target collection, as well as the obscure nature of the data, make *a priori* variable selection difficult. We have no immediate basis for distinguishing which variables in the MERRA-2 test collection might be the most important contributors to a final model. With 86 variables in play, even something as straightforward as correlation analysis provides little help, requiring nearly 3700 pair-wise comparisons in this case. Yet, going into this with no pre-vetting of the test collection whatsoever, we are struck by the ecological and biological relevance of the predictors selected by MERRA/Max in the MERRA-2 use case scenario.

In the same way that the Earth's climate is ultimately driven by a balance between incoming and outgoing energy, so too is the natural history of Cassin's Sparrow linked to the energetics of the species' diurnal and seasonal activities and the locations where those activities occur [58]. Viewed this way, it makes ecological sense to see that MERRA/Max identified M39 (surface net downward shortwave flux assuming a clear day) as the most important variable for modeling Cassin's Sparrow potential habitat in the MERRA-2 test collection. Likewise, a unique aspect of Cassin's Sparrow's breeding biology is an energetically demanding skylark display in which males define and defend territories and secure mates by aerial flight songs. Wind has a pronounced impact on this behavior [58]. While its effectiveness as a proxy for surface conditions in this setting is unknown, it is notable that MERRA/Max identified a lowlevel zonal wind component, M03 (northward wind), as a top contributor. Finally, given Cassin's Sparrow's ground-dwelling habit and the importance of low-level environmental conditions to almost all aspects of the species' life, it is not surprising to see M05 (specific humidity), M04 (temperature), and M81 (bare soil evaporation) in MERRA/Max's selection set.

These results must be interpreted with caution. After all, even models based on meaningless variables can be classified as excellent according to widely used evaluation metrics [123], and high predictive accuracy does not necessarily connote robust inferential capacity [17]. What is more, MERRA-2 variables represent the low-level physical drivers of many of the Earth system's biological processes [10, 124]: an interpretation of ecological plausibility could be made for almost any of the MERRA-2 variables. That being said, studies have shown that MaxEnt's ranking of variable importance can capture biologically realistic assessments of factors governing range boundaries when models are built using best-practice procedures and variables are ranked based on permutation importance [7, 17]. And, with Cassin's Sparrow, we have a species whose behavioral and energetic ecology has been studied in significant detail [58]. Of the many potential contributors in the MERRA-2 collection, the types of variables selected by MERRA/Max are known to be particularly important environmental influences for the species and are notably consistent with our mechanistic, process-based understanding of the bird's natural history [52, 56, 58, 61, 125]. Further, in the use case scenario's final MaxEnt model, we see a reasonable habitat suitability map based on our understanding of Cassin's Sparrow's current range (Fig 4B), a relatively robust set of metrics (AUC 0.83, PCC 0.72, TSS 0.44), and the top four contributing variables accounting for a significant proportion (i.e., 97%) of overall permutation importance (Fig 3B). While an interpretation of causality between selected variables and species occurrence may not be supported by the data currently at hand, it does appear that MERRA/Max and the Monte Carlo selection method have detected a signal in the MERRA-2 data that has both biological and statistical significance for Cassin's Sparrow [126, 127].

What about the larger question regarding new research opportunities? A particular type of question that might be better addressed with MERRA/Max's combination of data and

technology concerns the conservation status of arid-adapted birds. Accurate assessments are often difficult with these species [85, 112–118, 128]. With Cassin's Sparrow, for example, numerous studies over the past half-century have painted a confusing picture. Some find evidence for a retraction of viable habitat and declining regional populations [61], others find mixed results and too little data to establish with confidence an overall conservation status [56], and many sources identify the species as stable and of little immediate worry [129–132]. In recent work, of nine grassland birds of conservation concern, Cassin's Sparrow was the only species to project gains in suitable habitat over the next fifty years [75]. This ambiguous picture is not unique to Cassin's Sparrow; however, in this case, the species' itinerant breeding habit no doubt contributes to the confusion: it is simply impossible to know what part of the bird's population one is seeing at any given time.

Understanding the conservation status of a bird like Cassin's Sparrow means being able to tease apart the species' physiological capacity for seasonal response to weather from the species' longer-term, adaptive response to a changing climate's effect on the landscape [116]. Ultimately, one would like to distinguish short-term transformations in the bird's bioclimatic niche within a coherent, long-running temporal framework. Historical and multi-temporal scale modeling are not new [133–136]; however, with four decades of climate attributes modeled on an hourly basis, reanalyses, such as MERRA-2, are uniquely able to provide the hightemporal resolution, longitudinal environmental data for this. A technology like MERRA/Max transforms MERRA-2 into a viable experimental sandbox. And, thanks to long-running citizen-scientist efforts, such as the U.S. Geological Survey's North American Breeding Bird Survey (BBS) [137, 138]; Cornell University's eBird, Great Backyard Bird Count Surveys, and other projects [74, 139, 140]; Audubon's Christmas Bird Counts [141, 142]; and the wealth of online museum specimen records in resources like the Global Biodiversity Information Facility (GBIF) [143], the NSF-funded VertNet databases [144], and the U.S. Geological Survey's BISON species occurrence database [145], there now exists widespread availability of avian observations that provide good coverage for the MERRA-2 time span, making multi-temporal scale investigations like this possible [146].

Another dimension of conservation research that could potentially be advanced with a technology like MERA/Max is the modeling of rare or endangered species. Rare species are among the most in need of predictive distribution modeling but are often the most difficult to model. Known as the "rare species modeling paradox" [147], these species generally have a low number of occurrence records, which can lead to model over-parameterization and overfitting if too many predictors are used [11, 148, 149]. A new strategy using ensembles of small models (ESMs) was recently developed to overcome this limitation. It involves fitting many two-variable models, filtering the results against a weighted AUC-based performance threshold, then averaging the remaining models to produce an ensemble average model. The approach is particularly useful when applied to rare species, because it simultaneously winnows the starting pool of predictors while generating a final model in which the number of predictors has been kept low in each of the ensemble's contributing models [150, 151].

MERRA/Max also uses bivariate ensembles in its Monte Carlo approach to variable selection. In contrast to the ESM method, however, MERRA/Max discards its ensemble models after tallying the permutation importance of each model's two variables, producing in the end a small selection of top contributing variables for further consideration. By separating variable selection from final model construction, MERRA/Max provides the modeler with greater latitude in the overall ENM process, offering, in a sense, a supervised approach that could enable more carefully crafted results.

Finally, spatiotemporal projection is a critical element of conservation research that could also potentially benefit from MERRA/Max. ENMs are commonly used to predict the impact of

climate change on biodiversity. The reliability of those predictions, however, depends on a model's transferability in space and time, which, in turn, is influenced by variable selection [152–154]. While the work presented here has focused exclusively on performance evaluations within the calibration range, it is important to note that MERRA-2's selected variables, along with similar GCM outputs, form the basis for the IPCC's research activities. As a consequence, models using MERRA/Max-selected variables are particularly well-suited for extrapolative studies using data from IPCC's Global Projection scenarios [155–159].

