

Estimating Improved Partitioning Schemes for Ultraconserved Elements

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Abstract

Ultraconserved (UCEs) are popular markers for phylogenomic studies. They are relatively simple to collect from distantly-related organisms, and contain sufficient information to infer relationships at almost all taxonomic levels. Most studies of UCEs use partitioning to account for variation in rates and patterns of molecular evolution among sites, for example by estimating an independent model of molecular evolution for each UCE. However, rates and patterns of molecular evolution vary substantially within as well as between UCEs, suggesting that there may be opportunities to improve how UCEs are partitioned for phylogenetic inference. We propose and evaluate new partitioning methods for phylogenomic studies of UCEs: Sliding-Window Site Characteristics (SWSC), and UCE Site Position (UCESP). The first method uses site characteristics such as entropy, multinomial likelihood, and GC content to generate partitions that account for heterogeneity in rates and patterns of molecular evolution within each UCE. The second method groups together nucleotides that are found in similar physical locations within the UCEs. We examined the new methods with seven published data sets from a variety of taxa. We demonstrate the UCESP method generates partitions that are worse than other strategies used to partition UCE data sets (e.g., one partition per UCE). The SWSC method, particularly when based on site entropies, generates partitions that account for within-UCE heterogeneity and leads to large increases in the model fit. All of the methods, code, and data used in this study, are available from <https://github.com/Tagliacollo/PartitionUCE>. Simplified code for implementing the best method, the SWSC-EN, is available from <https://github.com/Tagliacollo/PFinderUCE-SWSC-EN>.

Key words: phylogenomics, UCEs, partitioning, partitioning methods, sliding-window site characteristics, PartitionFinder.

Introduction

Ultraconserved Elements (UCEs) are becoming one of the most popular DNA markers used for phylogenomic studies (Crawford et al. 2012; Faircloth et al. 2013, 2015; Jarvis et al. 2014; Meiklejohn et al. 2016). UCEs are highly conserved regions of DNA found throughout the genomes of many distant-related species. The functions of UCEs are not completely understood; however, their frequent proximity to transcriptional regulators or developmental genes has led to suggestions that they are directly involved in transcriptional regulation (Woolfe et al. 2005; Pennacchio et al. 2006). The conserved core regions of UCEs contain little variation, but the adjacent flanking regions contain more variation, and numerous studies have demonstrated that this variation is useful for inferring phylogenetic relationships between individuals, species, and higher clades (Crawford et al. 2012; Faircloth et al. 2012, 2015; Smith et al. 2014; Harrington et al. 2016; Moyle et al. 2016).

The accuracy of phylogenetic inferences often depends on choosing an appropriate model of molecular evolution, and many studies have demonstrated that accounting for variation in rates and patterns of evolution among sites is of

primary importance (Shapiro et al. 2006; Li et al. 2008; Ho and Lanfear 2010; Lanfear et al. 2012, 2014). There are multiple methods that account for variation of molecular evolution among sites (Le et al. 2008; Zhang and Townsend 2009; Goremykin et al. 2010; Cummins and McInerney 2011; Soubrier et al. 2012). The simplest and most widely-used in phylogenomics are the partitioning methods, which rely on defining groups of sites that have evolved under similar conditions. Specifically, partitioning methods assume a priori that each group of sites has evolved under the same Markov model of DNA sequence evolution, but that different groups may have evolved under different models. Partitioning has been shown to improve estimates of topologies, branch lengths, and divergence dates (Lanfear et al. 2014; Kainer and Lanfear 2015; Hoff et al. 2016).

Most phylogenomic studies of UCE markers use partitioning to account for variation in rates and patterns of evolution across sites. Typically, researchers choose one of two strategies: either they assign all UCEs in the alignment to a single partition (e.g., Faircloth et al. 2015), or they assign each UCE to a separate partition (e.g., Faircloth et al. 2013). The former strategy makes the assumption that every site in the

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alignment has evolved under a common Markov process, which is inadvisable and has been shown to increase the risk of inferring unreliable phylogenetic trees (Kainer and Lanfear 2015). The latter strategy accounts for variation in rates and patterns of evolution between UCEs, but makes the assumption that all of the sites within each UCE have evolved under the same Markov process. Given that the rates of molecular evolution within each UCE are known to vary predictably, with near-zero variation in the conserved core of the UCE increasing to large amounts of variation at the edges, it seems sensible to ask whether it is possible to improve upon the assumption that all sites within each UCE have evolved under a single Markov model.

