

Bézier interpolation improves the inference of dynamical models from data

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Many dynamical systems, from quantum many-body systems to evolving populations to financial markets, are described by stochastic processes. Parameters characterizing such processes can often be inferred using information integrated over stochastic paths. However, estimating time-integrated quantities from real data with limited time resolution is challenging. Here, we propose a framework for accurately estimating time-integrated quantities using Bézier interpolation. We applied our approach to two dynamical inference problems: Determining fitness parameters for evolving populations and inferring forces driving Ornstein-Uhlenbeck processes. We found that Bézier interpolation reduces the estimation bias for both dynamical inference problems. This improvement was especially noticeable for data sets with limited time resolution. Our method could be broadly applied to improve accuracy for other dynamical inference problems using finitely sampled data.

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I. INTRODUCTION

Stochastic processes are ubiquitous in nature. In biology, the evolution of genetic sequences can be formulated as a stochastic process. The Wright-Fisher (WF) model [1], a discrete-time stochastic process, has been used to study the evolution of organisms from viruses [2–4] to humans [5]. Models such as the Ornstein-Uhlenbeck (OU) process [6,7] have been applied to describe a wide range of phenomena, from the fluctuation of currency exchange rates [8] and cell migration [9] to driven quantum many-body systems [10].

Appropriate model parameters are needed to accurately describe the behavior of real systems. To infer such parameters from data, it is often necessary to compute statistics over a *path*, i.e., a complete realization of the stochastic processes. For example, the restoring force of the OU process can be estimated by taking the ratio of the deviation from the equilibrium position and the magnitude of the intrinsic fluctuations, both integrated over a stochastic path [11,12].

However, real data often consists of incomplete, occasional measurements of a system, which may also be limited by experimental constraints. This makes it more difficult to accurately estimate model parameters since statistics over the path must be estimated from incomplete information.

Here, we propose a tractable nonlinear interpolation framework using Bézier curves [13–16]. In addition to incorporating nonlinearity, this approach has the added advantage of conserving sums of categorical variables. This property can be especially useful for conserved quantities such as probabilities.

We applied Bézier interpolation to two example problems: inferring natural selection in evolving populations through the WF model and inferring restoring forces for OU processes.

Our method reduces estimation bias and improves the precision of model inferences. Furthermore, we show that the autocorrelation function of statistics over a path identifies time scales over which nonlinear interpolation is particularly effective, which is consistent with our observations in simulations. We show that Bézier interpolation can generically improve solutions of dynamical inference problems by accurately estimating statistics over stochastic paths. We expect that this nonlinear interpolation method can improve a wide range of dynamical inference problems beyond the specific examples we consider, such as parameter estimation for stochastic differential equations.

II. RESULTS

A. Bézier interpolation

Consider a function $x(t)$ sampled at discrete times t_k for $k \in \{0, 1, \dots, K\}$. Then the interpolated value of the function $x_B^{(k)}(t)$ between two successive discrete time points t_k and t_{k+1} is given by

$$x_B^{(k)}(t) = \sum_{n=0}^P \beta_n \left(\frac{t - t_k}{t_{k+1} - t_k} \right) \phi_n^{(k)}[(x(t_{k'}))_{k'=0}^K]. \quad (1)$$

Here, β_n is the n th Bernstein basis polynomial of degree P , with $\beta_n(\tau) = \binom{P}{n} \tau^n (1-\tau)^{P-n} \geq 0$. The control points $\phi_n^{(k)}[(x(t_{k'}))_{k'=0}^K]$ depend on the ensemble of data points $(x(t_k))_{k=0}^K$ and determine the outline of the interpolation curves.

For simplicity, we consider cubic ($P = 3$) interpolation, but our approach can be extended to polynomials of different degrees P . We impose the following conditions to ensure that the segment at each interval $[t_k, t_{k+1}] \forall k$ is seamlessly connected,

$$\phi_0^{(k)}[(x(t_{k'}))_{k'=0}^K] = x(t_k), \quad \phi_3^{(k)}[(x(t_{k'}))_{k'=0}^K] = x(t_{k+1}).$$

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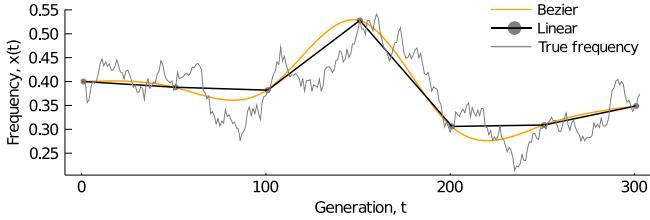


FIG. 1. Bézier interpolation generates smooth curves. Cubic Bézier curves smoothly interpolate between discretely sampled frequency trajectories generated from a Wright-Fisher model. *Simulation parameters.* $L = 50$ sites, population size $N = 10^3$, mutation rate $\mu = 10^{-3}$, with simulations over $T = 300$ generations. Data points are sampled every 50 generations and interpolated using cubic Bézier and linear interpolation.