We close this section by considering briefly the most apparent technical shortcoming in the MERRA-2 use case: the inherently coarse spatial resolution of reanalysis data. Given that species' responses to the environment are scale dependent, there is a recognized need within the ecological research community for higher resolution reanalysis products, to which various efforts are now responding [158, 160–162]. Increasing evidence, however, shows that both coarse and fine resolution variables are important across scales [163]. In the context of rapid prescreening, in particular, we feel that MERRA-2's coarse resolution data serves an important purpose. Selection times are fast with coarse-resolution data, MERRA/Max-selected variables are relevant, and many resolution shortcomings in the selected variables can be addressed in the refinement step, either by downscaling variables of interest or going to an alternative source for a higher resolution product, such as remote sensing data [164, 165] or NatureServe's high-resolution data sets [166].

(3) Can MERRA/Max improve the ecological niche modeling process?. Finally, in evaluating potential benefits to science, we can consider whether a tool like MERRA/Max could improve the work practices of ecological modeling. Here again, we think it can. There is heightened awareness of the significance of dimensionality in understanding environmental spaces and the importance of variable selection in modeling those spaces [15, 18, 167]. This awareness is accompanied by a recognition that logistic difficulties often preclude examining large numbers of variables, which has led to a search for alternative means of variable selection and calls for process automation [10, 17, 18, 88, 168]. A comprehensive review of these approaches is beyond the scope of this paper. In general, however, they include greater use of biological insight and expert knowledge in the selection of predictor variables; reliance on manual or statistical analysis of the published literature to identify predictors; use of statistical algorithms for variable prescreening based on cluster analysis, collinearity reduction, or calibration-/projection-range analogue analysis; and various types of classical principal component analysis (PCA) [153].

To understand where a technology fits within this conceptual framework, it is important to note that any selection process that involves human intervention is difficult or impossible to automate; the use of in-core statistical software tools are inherently unscalable to large data sets; whether implemented as in-core or out-of-core, compute-intensive, exponential time algorithms, such as PCA, are likewise not scalable; and mathematical approaches to feature reduction that operate on predictors in isolation from the feature of interest, i.e., the dependent variable, are not directly influenced by the underlying biology or ecology of the species being studied, which may limit the insights one might otherwise gain in the modeling process. MERRA/Max attempts to overcome all these limitations.

Finally, Cobos et al. [92] provide a helpful framework for understanding where a technology like MERRA/Max could fit in the overall ENM workflow (Fig 5). The work of ENM can be thought of as a multi-step process ranging from initial data preparation and cleaning, to model calibration, final model construction, model evaluation, and the assessment of extrapolation risk. Among the tasks associated with data cleaning, the selection of viable predictors is crucial, time-consuming, and the place where a means for rapid, automatic, preselection could

Initial data preparation	Data cleaning	Organize data	Model calibration	Final model construction and evaluation	Extrapolation risk analysis
Occurrence data Environmental data	Occurrence data filtering Reducing autocorrelation Calibration area delimitation (M) Variable preparation Variable selection	Prepare occurrences - Training datasets - Testing dataset - Full dataset (training + testing) - Independent set of data for final evaluation Predictors in the species M Multiple candidate sets	Candidate model creation Candidate model evaluation Model selection	Final models with no transfers Final models with transfers Final model evaluation	Identification of high- extrapolation risk areas

Ecological niche modeling process

Fig 5. Ecological niche modeling (ENM) process. Schematic description of the ENM process. Color bars under each step reflect an approximate amount of time that may be needed, ranging from low (blue) to high (red). The use of MERRA/Max to prescreen a large collection of predictors could support variable selection in the data cleaning step. Image provided by [92] and adapted for use here under a CC-BY license.

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improve the overall workflow, especially if it enabled exploration of a large universe of predictors.

Future work

Our plans for future work are being shaped by a vision where the technical complexities described here are abstracted away from the end-users, and low cost, easy access, and simplicity make MERRA/Max a practical and useful tool for the conservation research community. On the technology front, our next steps will focus on developing a "cloud bursting" capability that will allow MERRA/Max ensembles to migrate from NASA's private ADAPT science cloud to a public commercial cloud in response to resource demands that outstrip local capacity. This will allow us to scale MERRA/Max to larger data sets and more demanding science questions. For example, we would like to make better use of our experimental MERRA-2 test collection, which spans 40 years and includes monthly and weekly maximum, minimum, and average values for each of the collection's 86 variables, a total of N = 660,480 files. Beyond that, expanding MERRA/Max to accommodate all of MERRA-2's 600-plus variables is a more challenging long-term goal that could open interesting new research opportunities for the ENM community. Other technical improvements on the horizon include the use of refined selection criteria and automatic stopping rules that would enable true convergence in the selection process. Finally, given that each step in our use case scenarios is carried out by a program that could readily participate in an orchestrated workflow, we would like to know whether a practical level of automation for the entire MaxEnt-enabled ENM process might be possible.

On the science front, we plan to pursue two distinct threads of development. First, we want to follow up on the example posed in the previous section and see if MERRA/Max and the MERRA-2 test collection could provide a better understanding of the true conservation status of Cassin's Sparrow by temporally filtering occurrence data and environmental predictors to the breeding season and evaluating fine-grained changes in the species' bioclimatic niche over the past four decades. Pursuing that question for Cassin's Sparrow alongside other grassland birds of conservation concern, such as those studied by Salas et al. [75], would provide the additional benefit of extending our experiences to other species and would allow us to test the ecological and conservation applicability of this technology, which we view as an important next step.

The second thread of science development will examine the extensibility of this approach to other types of problems. For example, we have used MERRA/Max and MaxEnt to study the hydrological cycle in NASA's Arctic-Boreal Vulnerability Experiment (ABoVE) [169]. Changes to the hydrological cycle in the Arctic are particularly complex, because observed and projected warming directly impacts permafrost and leads to variable responses in surface water extent [170]. In preliminary work, using the locations of observed increases and decreases in surface water extent as dependent variables (in essence, treating them as "pseudo" species occurrences), the technologies and techniques described here successfully replicated observed patterns of surface water change [171]. If these findings are validated by further experimentation, the view of how MERRA/Max, the MERRA-2 reanalysis, and MaxEnt might be applied to studies of climate change and its impact on the Earth system becomes significantly broadened.

Conclusions

In this paper, we have described a prototype system called MERRA/Max that implements a feature selection approach to dimensionality reduction that is specifically intended to enable direct use of GCM outputs in ENM. The system accomplishes this reduction through a Monte Carlo optimization in which many independent MaxEnt runs operating on a species occurrence file and a small set of variables randomly selected from a large collection of variables converges on an estimate of the top contributing predictors in the larger collection. These top predictors become candidates for consideration in the variable selection step of the ENM process. MERRA/Max's Monte Carlo algorithm operates on files stored in the underlying filesystem and is thus scalable to large data sets. We implemented its program components using open-source and commercial off-the-shelf software. These components can run independently as parallel processes in a high-performance cloud computing environment to yield near real-time performance.