Our focus in this paper is on asking whether it is possible to estimate partitioning schemes for UCE data sets which improve upon the assumption that all sites in each UCE have evolved under a single Markov model. One way to approach this would be to use a recently-proposed k-means clustering approach which iteratively divides alignments into groups comprised of nucleotide sites sharing similar substitution rates (Frandsen et al. 2015). However, it has recently come to light that this approach systematically generates a partition comprised of all the invariant sites in the data set, which can subsequently mislead phylogenetic inference methods (Baca et al. 2017). We sought instead to develop approaches that avoid this problem, leverage the known molecular evolutionary patterns of UCEs, and allow us to assess whether it is better to group all sites in a UCE into a single partition, or to split each UCE into more than one group of sites.

This study proposes and evaluates two partitioning methods for phylogenomic studies of UCEs: UCE Site Position (UCESP) and Sliding-Window Site Characteristics (SWSC). The UCESP method groups nucleotide sites across UCEs using their physical location within the UCE, for example by grouping all of the central sites of each UCE into a single partition. The SWSC method uses proxies of rates and patterns of molecular evolution [e.g., entropy (EN), multinomial likelihoods (MUs), GC content] and a sliding-window approach to determine whether a central region of a UCE (which may loosely correspond to what is often called the ‘core’) evolves in a different way to the two flanking regions. This approach naturally splits each UCE into three data blocks corresponding to the underlying structure of these phylogenomic markers (i.e., conserved cores and more variable flanks).

Each of these methods conducts partitioning independently of estimating phylogenetic trees. One strength of these

methods is their specificity to UCEs, whose use is increasing in phylogenomics. A key aim of phylogenetic model selection is to find models that capture the key biological features of the underlying data with a modest number of parameters (Kolaczowski and Thornton 2004; Steel 2005). Here, we seek to achieve this by leveraging the known biological properties of UCEs themselves. In this respect, our approach is similar to biologically inspired models such as codon-based partitioning, the power of which is largely derived from building on the known biological properties of protein-coding DNA sequences.

New Approaches

Partitioning methods are based on the assumption that sites within a partition have evolved under similar conditions. We first sought to ascertain whether this is the case for UCEs by measuring three properties of each site in each UCE: the EN of a site, which can serve as a rough proxy for the rate of evolution of that site in the absence of a known phylogenetic tree; the GC content (GC) of a site; and the MU of a site, which describes the likelihood of observing a particular site pattern given the observed base frequencies of a particular UCE. Visualizations of the patterns of EN, GC, and MU with each of the 1000s of UCEs across seven diverse data sets (table 1, fig. 1) show that all three properties contain considerable and predictable variation within UCEs. EN and MU are low in the central region of the UCE and higher in the flanks, and the GC is high in the central region and lower in the flanks. Figure 1 shows that this variation can be very large—for example, in many cases GC content is <20% at the end of a UCE but >50% in the center.

Commonly-used models of molecular evolution (e.g., GTR models) can account for variation in rates of molecular evolution among sites within a partition using approximations such as the gamma distribution, proportion of invariant sites, and free-rates models. However, all commonly-used models of molecular evolution assume that the base frequencies (and thus GC) of all sites in a single partition are drawn from a single distribution. Large and predictable variation of GC within UCEs appears to be the rule rather than the exception (fig. 1). Together with the predictable variation in EN (fig. 1) this suggests that dividing UCEs up into more than one partition might improve models of molecular evolution by accounting for variation in GC and rates of molecular evolution among sites in ways that are not possible with standard unpartitioned Markov models.

Table 1. Data Set Names and References, Clade Names, and Summary Information of the UCE Alignments Used to Evaluate the Two New Partitioning Methods.

UCE data set	Clade (scientific name)	Clade (common name)	Size (bp)	UCEs	OTUs	GC%	References
Branstetter	Aculeata	Stinging wasps	183747	807	187	46	Branstetter et al. (2017)
Crawford	Sauria	Diapsids	465241	1143	10	38	Crawford et al. (2012)
Harrington	Pleuronectiformes	Flatfishes	235232	999	55	44	Harrington et al. (2016)
McCormack	Neoaves	Birds	539526	1537	33	37	McCormack et al. (2013)
Meiklejohn	Phasianidae	Gallopeasants	599627	1479	18	43	Meiklejohn et al. (2016)
Moyle	Oscines	Songbirds	375172	515	106	40	Moyle et al. (2016)
Prebus	Temnothorax	Acorn ants	1561581	2098	50	44	Prebus (2017)