Other internal points $\{(\phi_1^{(k)}, \phi_2^{(k)})\}_{k=0}^{K-1}$ are obtained by solving an optimization problem that reflects continuity and smoothness constraints imposed on the curves (see Supplemental Material [17]) (Fig. 1).

B. Wright-Fisher model of evolution

The WF model [1] is a classical model in evolutionary biology. In this model, a population of N individuals evolves over discrete generations under the influence of random mutations and natural selection. Each individual is represented by a genetic sequence of length L . For simplicity, we assume that each site in the genetic sequence is occupied by a mutant (1) or wild-type (0) nucleotide. There are thus $M = 2^L$ possible *genotypes* (i.e., genetic sequences) in the population.

The state of the population is described by a genotype frequency vector $\mathbf{z}(t) = (z_a(t))_{a=1}^M$, where $z_a(t)$ represents the frequency of individuals with genotype a in the population at time t . In the WF model, the probability of obtaining a genotype frequency vector \mathbf{z}' in the next generation is multinomial, with the succession probability of genotype a

$$p_a(\mathbf{z}(t)) \propto f_a z_a(t) + \sum_{b|b \neq a} [\mu_{ba} z_b(t) f_b - \mu_{ab} z_a(t) f_a]. \quad (2)$$

In (2), f_a denotes the *fitness* of genotype a . Individuals with higher fitness values reproduce more readily than those with lower fitness values. Here μ_{ab} is the probability to mutate from genotype a to genotype b .

Fitness values can be estimated from data by identifying the f_a that are most likely to generate the observed evolutionary history of a population, but this is challenging due to the enormous size of the genotype space. The problem can be simplified by assuming that fitness values are additive, $f_a = 1 + \sum_{i=1}^L \sigma_i^a s_i$, where $\sigma_i^a = 1$ if the nucleotide at site i in genotype a is a mutant and 0 otherwise. The s_i are referred to as *selection coefficients*, which are positive if the mutation at site i is beneficial for reproduction and negative if mutation at site i is deleterious. Similarly, the mutation rate μ_{ab} can be simplified to a constant μ if genotypes a and b differ from one another by only a single mutation and zero otherwise.

Sohail *et al.* solved this problem analytically in the limit that the population size $N \rightarrow \infty$, while the selection coefficients s_i and mutation rate μ scale as $1/N$ (Ref. [4]). In this

case, the maximum *a posteriori* vector of selection coefficients $\hat{\mathbf{s}} = (\hat{s}_i)_{i=1}^L$ is

$$\hat{\mathbf{s}} = \left(\int_{t_0}^{t_K} dt \mathbf{C}(t) + \gamma \mathbf{I} \right)^{-1} \times \left[\mathbf{x}(t_K) - \mathbf{x}(t_0) - \mu \int_{t_0}^{t_K} dt [1 - 2\mathbf{x}(t)] \right], \quad (3)$$

where the time of observation runs from t_0 to t_K . In (3), $\mathbf{x}(t) = (x_i(t))_{i=1}^L$ is a vector of mutant frequencies (i.e., the number of individuals in the population with a mutation at site i at time t) and $\mathbf{C}(t)$ is the covariance matrix of mutant frequencies at time t . Here γ is the precision of a Gaussian prior distribution for the selection coefficients with mean zero and \mathbf{I} is the identity matrix.

Extensive past work has also considered numerical solutions to this problem [3,5,18–22], though the analytical formula in (3) typically outperforms numerical approaches [4]. Sohail *et al.* referred to (3) as the marginal path likelihood (MPL) estimate for the selection coefficients, obtained by maximizing the posterior probability of an evolutionary history with respect to the selection coefficients. The MPL approach has also been extended to consider more complex evolutionary models [23], missing covariance data [24], and epidemiological dynamics [25].

C. Bézier interpolation for WF model inference

In practice, Eq. (3) is not straightforward to evaluate because sequence data comes at discrete times $(t_k)_{k=0}^K$. However, Bézier interpolation allows us to analytically integrate both mutant frequency trajectories $\mathbf{x}(t)$ and covariances $\mathbf{C}(t)$, obtained by interpolating frequencies and computing $C_{ij}(t) = x_{ij}(t) - x_i(t)x_j(t)$. Here $x_{ij}(t)$ is the frequency of individuals in the population at time t that have mutations at both sites i and j .