Within this framework, variable selection is guided by the indirect biological influences injected into MERRA/Max's feature reduction process by the species occurrence files. We find evidence for this tailoring of results in our use case scenarios. In preliminary tests using a single bird species and observations from a single year, MERRA/Max selected reasonable and familiar climatological predictors from the classic Bioclim collection of environmental variables. MERRA/Max also selected biologically and ecologically plausible predictors from a larger and much more diverse set of environmental variables derived from NASA's MERRA-2 reanalysis. Our experience is limited at this point, but we feel that these results point to a technological approach that could expand the use of GCM outputs in ENM, foster exploratory experimentation with otherwise difficult-to-use climate data sets, streamline the modeling process, and, eventually, enable automated bioclimatic modeling as a cloud service.

Supporting information

S1 Appendix. MERRA/Max parameterization. Overview of MERRA/Max's default screening parameters and supporting documentation. (PDF)

S1 File. Data and scripts. Compressed file folder containing input data and example scripts used in the study. (ZIP)

(ZIP

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References

- 1. Peterson AT, Soberón J, Pearson RG, Anderson RP, Martínez-Meyer E, Nakamura M, et al. Ecological niches and geographic distributions (MPB-49). Princeton University Press; 2011.
- Schmidt-Lebuhn AN, Knerr NJ, Miller JT, Mishler BD. Phylogenetic diversity and endemism of Australian daisies (Asteraceae). Journal of Biogeography. 2015; 42: 1114–1122. https://doi.org/10.1111/jbi. 12488
- Cardoso-Leite R, Vilarinho AC, Novaes MC, Tonetto AF, Vilardi GC, Guillermo-Ferreira R. Recent and future environmental suitability to dengue fever in Brazil using species distribution model. Transactions of The Royal Society of Tropical Medicine and Hygiene. 2014; 108: 99–104. <u>https://doi.org/10.1093/ trstmh/trt115</u> PMID: 24463584
- Franklin J. Species distribution models in conservation biogeography: developments and challenges. Diversity Distrib. 2013; 19: 1217–1223. https://doi.org/10.1111/ddi.12125
- Jiménez-Valverde A, Peterson AT, Soberón J, Overton JM, Aragón P, Lobo JM. Use of niche models in invasive species risk assessments. Biol Invasions. 2011; 13: 2785–2797. <u>https://doi.org/10.1007/</u> s10530-011-9963-4
- Muttaqin LA, Murti SH, Susilo B. MaxEnt (Maximum Entropy) model for predicting prehistoric cave sites in Karst area of Gunung Sewu, Gunung Kidul, Yogyakarta. In: Wibowo SB, Rimba AB, A. Aziz A, Phinn S, Sri Sumantyo JT, Widyasamratri H, et al., editors. Sixth Geoinformation Science Symposium. Yogyakarta, Indonesia: SPIE; 2019. p. 3. https://doi.org/10.1117/12.2543522
- Searcy CA, Shaffer HB. Do Ecological Niche Models Accurately Identify Climatic Determinants of Species Ranges? The American Naturalist. 2016; 187: 423–435. <u>https://doi.org/10.1086/685387</u> PMID: 27028071
- Harris RMB, Grose MR, Lee G, Bindoff NL, Porfirio LL, Fox-Hughes P. Climate projections for ecologists: Climate projections for ecologists. Wiley Interdisciplinary Reviews: Climate Change. 2014; 5: 621–637. https://doi.org/10.1002/wcc.291

- Edwards PN. A Vast Machine: Computer Models, Climate Data, and the Politics of Global Warming. Cambridge, MA: MIT Press; 2010.
- Cavanagh RD, Murphy EJ, Bracegirdle TJ, Turner J, Knowland CA, Corney SP, et al. A Synergistic Approach for Evaluating Climate Model Output for Ecological Applications. Frontiers in Marine Science. 2017; 4: 308. https://doi.org/10.3389/fmars.2017.00308
- Radosavljevic A, Anderson RP. Making better MaxEnt models of species distributions: complexity, overfitting and evaluation. Araújo M, editor. Journal of Biogeography. 2014; 41: 629–643. https://doi. org/10.1111/jbi.12227
- Heinze G, Wallisch C, Dunkler D. Variable selection–A review and recommendations for the practicing statistician. Biometrical Journal. 2018; 60: 431–449. https://doi.org/10.1002/bimj.201700067 PMID: 29292533
- Duan Y, Edwards JS, Dwivedi YK. Artificial intelligence for decision making in the era of Big Data–evolution, challenges and research agenda. International Journal of Information Management. 2019; 48: 63–71. https://doi.org/10.1016/j.ijinfomgt.2019.01.021
- Schnase JL, Carroll ML, Gill RL, Tamkin GS, Li J, Strong SL, et al. Toward a Monte Carlo approach to selecting climate variables in MaxEnt. PLOS ONE. 2021; 16: e0237208. <u>https://doi.org/10.1371/journal.pone.0237208</u> PMID: 33657125
- Araújo MB, Guisan A. Five (or so) challenges for species distribution modelling. Journal of Biogeography. 2006; 33: 1677–1688. https://doi.org/10.1111/j.1365-2699.2006.01584.x
- Araújo MB, Anderson RP, Barbosa AM, Beale CM, Dormann CF, Early R, et al. Standards for distribution models in biodiversity assessments. Science Advances. 2019; 5: eaat4858. <u>https://doi.org/10. 1126/sciadv.aat4858 PMID: 30746437</u>
- Smith AB, Santos MJ. Testing the ability of species distribution models to infer variable importance. Ecography. 2020; 43: 1801–1813. https://doi.org/10.1111/ecog.05317
- Cobos ME, Peterson AT, Osorio-Olvera L, Jiménez-García D. An exhaustive analysis of heuristic methods for variable selection in ecological niche modeling and species distribution modeling. Ecological Informatics. 2019; 53: 100983. https://doi.org/10.1016/j.ecoinf.2019.100983
- Peterson AT, Cobos ME, Jiménez-García D. Major challenges for correlational ecological niche model projections to future climate conditions: Climate change, ecological niche models, and uncertainty. Annals of the New York Academy of Sciences. 2018; 1429: 66–77. <u>https://doi.org/10.1111/nyas.</u> 13873 PMID: 29923606
- Elith J, Phillips SJ, Hastie T, Dudík M, Chee YE, Yates CJ. A statistical explanation of MaxEnt for ecologists. Diversity and distributions. 2011; 17: 43–57.
- 21. Phillips SJ, Anderson RP, Dudík M, Schapire RE, Blair ME. Opening the black box: An open-source release of Maxent. Ecography. 2017; 40: 887–893.
- 22. Phillips SJ. A Brief Tutorial on Maxent. AT&T Research. 2005; 190: 231–259.
- Gelaro R, Mccarty W, Su MJ, Todling R, Molod A, Takacs L, et al. The Modern-Era Retrospective Analysis for Research and Applications, Version 2 (MERRA-2). JOURNAL OF CLIMATE. 2017; 30: 36. https://doi.org/10.1175/JCLI-D-16-0758.1 PMID: 32020988
- 24. Reanalyses.org Home Page. 2021 [cited 12 Mar 2021]. Available: https://reanalyses.org/
- Schnase JL, Smith JA, Stohlgren TJ, Graves S, Trees C. Biological Invasions: a Challenge in Ecological Forecasting. IEEE International Geoscience and Remote Sensing Symposium. 2002. https://doi. org/10.1109/igarss.2002.1024961
- Beauchamp VB, Koontz SM, Suss C, Hawkins C, Kyde KL, Schnase JL. An Introduction to Oplismenus Undulatifolius (Ard.) Roem. & Schult (Wavyleaf Basketgrass), a Recent Invader in Mid-Atlantic Forest Understories 1,2. The Journal of the Torrey Botanical Society. 2013; 140: 391–413. https://doi. org/10.3159/torrey-d-13-00033.1
- Morisette JT, Jarnevich CS, Ullah A, Cai W, Pedelty JA, Gentle JE, et al. A Tamarisk Habitat Suitability Map for the Continental United States. Frontiers in Ecology and the Environment. 2006; 4: 11–17. https://doi.org/10.1890/1540-9295(2006)004
- Schnase JL, Carroll ML, Weber KT, Brown ME, Gill RL, Wooten M, et al. RECOVER: An Automated, Cloud-Based Decision Support System for Post-Fire Rehabilitation Planning. ISPRS—International Archives of the Photogrammetry, Remote Sensing and Spatial Information Sciences XL-1. 2014. pp. 363–70. https://doi.org/10.5194/isprsarchives-xl-1-363-2014
- 29. Schnase JL, Most N, Gill R, Ma P. The Invasive Species Forecasting System. 2009 17th International Conference on Geoinformatics. 2009. https://doi.org/10.1109/geoinformatics.2009.5293333
- Hu F, Yang C, Schnase JL, Duffy DQ, Xu M, Bowen MK, et al. ClimateSpark: An in-memory distributed computing framework for big climate data analytics. Computers & Geosciences. 2018; 115: 154–166. https://doi.org/10.1016/j.cageo.2018.03.011