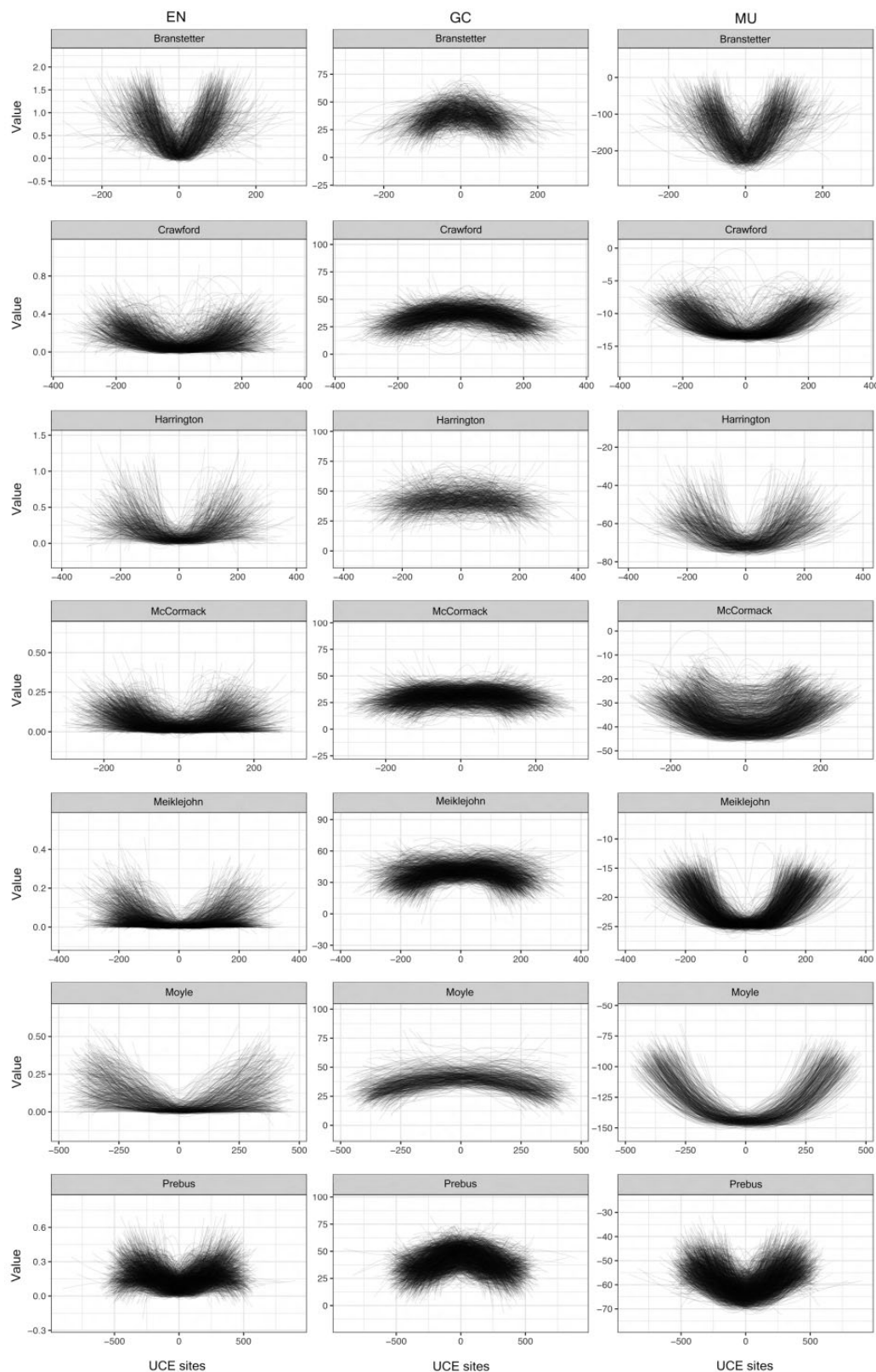


FIG. 1. Rates and patterns of evolution vary within UCEs. These plots highlight the predictable and sometimes dramatic variation in rates and patterns of molecular evolution within UCEs. Each line in each plot represents a single UCE. Each row represents a single data set (see [table 1](#)), and each column represents a metric measured for each site in a UCE. EN denotes the EN of a site, GC denotes the GC content of a site, and MU denotes the multinomial likelihood of a site. A value of 0 on the X-axis represents the central site of each UCE. Values among neighboring sites in each UCE are smoothed using the `geom_smooth()` function of `ggplot2`.