To assess the performance of Bézier interpolation for inferring selection in the WF model, we generated a test data set by running 100 replicate simulations of WF evolution with identical parameters [Fig. 2(a)]. We then inferred selection coefficients from this data using MPL with linear and Bézier interpolation, applied to data sampled at discrete intervals $\Delta t = 75$ generations apart. While MPL with linear interpolation readily distinguishes between beneficial, neutral, and deleterious parameters, the inferred selection coefficients are shrunk towards zero. However, parameters inferred using Bézier interpolation are distributed around their true values [Fig. 2(b)]. Bézier interpolation reduces estimation bias due to long intervals between observation intervals by producing better estimates of underlying covariances (which we will quantify below). Here we used a regularization strength of $\gamma = 0.1$, but similar results are obtained with different choices for the regularization (see Supplemental Material [17]).

Next we studied how Bézier interpolation affects our ability to classify mutations as beneficial or deleterious, which we evaluated by ranking mutations according to their inferred selection coefficients. We quantified classification accuracy using positive predictive value (PPV), $PPV = TP/(TP + FP)$, where TP and FP are the numbers of true positive and false positive predictions.

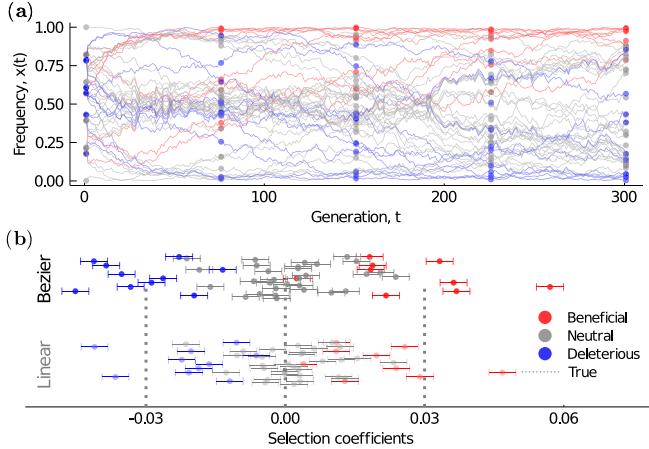


FIG. 2. Bézier interpolation reduces bias in estimated selection coefficients. (a) Wright-Fisher simulation with selection and mutation. Each trajectory drawn as a solid line is true complete data, and filled circles are a subset of the complete data, which is observed every $\Delta t = 75$ generations and used for selection coefficient prediction. (b) Selection coefficients for the frequency trajectories in (a) were estimated by MPL with Bézier and linear interpolation. MPL with Bézier interpolation greatly reduces estimation bias for inferred selection coefficients when the time interval between sampled observations is large. *Simulation parameters.* $L = 50$ sites with 10 beneficial, 10 deleterious, and 30 neutral mutations with selection coefficients of $s = 0.03$, $s = -0.03$, and $s = 0$, respectively. Other parameters of the WF models are the same as in Fig. 1.

The PPV curves for beneficial or deleterious mutations estimated by MPL with Bézier interpolation are higher than those with linear interpolation, indicating more accurate classification [Figs. 3(a) and 3(b)]. This can be understood by observing reduced overlap between the distribution of inferred

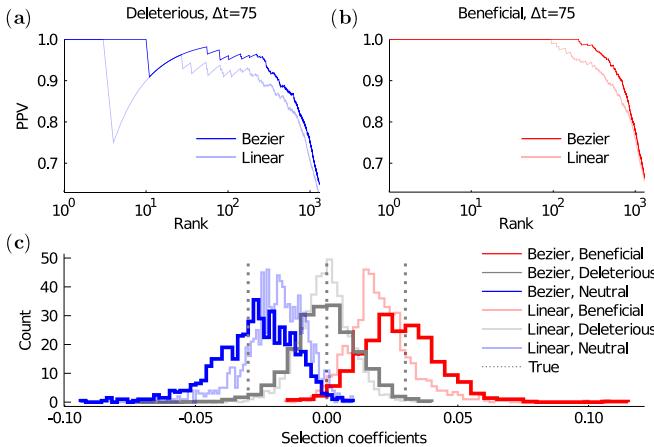


FIG. 3. MPL with Bézier interpolation improves prediction precision and reduces estimation bias. (a),(b) When the observation time interval is longer ($\Delta t = 75$), the PPV curve for Bézier interpolation is universally higher than the curve for linear interpolation for both deleterious and beneficial cases. (c) The selection coefficient distributions estimated by MPL with linear interpolation visibly shrank toward zero and were biased, while distributions estimated by MPL with Bézier interpolation did not considerably shrink and have the mean values near the true selection values.