- Schnase JL, Lee TJ, Mattmann CA, Lynnes CS, Cinquini L, Ramirez P. M., et al. Big Data Challenges in Climate Science: Improving the next-generation cyberinfrastructure. IEEE Geoscience and Remote Sensing Magazine. 2016; 4: 10–22. https://doi.org/10.1109/MGRS.2015.2514192 PMID: 31709380
- 32. Schnase JL. Climate Analytics as a Service. Cloud Computing in Ocean and Atmospheric Sciences. 2016. pp. 187–219. https://doi.org/10.1016/b978-0-12-803192-6.00011–6
- Schnase JL, Cushing J, Frame M, Frondorf A, Landis E, Maier D, et al. Information technology challenges of biodiversity and ecosystems informatics. Information Systems. 2003; 28: 339–345. <u>https://doi.org/10.1016/S0306-4379(02)00070-4</u>
- Schnase JL, Duffy DQ, McInerney MA, Webster WP, Lee TJ. Climate Analytics as a Service. Proceedings of the 2014 Conference on Big Data from Space (BiDS. Frascati: European Space Agency (ESA; 2014. pp. 90–94. https://doi.org/10.2788/1823
- Schnase JL, Duffy DQ, Tamkin GS, Nadeau D, Thompson JH, Grieg CM, et al. MERRA Analytic Services: Meeting the Big Data Challenges of Climate Science through Cloud-Enabled Climate Analyticsas-a-Service. Computers, Environment and Urban Systems. 2017; 61: 198–211. <u>https://doi.org/10.1016/j.compenvurbsys.2013.12.003</u>
- Carriere L, Potter GL, Hertz J, Peters J, Maxwell TP, Strong S, et al. CREATE-IP and CREATE-V: Data and Services Update. AGU Fall Meeting Abstracts. 2017. pp. IN21D-0064.
- Cinquini L, Crichton D, Mattmann C, Harney J, Shipman G, Wang F, et al. The Earth System Grid Federation: An open infrastructure for access to distributed geospatial data. Future Generation Computer Systems. 2014; 36: 400–417.
- Maxwell TP, Potter GL, Carriere L, Duffy D. The Earth Data Analytic Services Framework. AGU Fall Meeting Abstracts. 2019. pp. IN13B-0720.
- Tamkin G, Schnase JL, Duffy D, Li J, Strong S, Thompson JH. The NASA Reanalysis Ensemble Service-Advanced Capabilities for Integrated Reanalysis Access and Intercomparison. AGU Fall Meeting Abstracts. 2017. pp. IN21D-0065.
- GES DISC—Goddard Earth Science Data and Information Services Center. [cited 26 May 2021]. Available: https://disc.gsfc.nasa.gov/
- NASA Case Study–Amazon Web Services (AWS). In: Amazon Web Services, Inc. [Internet]. 2021 [cited 26 May 2021]. Available: https://aws.amazon.com/partners/success/nasa-image-library/
- 42. Research and Technical Computing on Amazon Web Services (AWS). In: Amazon Web Services, Inc. [Internet]. 2021 [cited 26 May 2021]. Available: https://aws.amazon.com/government-education/ research-and-technical-computing/
- Google Cloud offers global support for academic research. In: Google [Internet]. 2019 [cited 26 May 2021]. Available: https://blog.google/products/google-cloud/google-cloud-offers-global-support-foracademic-research/
- 44. Our head's in the cloud, but we're keeping the earth in mind. In: Google Cloud Blog [Internet]. 2019 [cited 26 May 2021]. Available: https://cloud.google.com/blog/topics/google-cloud-next/our-heads-inthe-cloud-but-were-keeping-the-earth-in-mind/
- Cloud Computing Services | Microsoft Azure. 2021 [cited 26 May 2021]. Available: https://azure. microsoft.com/en-us/
- 46. Data Science Virtual Machines | Microsoft Azure. 2021 [cited 26 May 2021]. Available: https://azure. microsoft.com/en-us/services/virtual-machines/data-science-virtual-machines/
- 47. ADAPT. In: ADAPT | NASA Center for Climate Simulation [Internet]. [cited 15 Mar 2021]. Available: https://www.nccs.nasa.gov/systems/ADAPT
- 48. R: The R Project for Statistical Computing. [cited 22 May 2020]. Available: https://www.r-project.org/
- 49. Muscarella R, Galante PJ, Soley-Guardia M, Boria RA, Kass JM, Anderson MU and RP. ENMeval: Automated Runs and Evaluations of Ecological Niche Models. 2020. Available: <u>https://CRAN.R-project.org/package=ENMeval</u>
- 50. Muscarella R, Galante PJ, Soley-Guardia M, Boria RA, Kass JM, Uriarte M, et al. ENMeval: An R package for conducting spatially independent evaluations and estimating optimal model complexity for Maxent ecological niche models. Methods in Ecology and Evolution. 2014; 5: 1198–1205. https://doi.org/10.1111/2041-210X.12261
- 51. Maxent Version 3.4.1 Download Site. In: Maxent Version 3.4.1 Download Site [Internet]. [cited 22 May 2020]. Available: https://biodiversityinformatics.amnh.org/open_source/maxent/
- Dunning, Jr. JB, Bowers, Jr. RK, Suter SJ, Bock CE. Cassin's Sparrow (Peucaea cassinii), Version 1.0. In: Birds of the World (P. G. Rodewald, Editor) [Internet]. 2020 [cited 22 May 2020]. Available: https://doi.org/10.2173/bow.casspa.01