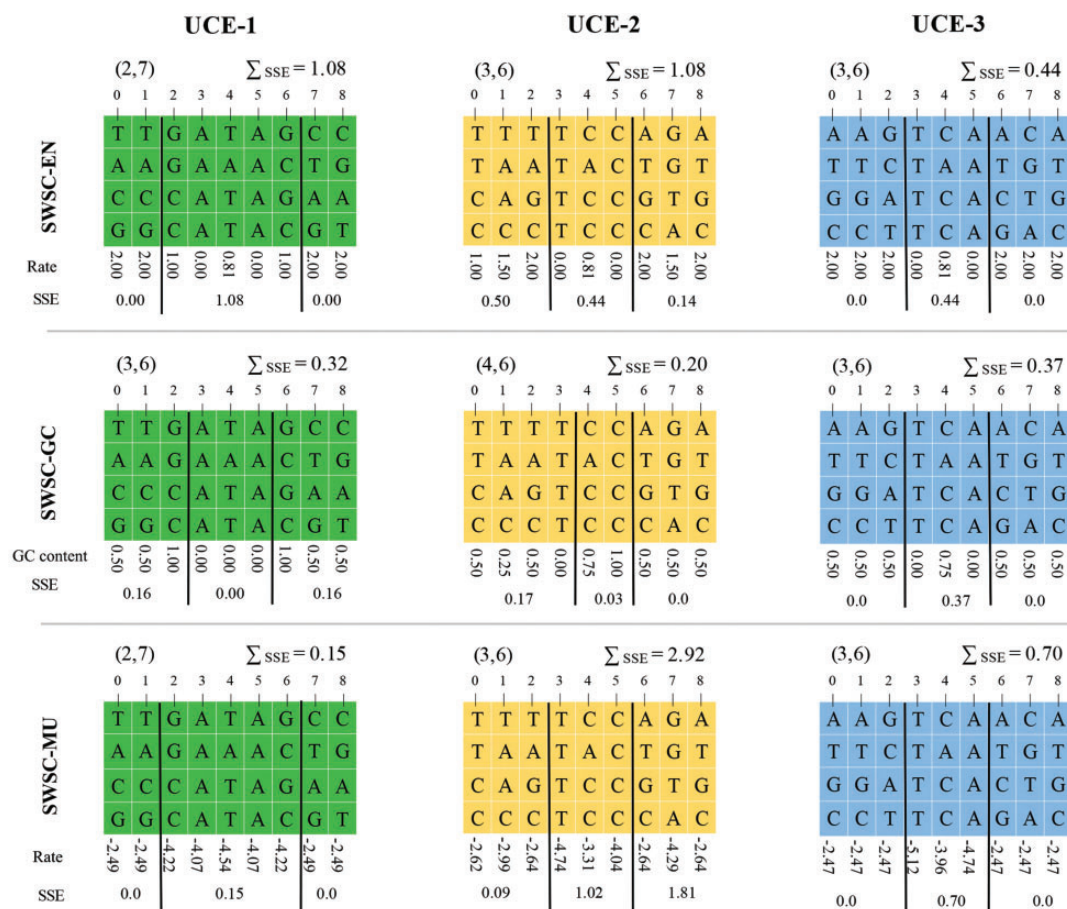


Fig. 2. SWSC method for partitioning UCEs. Schematic diagrams illustrating major steps of the SWSC algorithm. This diagram includes three hypothetical alignments: UCE-1 (green), UCE-2 (yellow), and UCE-3 (blue). The alignments include the same patterns of molecular evolution (i.e., EN, GC, and MU rates) as seen in the UCE markers. Note the alignments are comprised of conserved cores and variable flanking regions. The SWSC algorithm includes three steps. First, it proposes all combinations of three-window models in the alignments. It delimitates windows by locating all conceivable pairs of nucleotide sites in the alignments. Second, it estimates site-wise molecular evolution and nucleotide proportions/counts across the alignments. We used three alternatives properties of UCEs: (1) EN (SWSC-EN), (2) GC content (SWSC-GC), and (3) MUs (SWSC-MU). See further details in the New Approaches section. Third, it calculates the SSE statistics across windows and sums them up to obtain the sum of SSEs for every three data block model. The SWSC algorithm selects the best models by minimizing the squared residuals among three windows. The diagrams illustrate the best three-window models as indicated by the SWSC-GC, SWSC-EN, and SWSC-MU, respectively. The vertical bars delimit the three data blocks. The windows (i.e., size of the data blocks) are indicated by the pair of numbers in parenthesis.

We propose two new methods to automatically divide individual UCEs into multiple partitions based on their characteristic patterns of molecular evolution: SWSC, and UCESP. These two methods represent pragmatic approaches to potentially improve on treating each UCE as a single partition. The SWSC uses a sliding-window approach to divide each UCE into three data blocks. The choice of three data blocks is motivated by the observed patterns of variation in figure 1: all three measured site characteristics show that predictable variation focused on the center of each UCE. The SWSC is graphically summarized in figure 2. We describe its algorithm in detail in the Materials and Methods. The UCESP method takes a different approach, and groups nucleotides sharing similar locations in the UCEs into data blocks. In contrast to the SWSC, which divides the sites of each UCE into three data blocks, the UCESP places almost every site of a UCE into a different data block that contains similarly-positioned sites

from almost every other UCE. This approach is motivated by the relatively predictable variation in the three measured site characteristics within UCEs and across diverse data sets (table 1, fig. 1). The UCESP is graphically summarized in figure 3. We describe its algorithm in detail in the Materials and Methods.