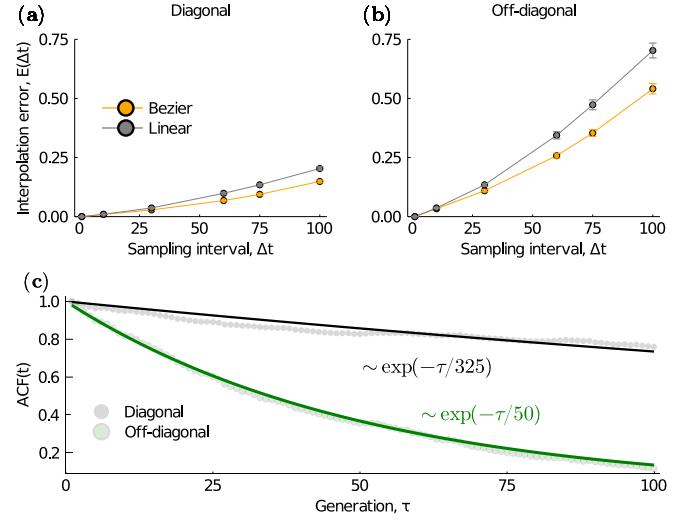


FIG. 4. Bézier method suppresses interpolation error, especially for off-diagonal pairwise covariances. (a) Sampling time interval dependence for interpolation errors $\mathcal{E}(\Delta t)$ for diagonal covariances and (b) for off-diagonal pairwise covariances. We simulated WF dynamics using the model described in Fig. 2 and generated data sets that evolved to the 300th generation for each trial. For example, when $\Delta t = 100$, results only use data from $t = 0, 100, 200$, and 300. (c) The autocorrelation of off-diagonal covariance elements decays faster than diagonal ones. To simplify the analysis, we evaluated the autocorrelation function from generation $t = 50$. The diagonal autocorrelation shows nonmonotonic decay after long times due to mutant frequencies that approach the frequency boundaries (i.e., 0 and 1).

selection coefficients for beneficial, neutral, and deleterious mutations using Bézier interpolation [Fig. 3(c)].

D. Recovery of rapidly decaying correlations underlies improved accuracy

To understand why MPL with Bézier interpolation yields more accurate inferences, we studied errors between true and estimated parameters as a function of the time interval Δt between samples. For arbitrary matrices M we define an error function $\mathcal{E}(\Delta t) = \|\mathcal{I}_M(\Delta t) - \mathcal{I}_M(1)\|/\|\mathcal{I}_M(1)\|$, normalizing by the matrix norm $\|\mathcal{I}_M(1)\|$, which corresponds to perfect sampling for the WF model. Here, $\mathcal{I}_M(\Delta t)$ is a time integral depending on the type of integration (piecewise constant, linear, and Bézier). For example, for piecewise constant integration, it will be $\mathcal{I}_M(\Delta t) = \sum_{k=1}^{\lfloor T/\Delta t \rfloor} M(k\Delta t)\Delta t$. In the discussion below we apply the L_2 norm, $\|M\| = \sqrt{(\sum_{i,j} M_{ij}^2)}$, but other conventions could also be considered.

Using the metric defined above, we found that Bézier interpolation yields better estimates for both the diagonal and off-diagonal terms of the mutant frequency covariance matrix. However, the error for the off-diagonal covariances is larger and increases much more rapidly with increasing Δt than the error for the diagonal variances [Figs. 4(a) and 4(b)]. The reduction in error for Bézier interpolation is more substantial for off-diagonal terms compared to diagonal ones. Consistent with this observation, Bézier interpolation yields smaller improvements in performance for a simple version of MPL

in which the off-diagonal terms of the integrated covariance matrix are ignored (see Supplemental Material [17]) [referred to as the single locus (SL) method in Ref. [4]].

To study the time scale τ on which nonlinear effects become important and Bézier interpolation is advantageous, we modeled the covariance elements using a simple Langevin equation, $\dot{z}(t) = -\lambda z(t) + \xi(t)$. Here $z(t)$ represents an element of the covariance matrix, $\lambda > 0$ is a damping coefficient, and $\xi(t)$ is a standard white noise with $\langle \xi(t) \rangle = 0$ and $\langle \xi(t) \xi(t + \tau) \rangle = 2\delta(\tau)$. Following this approach, a linear approximation should describe the evolution of $z(t)$ accurately if $\lambda \Delta t \ll 1$; otherwise, nonlinear nature of the $z(t)$ becomes significant and at this point the linear approximation cannot capture the actual evolution of $z(t)$.