- **53.** Fick SE, Hijmans RJ. WorldClim 2: new 1-km spatial resolution climate surfaces for global land areas. International Journal of Climatology. 2017; 37: 4302–4315. https://doi.org/10.1002/joc.5086
- 54. Worldclim bioclimatic variables. 2020 [cited 22 May 2020]. Available: https://worldclim.org/data/ worldclim21.html
- 55. GBIF.org (21 February 2019) GBIF Occurrence Download https://doi.org/10.15468/dl.0s8yak.
- Ruth JM. Cassin's Sparrow Status Assessment and Conservation Plan. Biological Technical Publication BTP-R6002-2000. Denver, CO: U.S. Department of the Interior, Fish and Wildlife Service; 2000.
- 57. Schnase JL, Maxwell TC. Use of song patterns to identify individual male Cassin's Sparrows. Journal of Field Ornithology. 1989; 60: 12–19.
- Schnase JL, Grant WE, Maxwell TC, Leggett JJ. Time and energy budgets of Cassin's sparrow (Aimophila cassinii) during the breeding season: evaluation through modelling. Ecological Modelling. 1991; 55: 285–319.
- 59. GDAL/OGR Geospatial Data Abstraction Software Library. Open Source Geospatial Foundation; 2020. Available: https://gdal.org/
- Hijmans RJ, Phillips S, Elith J, Leathwick J. dismo: Species Distribution Modeling. 2017. Available: https://CRAN.R-project.org/package=dismo
- **61.** Lynn J. Cassin's Sparrow (Aimophila cassinii): A Technical Conservation Assessment. USDA Forest Service, Species Conservation Project. Rocky Mountain Region. 2006; 46.
- **62.** Pradhan P. Strengthening MaxEnt modelling through screening of redundant explanatory bioclimatic variables with variance inflation factor analysis. Researcher. 2016; 8: 29–34.
- Warren DL, Glor RE, Turelli M. ENMTools: a toolbox for comparative studies of environmental niche models. Ecography. 2010 [cited 27 Mar 2020]. https://doi.org/10.1111/j.1600-0587.2009.06142.x
- Akaike H. A new look at the statistical model identification. IEEE Transactions on Automatic Control. 1974; 19: 716–723. https://doi.org/10.1109/TAC.1974.1100705
- Intergovernmental Panel on Climate Change (IPCC). 2021 [cited 14 Mar 2021]. Available: https://www.ipcc.ch/
- Bosilovich MG, Lucchesi R, Suarez M. MERRA-2: File Specification. GMAO Office Note. 2016; 9: 1– 73.
- 67. MERRA-2. 2020 [cited 19 Mar 2020]. Available: https://gmao.gsfc.nasa.gov/reanalysis/MERRA-2/
- Beaumont LJ, Hughes L, Poulsen M. Predicting species distributions: use of climatic parameters in BIOCLIM and its impact on predictions of species' current and future distributions. Ecological Modelling. 2005; 186: 251–270. https://doi.org/10.1016/j.ecolmodel.2005.01.030
- Booth TH, Nix HA, Busby JR, Hutchinson MF. BIOCLIM: the first species distribution modelling package, its early applications and relevance to most current MaxEnt studies. Franklin J, editor. Diversity Distrib. 14AD; 20: 1–9. https://doi.org/10.1111/ddi.12144
- Hijmans RJ, Cameron SE, Parra JL, Jones PG, Jarvis A. Very high resolution interpolated climate surfaces for global land areas. International Journal of Climatology. 2005; 25: 1965–1978. https://doi.org/ 10.1002/joc.1276
- 71. O'Donnell MS, Ignizio D a. Bioclimatic Predictors for Supporting Ecological Applications in the Conterminous United States. Reston, VA: US Geological Survey; 2012 p. 10. Report No.: 691. Available: https://pubs.usgs.gov/ds/691/
- 72. Vega GC, Pertierra LR, Olalla-Tárraga MÁ. Data from: MERRAclim, a high-resolution global dataset of remotely sensed bioclimatic variables for ecological modelling. Dryad; 2018. p. 8324359732 bytes. https://doi.org/10.1038/sdata.2018.70 PMID: 29664471
- 73. Hoyer S, Hamman JJ. xarray: N-D labeled Arrays and Datasets in Python. Journal of Open Research Software. 2017; 5: 10. https://doi.org/10.5334/jors.148
- (Peucaea cassinii)—Species Map—eBird. 2020 [cited 31 May 2020]. Available: <u>https://ebird.org/map/casspa</u>
- 75. Salas EAL, Seamster VA, Boykin KG, Harings NM, Dixon KW, Department of Fish, Wildlife and Conservation Ecology, New Mexico State University, Las Cruces, New Mexico 88003, USA. Modeling the impacts of climate change on Species of Concern (birds) in South Central U.S. based on bioclimatic variables. AIMS Environmental Science. 2017; 4: 358–385. https://doi.org/10.3934/environsci.2017.2. 358
- 76. Fielding AH, Bell JF. A review of methods for the assessment of prediction errors in conservation presence/absence models. Environmental Conservation. 1997; 24: 38–49. <u>https://doi.org/10.1017/S0376892997000088</u>