Both the SWSC and UCESP methods group together putatively similar sites into data blocks, but neither asks whether any of the resulting data blocks are similar to one another and so both risk over-partitioning the data. For this reason we use PartitionFinder 2 (Lanfear et al. 2016) to estimate optimal partitioning schemes by grouping together similar data blocks from the output of the SWSC and UCESP. Given the optimal partitioning scheme for each of the evaluated new method, we then define the best method for each data set as the one that results in the partitioning scheme with the best (i.e., lowest) AICc score.

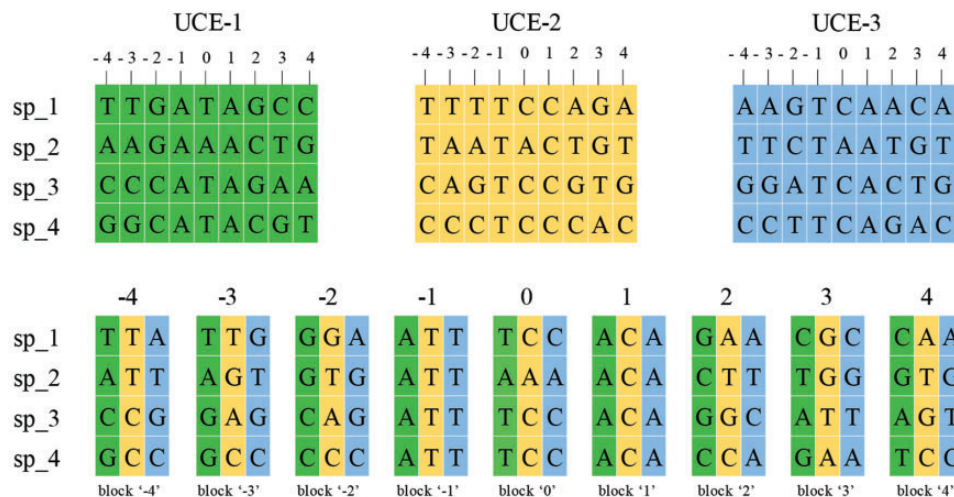


Fig. 3. UCESP for partitioning UCE alignments. Diagram illustrating the second new method of proposing partitions using the locations of nucleotide sites among UCEs. This method relies on the assumption the UCE regions (e.g., cores and flanking regions) have evolved under similar evolutionary processes. See further details in the New Approaches section. This diagram includes three hypothetical alignments: UCE-1 (green), UCE-2 (yellow), and UCE-3 (blue). The position of nucleotide sites in the alignments are indicated by the numbers of each site. The UCESP algorithm includes two steps. First, define the central site of the UCE as site 0 (zero) and sites to the left of this site are labeled with negative numbers, and sites to the right of this site are labeled with positive numbers. Second, it is to create one data block for each label, by combining all the sites with the same label into a single data block. Note that there are nine partitions, all sharing the same locations of sites among the UCEs.

Results

Visual Inspection of the Properties of UCEs

We measured three properties of UCE sites: the EN of a site, the GC content (GC) of a site, and the MU of a site. We observed that EN and MU are lower at the center of each UCE and higher at the edges, and that this pattern is reversed for GC (figure 1). Figure 1 also shows that the absolute values of EN, GC, and MU can vary substantially between data sets, as expected given that each data set represents a particular taxonomic clade sampled at a particular phylogenetic depth. For example, the mean EN varies roughly 6-fold from 0.03 in the gallopheasant data set (Meiklejohn et al. 2016) to 0.19 in the flatfish data set (Harrington et al. 2016), and the mean GC content varies from 37% in the bird data set to 44% in the stinging wasp data set (table 1).

The SWSC Algorithm Divides UCEs into Data Blocks That Reflect Site Properties

The SWSC partitioning method uses a site property (i.e., EN, GC, or MU) to split each UCE into three data blocks (fig. 4). As expected given our observations in figure 1, the EN and MU are lower in the central data blocks and higher in the edge data blocks, and GC shows the opposite pattern (table 2). In a small proportion of cases (~16% or 910 UCEs across all seven data sets) a UCE could not be split into three data blocks because there was no solution that satisfied the criteria that each data block had to contain at least 50 sites, all four nucleotides, and at least one variable site (see Materials and Methods). Most of the UCEs set to single partitions were smaller than 150 bp and were found in the stinging wasp data set (table 1, fig. 4).