The damping coefficient λ can be estimated by computing the autocorrelation function (ACF) of the covariance matrix elements, which can be matched to expectations from the Langevin equation, $\langle x(t)x(t + \tau) \rangle \propto \exp(-\lambda\tau)$. In our simulations, the exponents of the ACF for diagonal and off-diagonal terms are around $\lambda_d \sim 1/325$ and $\lambda_o \sim 1/50$, respectively [Fig. 4(c)]. When the time between sampling events is $\Delta t = 75$, where Bézier interpolation clearly has an advantage (Fig. 3), for diagonal and off-diagonal covariances we have $\lambda_d \Delta t \sim 0.23$ and $\lambda_o \Delta t = 1.5$, respectively. At this point, $\lambda_o \Delta t$ is $O(1)$, indicating the onset of nonlinearity for off-diagonal terms. Consistent with this observation, for this value of Δt , Bézier interpolation has notably lower error for off-diagonal covariances than linear interpolation, while errors for the diagonal terms are comparable.

While we focused specifically on the WF model in this example, the principle of autocorrelations and transitioning between linear and nonlinear behavior is general. This can allow us to anticipate the benefit of nonlinear interpolation for a wide range of problems.

E. Inference of forces in Ornstein-Uhlenbeck processes

We further applied Bézier interpolation to accurately infer the collective forces in Ornstein-Uhlenbeck (OU) processes, which plays important roles in various fields such as physics, biology, and mathematical finance [11,26–28]. Data has been used to infer the parameters of OU processes describing phenomena including cell migration [29], coevolution of species [30], and currency exchange rates [31], to name a few examples.

We consider the following multivariate OU process:

$$d\mathbf{X}_t = \mathbf{J}\mathbf{X}_t dt + \Sigma^{1/2} d\mathbf{W}_t. \quad (4)$$

Here t is the time variable, L is the number of OU variables, $\mathbf{X}_t \in \mathbb{R}^L$, $\mathbf{J} \in \mathbb{R}^{L \times L}$ is a negative semidefinite matrix, Σ is a time-independent noise covariance, and \mathbf{W}_t is a Wiener process. We assume that the noise covariance matrix is constant over the evolution and given. Therefore, the unknown variable in the SDE in (4) is only the drift term, the interaction matrix \mathbf{J} .

One of the most commonly used approaches for inferring stochastic force in OU processes is maximizing the likelihood ratio or Radon-Nikodym derivative, which is the ratio of two probability measures [12,32] and is commonly employed in fields such as mathematical finance [11]. In our problem, the likelihood ratio is defined as the probability density obeying

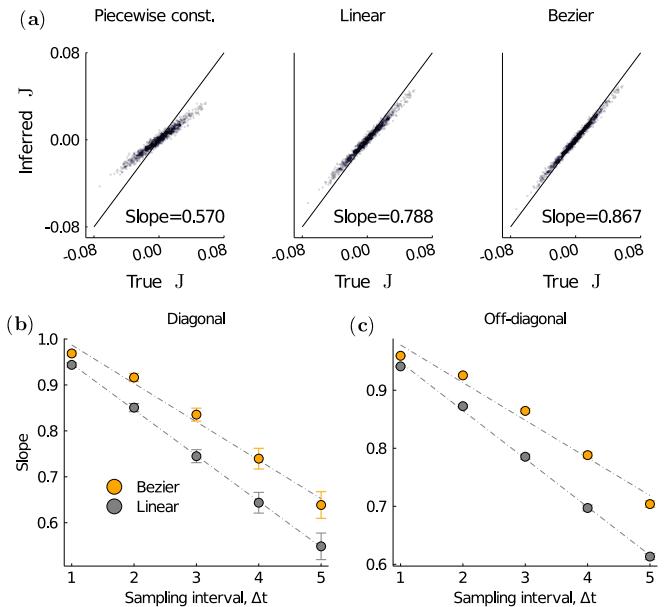


FIG. 5. Bézier interpolation can improve the inference accuracy of parameters in the OU process. (a) Comparison between true and inferred OU parameters using piecewise constant, linear, and Bézier interpolation. Linear regression slope values are included in each panel. Inferred interaction parameters using Bézier interpolation correspond most closely with the true parameters. (b) Dependence of the slope between true and inferred parameters on the time sampling interval $\Delta t = 1$, shown separately for the (b) diagonal and (c) off-diagonal interaction parameters of the J matrix. In both diagonal and off diagonal, the slope values decrease more gradually with increasing Δt for Bézier interpolation than for linear interpolation.

the dynamics of (4) with interactions divided by the probability density of a “null” model with no interactions. Here, we inferred OU interactions by directly maximizing the path likelihood, as described for the WF model. Interestingly, this leads to exactly the same solution as the one for the standard likelihood or Radon-Nikodym derivative methods (see Supplemental Material [17]).