- Warren DL, Seifert SN. Ecological niche modeling in Maxent: the importance of model complexity and the performance of model selection criteria. Ecological Applications. 2011; 21: 335–342. https://doi. org/10.1890/10-1171.1 PMID: 21563566
- 78. Allouche O, Tsoar A, Kadmon R. Assessing the accuracy of species distribution models: prevalence, kappa and the true skill statistic (TSS): Assessing the accuracy of distribution models. Journal of Applied Ecology. 2006; 43: 1223–1232. https://doi.org/10.1111/j.1365-2664.2006.01214.x
- 79. Frans VF. True Skill Statistic (TSS) Calculation across Multiple Maxent Runs. Michigan State University; 2018 p. 5.
- Warren DL, Glor RE, Turelli M. Environmental niche equivalency versus conservatism: quantitative approaches to niche evolution. Evolution. 2008; 62: 2868–2883. https://doi.org/10.1111/j.1558-5646. 2008.00482.x PMID: 18752605
- Schoener TW. The Anolis Lizards of Bimini: Resource Partitioning in a Complex Fauna. Ecology. 1968; 49: 704–726. https://doi.org/10.2307/1935534
- Rodgers JL, Nicewander WA. Thirteen Ways to Look at the Correlation Coefficient. The American Statistician. 1988; 42: 59–66. https://doi.org/10.2307/2685263
- Fink D, Auer T, Johnston A, Strimas-Mackey M, Robinson O, Ligocki S, et al. Cassin's Sparrow— Abundance map—eBird Status and Trends. In: eBird Status and Trends, Data Version: 2018; Released: 2020 [Internet]. 2020 [cited 5 Oct 2020]. Available: <u>https://ebird.org/ebird/science/status-and-trends/casspa/abundance-map</u>
- Araújo M, New M. Ensemble forecasting of species distributions. Trends in Ecology & Evolution. 2007; 22: 42–47. https://doi.org/10.1016/j.tree.2006.09.010 PMID: 17011070
- Huntley B, Barnard P, Altwegg R, Chambers L, Coetzee BWT, Gibson L, et al. Beyond bioclimatic envelopes: dynamic species' range and abundance modelling in the context of climatic change. Ecography. 2010 [cited 13 Mar 2020]. https://doi.org/10.1111/j.1600-0587.2009.06023.x
- Feng X, Park DS, Walker C, Peterson AT, Merow C, Papeş M. A checklist for maximizing reproducibility of ecological niche models. Nature Ecology & Evolution. 2019; 3: 1382–1395. <u>https://doi.org/10. 1038/s41559-019-0972-5 PMID: 31548646</u>
- Morales NS, Fernández IC, Baca-González V. MaxEnt's parameter configuration and small samples: are we paying attention to recommendations? A systematic review. PeerJ. 2017; 5: e3093. <u>https://doi.org/10.7717/peerj.3093</u> PMID: 28316894
- Zeng Y, Low BW, Yeo DCJ. Novel methods to select environmental variables in MaxEnt: A case study using invasive crayfish. Ecological Modelling. 2016; 341: 5–13. <u>https://doi.org/10.1016/j.ecolmodel.</u> 2016.09.019
- Qiao H, Soberón J, Peterson AT. No silver bullets in correlative ecological niche modelling: insights from testing among many potential algorithms for niche estimation. Kriticos D, editor. Methods in Ecology and Evolution. 2015; 6: 1126–1136. https://doi.org/10.1111/2041-210X.12397
- Ashraf U, Peterson AT, Chaudhry MN, Ashraf I, Saqib Z, Rashid Ahmad S, et al. Ecological niche model comparison under different climate scenarios: a case study of *Olea* spp. in Asia. Ecosphere. 2017; 8: e01825. https://doi.org/10.1002/ecs2.1825
- Guisan A, Zimmermann NE. Predictive habitat distribution models in ecology. Ecological modelling. 2000; 135: 147–186.
- Cobos ME, Peterson AT, Barve N, Osorio-Olvera L. kuenm: an R package for detailed development of ecological niche models using Maxent. PeerJ. 2019; 7: e6281. https://doi.org/10.7717/peerj.6281 PMID: 30755826
- Qiu J, Wu Q, Ding G, Xu Y, Feng S. A survey of machine learning for big data processing. EURASIP J Adv Signal Process. 2016; 2016: 67. https://doi.org/10.1186/s13634-016-0355-x
- Obermeyer Z, Emanuel EJ. Predicting the Future—Big Data, Machine Learning, and Clinical Medicine. N Engl J Med. 2016; 375: 1216–1219. https://doi.org/10.1056/NEJMp1606181 PMID: 27682033
- 95. Bailly S, Meyfroidt G, Timsit J-F. What's new in ICU in 2050: big data and machine learning. Intensive Care Medicine. 2018; 44: 1524–1527. https://doi.org/10.1007/s00134-017-5034-3 PMID: 29279970
- 96. van der Maaten L, Postma E, van den Herik J. Dimensionality Reduction: A Comparative Review. The Netherlands: Tilburg University; 2009 p. 36. Report No.: TiCC TR 2009–005. Available: <u>https://members.loria.fr/moberger/Enseignement/AVR/Exposes/TR_Dimensiereductie.pdf</u>
- Guyon I, Elisseeff A. An Introduction to Variable and Feature Selection. Journal of Machine Learning Research. 2003; 3: 1157–1182.
- Espadoto M., Martins R. M., Kerren A., Hirata N. S. T., Telea A. C. Towards a Quantitative Survey of Dimension Reduction Techniques. IEEE Transactions on Visualization and Computer Graphics. 2019; 1–1. https://doi.org/10.1109/TVCG.2019.2944182 PMID: 31567092

- 99. Feng X, Park DS, Liang Y, Pandey R, Papeş M. Collinearity in ecological niche modeling: Confusions and challenges. Ecology and Evolution. 2019; 9: 10365–10376. <u>https://doi.org/10.1002/ece3.5555</u> PMID: 31624555
- 100. Kroese DP, Brereton T, Taimre T, Botev ZI. Why the Monte Carlo method is so important today: Why the MCM is so important today. Wiley Interdisciplinary Reviews: Computational Statistics. 2014; 6: 386–392. https://doi.org/10.1002/wics.1314
- 101. Ito Y, Imai H, Duc TL, Negishi Y, Kawachiya K, Matsumiya R, et al. Profiling based Out-of-core Hybrid Method for Large Neural Networks. arXiv:190705013 [cs]. 2019 [cited 11 Jan 2021]. Available: <u>http://arxiv.org/abs/1907.05013</u>
- 102. Chen T, Xu B, Zhang C, Guestrin C. Training Deep Nets with Sublinear Memory Cost. arXiv:160406174 [cs]. 2016 [cited 11 Jan 2021]. Available: http://arxiv.org/abs/1604.06174
- 103. Hanlon J. How To Solve The Memory Challenges Of Deep Neural Networks. In: TOPBOTS [Internet]. 2017 [cited 13 May 2021]. Available: https://www.topbots.com/how-solve-memory-challenges-deep-learning-neural-networks-graphcore/
- 104. Zhang B. A Solution to the Memory Limit Challenge in Big Data Machine Learning. In: Medium [Internet]. 2018 [cited 13 May 2021]. Available: https://petuum.medium.com/a-solution-to-the-memory-limitchallenge-in-big-data-machine-learning-49783a72088b
- 105. Bicer T, Chiu D, Agrawal G. A Framework for Data-Intensive Computing with Cloud Bursting. 2011 IEEE International Conference on Cluster Computing. 2011. pp. 169–177. <u>https://doi.org/10.1109/</u> CLUSTER.2011.21
- 106. Pham B, Jones RC, Shalaan M. Analysis of Cloud Bursting from the Openstack Infrastructure to AWS. 2020 IEEE Cloud Summit. 2020. pp. 114–118. https://doi.org/10.1109/IEEECloudSummit48914. 2020.00037
- Guo T, Sharma U, Shenoy P, Wood T, Sahu S. Cost-Aware Cloud Bursting for Enterprise Applications. ACM Trans Internet Technol. 2014;13. https://doi.org/10.1145/2602571
- 108. Oberholser HC. Cassin's Sparrow, "Aimophila cassinii" (Woodhouse). 1st ed. In: Kincaid EB Jr., editor. Bird Life of Texas. 1st ed. Austin: University of Texas Press; 1974. pp. 920–921.
- 109. Williams FC, LeSassier AL. Cassin's Sparrow. In: Austin OL, editor. Life Histories of North American Cardinals, Grosbeaks, Buntings, Towhees, Finches, Sparrows, and Allies, Order Passeriformes, Family Fringillidae: (in 3vols) Part 2, Genera Pipilo (part) Through Spizella. New York: Dover; 1968. pp. 981–990.
- 110. Woodhouse SW. Zonotrichia Cassinii, nobis. Proceedings of the Academy of Natural Science of Philadelphia. Philadelphia: Merrihow and Thompson; 1852. pp. 60–61. Available: <u>https://www.biodiversitylibrary.org/item/17888#page/7/mode/1up</u>
- 111. Ohmart RD. Dual breeding ranges in Cassin's sparrow (Aimophila cassinii). In: Hoff CC, Riedesel ML, editors. Physiological systems in semiarid environments. Albuquerque, NM: University of New Mexico Press; 1969. p. 105.
- 112. Norman JA, Christidis L. Ecological opportunity and the evolution of habitat preferences in an aridzone bird: implications for speciation in a climate-modified landscape. Scientific Reports. 2016; 6: 19613. https://doi.org/10.1038/srep19613 PMID: 26787111
- 113. Hubbard JP. Avian evolution in the aridlands of North America. The Living Bird. 1974; 155–196.
- 114. Sohl TL. The Relative Impacts of Climate and Land-Use Change on Conterminous United States Bird Species from 2001 to 2075. Romanach SS, editor. PLoS ONE. 2014; 9: e112251. https://doi.org/10. 1371/journal.pone.0112251 PMID: 25372571
- 115. Rosenberg KV, Dokter AM, Blancher PJ, Sauer JR, Smith AC, Smith PA, et al. Decline of the North American avifauna. Science. 2019; 366: 120. <u>https://doi.org/10.1126/science.aaw1313</u> PMID: 31604313
- 116. Reside AE, VanDerWal JJ, Kutt AS, Perkins GC. Weather, Not Climate, Defines Distributions of Vagile Bird Species. Hector A, editor. PLoS ONE. 2010; 5: e13569. <u>https://doi.org/10.1371/journal.pone.</u> 0013569 PMID: 21042575
- 117. Iknayan KJ, Beissinger SR. Collapse of a desert bird community over the past century driven by climate change. Proc Natl Acad Sci USA. 2018; 115: 8597. https://doi.org/10.1073/pnas.1805123115 PMID: 30082401
- Heenan CB, Seymour RS. The Effect of Wind on the Rate of Heat Loss from Avian Cup-Shaped Nests. Brigham RM, editor. PLoS ONE. 2012; 7: e32252. <u>https://doi.org/10.1371/journal.pone.</u> 0032252 PMID: 22389689
- 119. Liebmann B. Characteristics of North American Summertime Rainfall with Emphasis on the Monsoon. American Meteorological Society Jounal of Climate. 2008; 21: 1277–1294.