The SWSC-EN Method Outperforms Other Approaches

The SWSC approach using site EN to derive data blocks (SWSC-EN) outperformed all other methods on all seven empirical data sets (table 3, fig. 5). We compared seven approaches to partitioning UCEs, comprising the new methods proposed here and three partitioning methods widely used in phylogenomic studies: (1) SWSC-EN; (2) SWSC-GC; (3) SWSC-MU; (4) UCESP; (5) Single (i.e., all UCEs treated as one partition); (6) UCE (i.e., each UCE treated as a partition); (7) and PF-UCE (i.e., a partitioning scheme estimated by defining each UCE as an initial data block and optimizing the partitioning scheme in PartitionFinder). We used PartitionFinder 2 (Lanfear et al. 2016) to optimize the partitioning schemes for methods (1)–(4) and method (7), see Materials and Methods. In all cases, the SWSC-EN method received the lowest AICc score by a considerable margin. The closest AICc score to the SWSC-EN method for a single data set was more than 2,000 points higher, for the SWSC-MU approach on the Branstetter data set (table 3). The three standard approaches (Single, UCE, and PF-UCE) performed a great deal worse than all of the SWSC methods on all data sets. For example, the smallest difference between any of the three standard approaches and the SWSC-EN method was >60,000 AICc units for the Meiklejohn data set (table 3). The UCESP method performed poorly on all data sets, achieving the second-worst AICc scores in more than half of the seven data sets (table 3, fig. 5); that is worse only than treating the entire alignment as a single partition.

Comparison of Observed and Shuffled UCE Alignments

A potential concern with the SWSC and UCESP approaches is that they may achieve their improvement in AICc scores