The interaction matrix $\hat{\mathbf{J}}$ that best describes the data is given by

$$\hat{\mathbf{J}} = \left(\sum_{k=0}^{K-1} \Delta \mathbf{x}(t_k) \mathbf{x}(t_k)^\top \right) \left(\sum_{k=0}^{K-1} \Delta t_k \mathbf{x}(t_k) \mathbf{x}(t_k)^\top \right)^{-1}. \quad (5)$$

Here $(\mathbf{x}(t_k))_{k=0}^{K-1}$ is the observed trajectory following the OU process, $\Delta t_k = t_{k+1} - t_k$ is an observation interval, and $\Delta \mathbf{x}(t_k) = \mathbf{x}(t_{k+1}) - \mathbf{x}(t_k)$ is the amount of change during the k th interval.

To generate test data, we simulated the OU process using negative definite interaction matrices (see Supplemental Material [17]), which follows the construction of a Hopfield network [33]. Hopfield networks were first constructed to study associative memory [33] and have since been applied to inference problems in biology [34–37].

Interaction parameters estimated using Bézier interpolation matched better with the true, underlying parameters than those inferred using linear interpolation or a piecewise-constant assumption for the $\mathbf{x}(t)$ [Fig. 5(a)]. In particular, large

parameters inferred with linear interpolation or the piecewise-constant assumption tended to be underestimated. In addition, we found that the slope relating the true and inferred parameters decreases as the sampling interval Δt increases. However, the slope decreases more slowly for Bézier interpolation compared to linear interpolation [Figs. 5(b) and 5(c)]. Overall, OU interaction parameters inferred using Bézier interpolation more closely match the true, underlying parameters than those inferred with simpler interpolation approaches, with gains in performance that increase as data becomes more limited.

III. DISCUSSION

Here we developed a nonlinear interpolation method using Bézier curves that improves the inference of dynamical models from finite data. We applied our approach to two problems: The inference of natural selection in evolving populations and interactions in multivariate Ornstein-Uhlenbeck processes. Bézier interpolation makes inference more pre-

cise and reduces bias, especially for data sets that are more sparsely sampled.

Because of its generality, Bézier interpolation could be broadly applied to give more reliable results for dynamic inference problems. For example, our approach could be combined with methods to learn forces from nonequilibrium dynamics [38,39] or ones used to learn parameters of stochastic differential equations from finitely sampled data [6,11,40].

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All authors contributed to methods development, data analysis, interpretation of results, and writing the paper. K.S. performed simulations and computational analyses. J.P.B. supervised the project.