- 120. NOAA. The North American Monsoon. NOAA NWS Climate Prediction Center; 2019 p. 25. Available: https://www.cpc.ncep.noaa.gov/products/outreach/Report-to-the-Nation-Monsoon_aug04.pdf
- 121. Hansen H. Skylarking Cassin's Sparrows in Southeast Arizona. In: ABA Blog [Internet]. 2019 [cited 23 Jun 2021]. Available: <u>https://blog.aba.org/2019/08/skylarking-cassins-sparrows-in-southeast-arizona.</u> html
- 122. West AM, Kumar S, Brown CS, Stohlgren TJ, Bromberg J. Field validation of an invasive species Maxent model. Ecological Informatics. 2016; 36: 126–134. https://doi.org/10.1016/j.ecoinf.2016.11.001
- 123. Fourcade Y, Besnard AG, Secondi J. Paintings predict the distribution of species, or the challenge of selecting environmental predictors and evaluation statistics. Global Ecol Biogeogr. 2018; 27: 245– 256. https://doi.org/10.1111/geb.12684
- 124. Porter WP, Budaraju S, Stewart WE, Ramankutty N. Calculating Climate Effects on Birds and Mammals: Impacts on Biodiversity, Conservation, Population Parameters, and Global Community Structure. American Zoologist. 2000; 40: 597–630. https://doi.org/10.1668/0003-1569(2000)040[0597: cceoba]2.0.co;2
- **125.** Anderson JT, Conway WC. The flight song display of the Cassin's Sparrow (Aimophila cassinii): form and possible function. Bulletin of the Texas Ornithological Society. 2000; 33: 1–12.
- 126. Cuddington K, Fortin M-J, Gerber LR, Hastings A, Liebhold A, O'Connor M, et al. Process-based models are required to manage ecological systems in a changing world. Ecosphere. 2013; 4: 1–12. https://doi.org/10.1890/ES12-00178.1
- **127.** Kendall BE, Briggs CJ, Murdoch WW, Turchin P, Ellner SP, McCauley E, et al. Why do populations cycle? A synthesis of statistical and mechanistic modeling approaches. 1999; 80: 17.
- 128. Lipschutz ML. Effects of Drought and Grazzing on Land Bird Populations in South Texas. MS, Range and Wildlife Management, Texas A&M University-Kingsville. 2016.
- 129. Cassin's Sparrow—Whatbird.com. [cited 22 May 2021]. Available: https://identify.whatbird.com/obj/ 278/overview/cassins_sparrow.aspx
- Cassin's Sparrow. In: Audubon [Internet]. 2014 [cited 22 May 2021]. Available: <u>https://www.audubon.org/field-guide/bird/cassins-sparrow</u>
- 131. Cassin's Sparrow (Peucaea cassinii)—BirdLife species factsheet. [cited 22 May 2021]. Available: http://datazone.birdlife.org/species/factsheet/22721272
- Cassin's Sparrow Life History, All About Birds, Cornell Lab of Ornithology. [cited 22 May 2021]. Available: https://www.allaboutbirds.org/guide/Cassins_Sparrow/lifehistory
- 133. Tomašových A, Kidwell SM. The Effects of Temporal Resolution on Species Turnover and on Testing Metacommunity Models. The American Naturalist. 2010; 175: 587–606. <u>https://doi.org/10.1086/</u> 651661 PMID: 20302427
- 134. Dornelas M, Gotelli NJ, McGill B, Shimadzu H, Moyes F, Sievers C, et al. Assemblage time series reveal biodiversity change but not systematic loss. Science. 2014; 344: 296–299. <u>https://doi.org/10. 1126/science.1248484 PMID: 24744374</u>
- 135. Castillo-Escrivà A, Mesquita-Joanes F, Rueda J. Effects of the Temporal Scale of Observation on the Analysis of Aquatic Invertebrate Metacommunities. Front Ecol Evol. 2020; 8: 561838. https://doi.org/ 10.3389/fevo.2020.561838
- 136. Wisz MS, Broennimann O, Grønkjær P, Møller PR, Olsen SM, Swingedouw D, et al. Arctic warming will promote Atlantic–Pacific fish interchange. Nature Clim Change. 2015; 5: 261–265. https://doi.org/ 10.1038/nclimate2500
- 137. Sauer JR, Link WA, Fallon JE, Pardieck KL, Ziolkowski DJ. The North American Breeding Bird Survey 1966–2011: Summary Analysis and Species Accounts. North American Fauna. 2013; 79: 1–32. https://doi.org/10.3996/nafa.79.0001
- 138. North American Breeding Bird Survey. 2017. Available: https://www.pwrc.usgs.gov/bbs/.
- 139. eBird T. Global Big Day—8 May 2021—eBird. 2021 [cited 23 May 2021]. Available: https://ebird.org/ ebird/news/global-big-day-8-may-2021
- 140. Cornell Lab's Citizen Science Projects. In: Citizen Science [Internet]. [cited 23 Aug 2021]. Available: https://www.birds.cornell.edu/citizenscience/about-the-projects/
- Christmas Bird Count. In: Audubon [Internet]. [cited 23 May 2021]. Available: <u>https://www.audubon.org/conservation/science/christmas-bird-count</u>
- 142. About the Great Backyard Bird Count. In: Audubon [Internet]. 2015 [cited 23 May 2021]. Available: https://www.audubon.org/conservation/about-great-backyard-bird-count
- GBIF—Global Biodiversity Information Facility. 2020 [cited 22 May 2020]. Available: <u>https://www.gbif.org/</u>
- 144. VertNet. [cited 23 Aug 2021]. Available: http://vertnet.org/