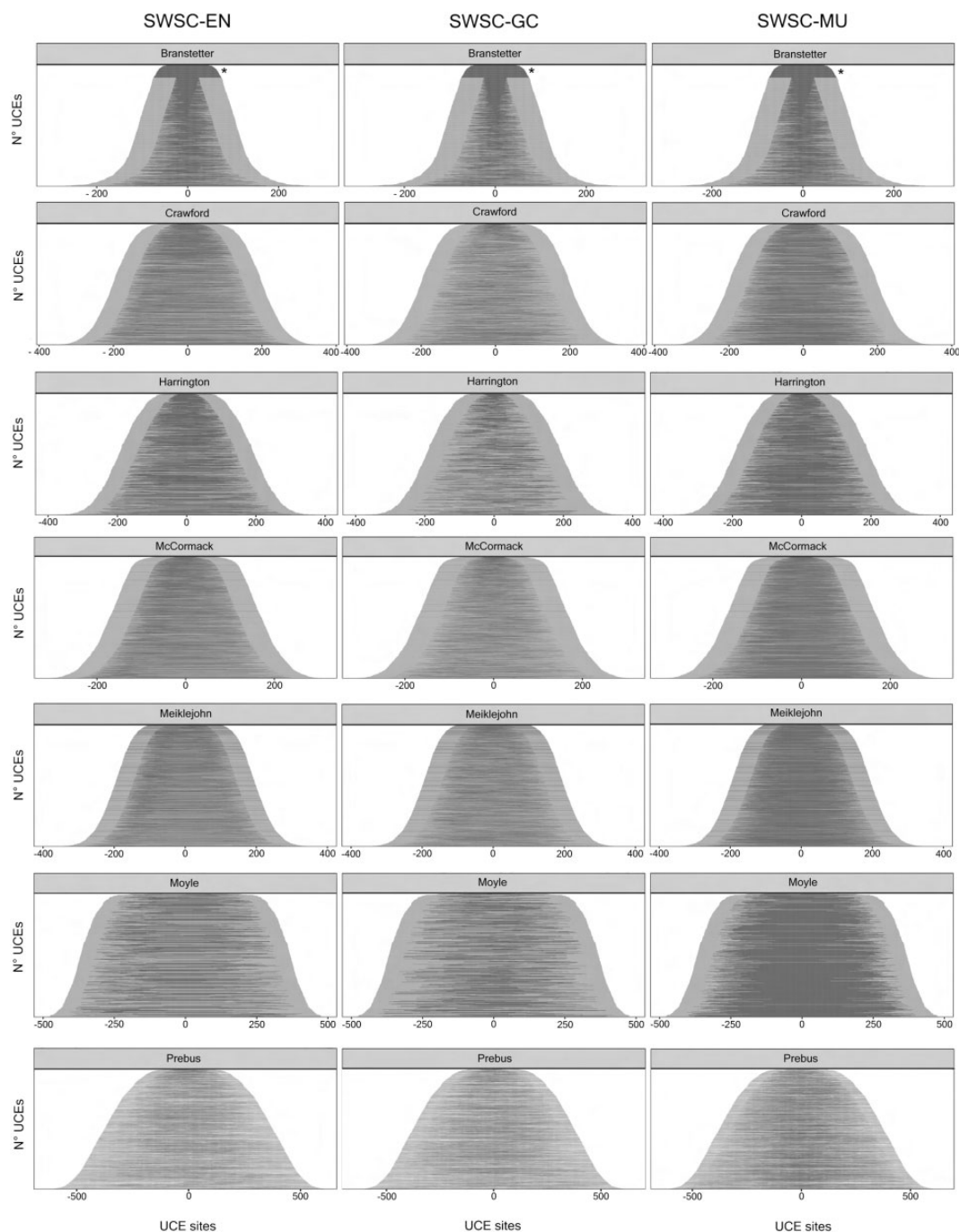


FIG. 4. Best three data blocks for each UCE. Graphical representations of the best three data block for each UCE of each data set (see [table 1](#)) as indicated by the SWSC method. Each line in each plot represents a single UCE split in three data blocks. Black indicates the position of the central window, which may loosely correspond to what is often called the core region of UCEs, and gray indicates the position of the edge windows, which may loosely correspond to what is often called the flanking regions of UCEs. Each row of panels represents a single data set (see [table 1](#)), and each column of panels represents a metric measured for each site in a UCE. EN denotes the entropy of a site, GC denotes the GC content of a site, and MU denotes the multinomial likelihood of a site. UCEs are organized by increasing length. Asterisks (*) indicate UCEs smaller than 150 bp, which were not split into three data blocks but kept as a single data block (see details in the Materials and Methods: Sliding-Window Site Characteristics).

simply by defining a large number of starting subsets for PartitionFinder. This would define a larger search space of potential partitioning schemes for a data set, potentially accounting for the improved AICc scores. To assess this possibility, we created 25 permutations of all 7 data sets, in which we shuffled the sites within each UCE. We then re-ran all of the SWSC-EN analyses on the permuted alignments. This

compares the observed SWSC-EN solution to one in which the assignment of sites to data blocks within each UCE is randomized, while other characteristics of the partitioning scheme such as the number and size of the starting subsets are held constant. Across all seven empirical data sets ([table 1](#)), and for all 25 permutations of the UCE alignments, the shuffled UCE alignments produced

Table 2. Means of EN and GC of Each Window and Data Set Produced by the SWSC Method.

	Branstetter	Crawford	Harrington	McCormack	Meiklejohn	Moyle	Prebus	Overall
SWSC-EN								
Left	0.828	0.212	0.278	0.097	0.055	0.140	0.182	0.256
Center	0.236	0.072	0.090	0.031	0.015	0.054	0.122	0.088
Right	0.831	0.145	0.267	0.059	0.036	0.090	0.181	0.230
SWSC-GC								
Left	32.91	31.39	40.12	28.42	36.10	33.61	36.58	34.16
Center	43.10	41.02	46.68	34.09	43.38	44.38	48.04	42.96
Right	32.24	30.99	40.47	28.75	36.61	33.5	36.94	34.21

Table 3. AICc Scores, Number of Subsets, and Parameters of Partitioning Schemes Inferred From Seven Partitioning Strategies Across Seven UCE Data Sets.

Data set	Partitioning strategy	AICc	ΔAICc	Subsets	Parameters
Branstetter	Single	17672426	−336594	1	380
	UCESP	17591017	−255186	158	1950
	UCE	17581458	−245627	807	8440
	PF+UCE	17575239	−239408	475	5120
	SWSC-GC	17415498	−79666	1018	10550
	SWSC-MU	17338096	−2264	948	9850
	SWSC-EN	17335832	0	948	9850
Crawford	Single	2481257	−100299	1	26
	UCESP	2458689	−77731	85	866
	UCE	2467719	−86761	1145	11466
	PF+UCE	2456063	−75104	222	2236
	SWSC-GC	2410728	−29770	456	4576
	SWSC-MU	2397171	−16213	461	4626
	SWSC-EN	2380958	0	452	4536
Harrington	Single	3833182	−145228	1	116
	UCESP	3800797	−112843	114	1246
	UCE	3796629	−108675	596	6066
	PF+UCE	3793471	−105517	267	2776
	SWSC-GC	3736560	−48606	513	5236
	SWSC-MU	3695067	−7112	493	5036
	SWSC-EN	3687954	0	493	5036
McCormack	Single	3192533	−137223	1	72
	UCESP	3166286	−110976	101	1072
	UCE	3162976	−107666	1541	15472
	PF+UCE	3147458	−92148	318	3242
	SWSC-GC	3088673	−33363	615	6212
	SWSC-MU	3094082	−38772	638	6442
	SWSC-EN	3055310	0	586	5922
Meiklejohn	Single	2339168	−100584	1	42
	UCESP	2314874	−76290	97	1002
	UCE	2316404	−77820	1479	14822
	PF+UCE	2299674	−61091	249	2522
	SWSC-GC	2255398	−16814	463	4662
	SWSC-MU	2254129	−15545	471	4742
	SWSC-EN	2238583	0	458	4612
Moyle	Single	6005672	−230918	1	218
	UCESP	5936152	−161398	200	2208
	UCE	5937377	−162623	515	5358
	PF+UCE	5935114	−160361	271	2918
	SWSC-GC	5839663	−64910	520	5408
	SWSC-MU	5820918	−46165	496	5168
	SWSC-EN	5774753	0	496	5168
Prebus	Single	20339794	−625451	1	106
	UCESP	20234314	−519970	178	1876
	UCE	20173076	−458732	2098	21076
	PF+UCE	20161619	−447276	806	8156
	SWSC-GC	19835806	−121462	1141	11506
	SWSC-MU	19766846	−52503	1138	11476
	SWSC-EN	19714343	0	1125	11346