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- [1] W. J. Ewens, *Mathematical Population Genetics: Theoretical Introduction* (Springer, Berlin, 2004), Vol. 1.
 - [2] M. Foll, Y.-P. Poh, N. Renzette, A. Ferrer-Admetlla, C. Bank, H. Shim, A.-S. Malaspinas, G. Ewing, P. Liu, D. Wegmann *et al.*, Influenza virus drug resistance: A time-sampled population genetics perspective, *PLoS Genet.* **10**, e1004185 (2014).
 - [3] A. Ferrer-Admetlla, C. Leuenberger, J. D. Jensen, and D. Wegmann, An approximate Markov model for the Wright–Fisher diffusion and its application to time series data, *Genetics* **203**, 831 (2016).
 - [4] M. S. Sohail, R. H. Y. Louie, M. R. McKay, and J. P. Barton, Mpl resolves genetic linkage in fitness inference from complex evolutionary histories, *Nat. Biotechnol.* **39**, 472 (2021).
 - [5] I. Mathieson and G. McVean, Estimating selection coefficients in spatially structured populations from time series data of allele frequencies, *Genetics* **193**, 973 (2013).
 - [6] S. M. Iacus, *Simulation and Inference for Stochastic Differential Equations: With R Examples* (Springer, Berlin, 2008), Vol. 486.
 - [7] D. T. Gillespie, Exact numerical simulation of the Ornstein–Uhlenbeck process and its integral, *Phys. Rev. E* **54**, 2084 (1996).
 - [8] G. O. Roberts, O. Papaspiliopoulos, and P. Dellaportas, Bayesian inference for non-gaussian ornstein–uhlenbeck stochastic volatility processes, *J. R. Stat. Soc.: Ser. B* **66**, 369 (2004).
 - [9] P. Dieterich, R. Klages, R. Preuss, and A. Schwab, Anomalous dynamics of cell migration, *Proc. Natl. Acad. Sci. USA* **105**, 459 (2008).
 - [10] P. Jung, Periodically driven stochastic systems, *Phys. Rep.* **234**, 175 (1993).
 - [11] P. C. B. Phillips and J. Yu, Maximum likelihood and gaussian estimation of continuous time models in finance, in *Handbook of Financial Time Series* (Springer, Berlin, 2009), pp. 497–530.
 - [12] R. S. Liptser and A. N. Shiriaev, *Statistics of Random Processes: General Theory* (Springer, Berlin, 1977), Vol. 394.
 - [13] R. T. Farouki, The Bernstein polynomial basis: A centennial retrospective, *Computer Aided Geometric Design* **29**, 379 (2012).
 - [14] J.-W. Choi, R. Curry, and G. Elkaim, Path planning based on bézier curve for autonomous ground vehicles, in *Advances in Electrical and Electronics Engineering-IAENG Special Edition of the World Congress on Engineering and Computer Science 2008* (IEEE, New York, 2008), pp. 158–166.
 - [15] K. R. Simba, N. Uchiyama, and S. Sano, Real-time obstacle-avoidance motion planning for autonomous mobile robots, in *Proceedings of the 2014 4th Australian Control Conference (AUCC)* (IEEE, New York, 2014), pp. 267–272.
 - [16] A. R. Forrest, Interactive interpolation and approximation by bézier polynomials, *Comput. J.* **15**, 71 (1972).
 - [17] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevE.107.024116> for details about (1) the data and code for the reported results, (2) the definition of Bézier interpolation, (3) analytical expressions of the integrated covariance matrix, (4) analysis of the normalizability and non-negativity of Bézier interpolation, (5) details of Ornstein–Uhlenbeck (OU) simulations, (6) maximum path-likelihood estimation for OU processes, (7) application of the Cameron–Martin–Girsanov theorem for OU inference, (8) application of Bézier interpolation for hiv-1 data, (9) the effects of regularization strength and sampling intervals, (10) numerical tests of the positive-semidefiniteness of interpolated covariance matrices, and (11) complementary simulations for inference of OU processes. It also includes Refs. [41–51].
 - [18] M. Lacerda and C. Seoighe, Population genetics inference for longitudinally-sampled mutants under strong selection, *Genetics* **198**, 1237 (2014).
 - [19] P. Tataru, M. Simonsen, T. Bataillon, and A. Hobolth, Statistical inference in the wright–fisher model using allele frequency data, *Syst. Biol.* **66**, e30 (2017).
 - [20] J. G. Schraiber, S. N. Evans, and M. Slatkin, Bayesian inference of natural selection from allele frequency time series, *Genetics* **203**, 493 (2016).