- 145. BISON—Biodiversity Information Serving Our Nation. [cited 23 Aug 2021]. Available: https://bison.usgs.gov/#home
- 146. Zhang C, Chen Y, Xu B, Xue Y, Ren Y. Improving prediction of rare species' distribution from community data. Sci Rep. 2020; 10: 12230. https://doi.org/10.1038/s41598-020-69157-x PMID: 32699354
- 147. Lomba A, Pellissier L, Randin C, Vicente J, Moreira F, Honrado J, et al. Overcoming the rare species modelling paradox: A novel hierarchical framework applied to an Iberian endemic plant. Biological Conservation. 2010; 143: 2647–2657. https://doi.org/10.1016/j.biocon.2010.07.007
- 148. Galante PJ, Alade B, Muscarella R, Jansa SA, Goodman SM, Anderson RP. The challenge of modeling niches and distributions for data-poor species: a comprehensive approach to model complexity. Ecography. 2018; 41: 726–736.
- 149. Wisz MS, Hijmans RJ, Li J, Peterson AT, Graham CH, Guisan A, et al. Effects of sample size on the performance of species distribution models. Diversity and distributions. 2008; 14: 763–773.
- 150. Breiner FT, Nobis MP, Bergamini A, Guisan A. Optimizing ensembles of small models for predicting the distribution of species with few occurrences. Isaac N, editor. Methods Ecol Evol. 2018; 9: 802– 808. https://doi.org/10.1111/2041-210X.12957
- 151. Breiner FT, Guisan A, Bergamini A, Nobis MP. Overcoming limitations of modelling rare species by using ensembles of small models. Anderson B, editor. Methods Ecol Evol. 2015; 6: 1210–1218. https://doi.org/10.1111/2041-210X.12403
- 152. Heikkinen RK, Marmion M, Luoto M. Does the interpolation accuracy of species distribution models come at the expense of transferability? Ecography. 2012; 35: 276–288. <u>https://doi.org/10.1111/j.1600-0587.2011.06999.x</u>
- 153. Petitpierre B, Broennimann O, Kueffer C, Daehler C, Guisan A. Selecting predictors to maximize the transferability of species distribution models: lessons from cross-continental plant invasions: Which predictors increase the transferability of SDMs? Global Ecology and Biogeography. 2017; 26: 275–287. https://doi.org/10.1111/geb.12530
- **154.** Engler J, Rödder D. Disentangling interpolation and extrapolation uncertainties in ecologial niche models: a novel visualization technique for the spatial variation of predictor variable colinearity. Biodiversity Informatics. 2012; 8: 30–40.
- 155. IPCC Data Distribution Centre (DDC). 2021 [cited 26 Oct 2021]. Available: http://www.ipcc-data.org/
- 156. IPCC Global Climate Projections. [cited 26 Oct 2021]. Available: <u>https://www.ipcc.ch/report/ar4/wg1/</u>global-climate-projections/
- 157. Earth System Grid Federation (ESGF). 2021 [cited 8 Nov 2021]. Available: https://esgf.llnl.gov/index. html
- 158. Copernicus Climate Data Store (CDS). 2021 [cited 8 Nov 2021]. Available: https://cds.climate. copernicus.eu/#I/home
- 159. Climate Model Projections. In: Data.gov [Internet]. [cited 8 Nov 2021]. Available: https://www.data.gov/climate/portals/
- Vega G C., Pertierra LR, Olalla-Tárraga MÁ. MERRAclim, a high-resolution global dataset of remotely sensed bioclimatic variables for ecological modelling. Scientific Data. 2017; 4. https://doi.org/10.1038/ sdata.2017.78 PMID: 28632236
- 161. von Storch H, Feser F, Geyer B, Klehmet K, Li D, Rockel B, et al. Regional reanalysis without local data: Exploiting the downscaling paradigm: Regional Reanalysis by Downscaling. J Geophys Res Atmos. 2017; 122: 8631–8649. https://doi.org/10.1002/2016JD026332
- 162. Sen Gupta A, Tarboton DG. A tool for downscaling weather data from large-grid reanalysis products to finer spatial scales for distributed hydrological applications. Environmental Modelling & Software. 2016; 84: 50–69. https://doi.org/10.1016/j.envsoft.2016.06.014
- 163. Austin MP, Van Niel KP. Improving species distribution models for climate change studies: variable selection and scale: Species distribution models for climate change studies. Journal of Biogeography. 2011; 38: 1–8. https://doi.org/10.1111/j.1365-2699.2010.02416.x
- 164. SEDAC—Socioeconomic Data and Applications Center. [cited 23 Aug 2021]. Available: https://sedac. ciesin.columbia.edu/theme/remote-sensing/data/sets/browse
- 165. Earthdata—NASA's Earth Science Data Systems (ESDS) Program. [cited 23 Aug 2021]. Available: https://earthdata.nasa.gov//
- 166. NatureServe. [cited 23 Aug 2021]. Available: https://www.natureserve.org/
- 167. Peterson AT, Nakazawa Y. Environmental data sets matter in ecological niche modelling: an example with Solenopsis invicta and Solenopsis richteri. Global Ecology and Biogeography. 2008;0: 071113201427001-??? https://doi.org/10.1111/j.1466-8238.2007.00347.x

- 168. Barbet-Massin M, Jetz W. A 40-year, continent-wide, multispecies assessment of relevant climate predictors for species distribution modelling. Heikkinen R, editor. Diversity and Distributions. 2014; 20: 1285–1295. https://doi.org/10.1111/ddi.12229
- 169. ABoVE—Arctic-Boreal Vulnerability Experiment. 2021 [cited 27 May 2021]. Available: https://above.nasa.gov/index.html?
- 170. Carroll ML, Loboda TV. The sign, magnitude and potential drivers of change in surface water extent in Canadian tundra. Environ Res Lett. 2018; 13: 045009. https://doi.org/10.1088/1748-9326/aab794
- 171. Carroll ML, Schnase JL, Gill RL, Tamkin GS, Li J, Maxwell TP, et al. MERRA/Max: A Machine Learning Approach to Stochastic Convergence with a Multi-Variate Dataset. IGARSS 2020 Virtual Symposium. IEEE; 2020. Available: https://igarss2020.org/view_paper.php?PaperNum=4013