- [21] M. Foll, H. Shim, and J. D. Jensen, WFABC: A Wright-Fisher ABC-based approach for inferring effective population sizes and selection coefficients from time-sampled data, *Mol. Ecol. Resources* **15**, 87 (2015).
- [22] A. Iranmehr, A. Akbari, C. Schlötterer, and V. Bafna, Clear: Composition of likelihoods for evolve and resequence experiments, *Genetics* **206**, 1011 (2017).
- [23] M. S. Sohail, R. H. Y. Louie, Z. Hong, J. P. Barton, and M. R. McKay, Inferring epistasis from genetic time-series data, *Mol. Biol. Evol.* **39**, msac199 (2022).
- [24] Y. Li and J. P. Barton, *Estimating Linkage Disequilibrium and Selection from Allele Frequency Trajectories*, Genetics (2023).
- [25] B. Lee, M. S. Sohail, E. Finney, S. F. Ahmed, A. A. Quadeer, M. R. McKay, and J. P. Barton, Inferring effects of mutations on SARS-CoV-2 transmission from genomic surveillance data, medRxiv 2021, <https://doi.org/10.1101/2021.12.31.21268591>.
- [26] J.-P. Bouchaud and R. Cont, A Langevin approach to stock market fluctuations and crashes, *Eur. Phys. J. B* **6**, 543 (1998).
- [27] O. Vasicek, An equilibrium characterization of the term structure, *J. Financ. Econ.* **5**, 177 (1977).
- [28] R. S. Mamon, Three ways to solve for bond prices in the vasicek model, *Adv. Decision Sci.* **8**, 1 (2004).
- [29] D. B. Brückner, A. Fink, C. Schreiber, P. J. F. Röttgermann, J. O. Rädler, and C. P. Broedersz, Stochastic nonlinear dynamics of confined cell migration in two-state systems, *Nat. Phys.* **15**, 595 (2019).
- [30] L. S. T. Ho and C. Ané, Intrinsic inference difficulties for trait evolution with ornstein-uhlenbeck models, *Methods Ecol. Evol.* **5**, 1133 (2014).
- [31] G. O. Roberts and J. S. Rosenthal, Optimal scaling of discrete approximations to langevin diffusions, *J. R. Stat. Soc.: Ser. B* **60**, 255 (1998).
- [32] H. Risken, *The Fokker-Planck Equation: Methods of Solution and Applications*, 2nd ed. (Springer-Verlag, Berlin, 1989).
- [33] J. J. Hopfield, Neural networks and physical systems with emergent collective computational abilities, *Proc. Natl. Acad. Sci. USA* **79**, 2554 (1982).
- [34] S. Cocco, R. Monasson, and M. Weigt, From principal component to direct coupling analysis of coevolution in proteins: Low-eigenvalue modes are needed for structure prediction, *PLoS Comput. Biol.* **9**, e1003176 (2013).
- [35] J. Tubiana, S. Cocco, and R. Monasson, Learning protein constitutive motifs from sequence data, *eLife* **8**, e39397 (2019).
- [36] K. Shimagaki and M. Weigt, Selection of sequence motifs and generative Hopfield-Potts models for protein families, *Phys. Rev. E* **100**, 032128 (2019).
- [37] K. Shimagaki and M. Weigt, Selection of sequence motifs and generative Hopfield-Potts models for protein families, *bioRxiv* 652784 (2019), <https://doi.org/10.1101/652784>.
- [38] A. Frishman and P. Ronceray, Learning Force Fields from Stochastic Trajectories, *Phys. Rev. X* **10**, 021009 (2020).
- [39] D. B. Brückner, P. Ronceray, and C. P. Broedersz, Inferring the Dynamics of Underdamped Stochastic Systems, *Phys. Rev. Lett.* **125**, 058103 (2020).
- [40] F. Ferretti, V. Chardès, T. Mora, A. M. Walczak, and I. Giardina, Building General Langevin Models from Discrete Datasets, *Phys. Rev. X* **10**, 031018 (2020).
- [41] E. H. Doha, A. H. Bhrawy, and M. A. Saker, Integrals of bernstein polynomials: An application for the solution of high even-order differential equations, *Appl. Math. Lett.* **24**, 559 (2011).
- [42] A. Altürk, Application of the bernstein polynomials for solving volterra integral equations with convolution kernels, *Filomat* **30**, 1045 (2016).
- [43] R. H. Cameron and W. T. Martin, Transformations of weiner integrals under translations, *Ann. Math.* **45**, 386 (1944).
- [44] I. V. Girsanov, On transforming a certain class of stochastic processes by absolutely continuous substitution of measures, *Theory Probab. Appl.* **5**, 285 (1960).
- [45] M. K. P. Liu, N. Hawkins, A. J. Ritchie, V. V. Ganusov, V. Whale, S. Brackenridge, H. Li, J. W. Pavlicek, F. Cai, M. Rose-Abrahams *et al.*, Vertical T cell immunodominance and epitope entropy determine HIV-1 escape, *J. Clin. Invest.* **123**, 380 (2013).
- [46] F. Zanini, J. Brodin, L. Thebo, C. Lanz, G. Bratt, J. Albert, and R. A. Neher, Population genomics of intrapatient HIV-1 evolution, *eLife* **4**, e11282 (2015).
- [47] R. Christensen, *Advanced Linear Modeling* (Springer, New York, 2019).
- [48] D. J. C. MacKay, D. J. C. M. Kay *et al.*, *Information Theory, Inference and Learning Algorithms* (Cambridge University Press, Cambridge, UK, 2003).
- [49] C. M. Bishop and N. M. Nasrabadi, *Pattern Recognition and Machine Learning* (Springer, Berlin, 2006), Vol. 4.
- [50] Q. Lin and C. Li, Kriging based sequence interpolation and probability distribution correction for gaussian wind field data reconstruction, *J. Wind Eng. Ind. Aerodyn.* **205**, 104340 (2020).
- [51] F. Morcos, A. Pagnani, B. Lunt, A. Bertolino, D. S. Marks, C. Sander, R. Zecchina, J. N. Onuchic, T. Hwa, and M. Weigt, Direct-coupling analysis of residue coevolution captures native contacts across many protein families, *Proc. Natl. Acad. Sci. USA* **108**, E1293 (2011).