NOTE.—(1) Single: treating all sites as belonging to a single subset; (2) UCE: one subset for each UCE alignment; (3) PF+UCE: each UCE was defined as a data block; (4) SWSC-GC: data blocks defined by the SWSC-GC algorithm; (5) SWSC-EN: data blocks defined by the SWSC-EN algorithm; (6) SWSC-MU: data blocks defined by the SWSC-MU algorithm; and (7) UCESP: data blocks defined by the UCESP algorithm. In (3)–(7), the final partitioning scheme was optimized in PartitionFinder 2. Bold: best partitioning scheme.

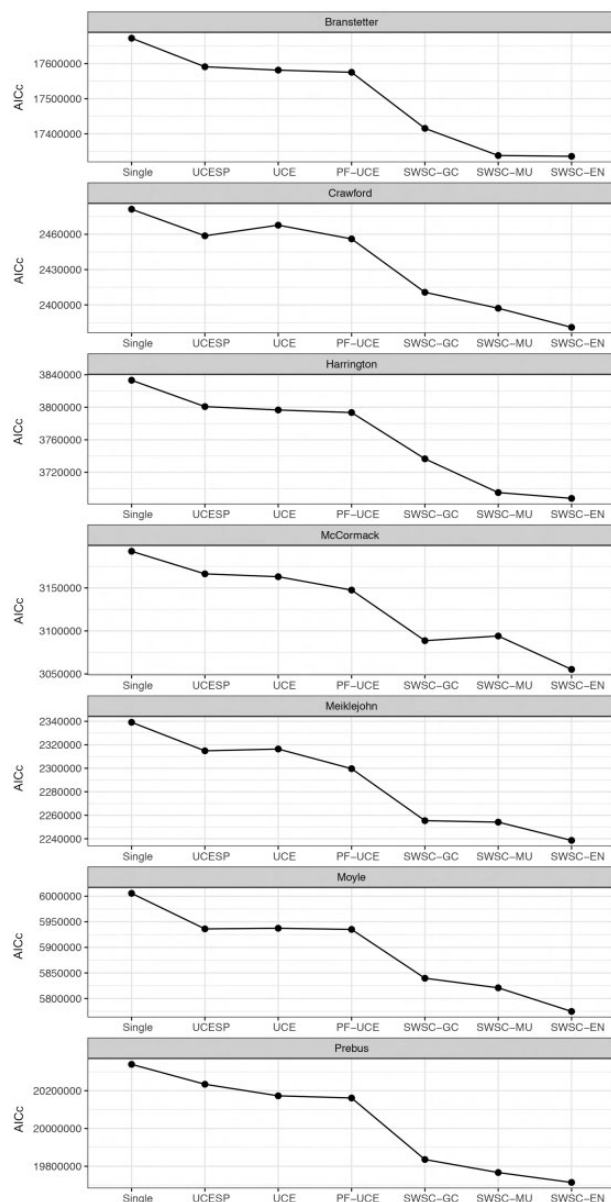


FIG. 5. The SWSC partitioning method leads to substantial improvements in model-fit. AICc scores of partitioning schemes derived from the SWSC outperform UCESP and other methods applied in phylogenomics. The plots show the AICc scores (Y-axis) of seven partitioning schemes (X-axis). The seven partitioning strategies were: (1) Single: treating all sites as belonging to a single subset; (2) UCE: one subset for each UCE alignment; (3) PF-UCES: each UCE was defined as a data block; (4) SWSC-GC: data blocks defined by the SWSC-GC algorithm; (5) SWSC-EN: data blocks defined by the SWSC-EN algorithm; (6) SWSC-MU: data blocks defined by the SWSC-MU algorithm; and (7) UCESP: data blocks defined by the UCESP algorithm. In (3)–(7), the final partitioning scheme was optimized in PartitionFinder 2. The AICc scores indicate that the SWSC, particularly the SWSC-EN, produces partitioning schemes that outperform all other partitioning strategies investigated here. Surprisingly, the partitioning schemes derived from the UCESP obtained AICc scores that were worse than partitioning schemes widely used in phylogenomics (i.e., UCE and UCE-PF).

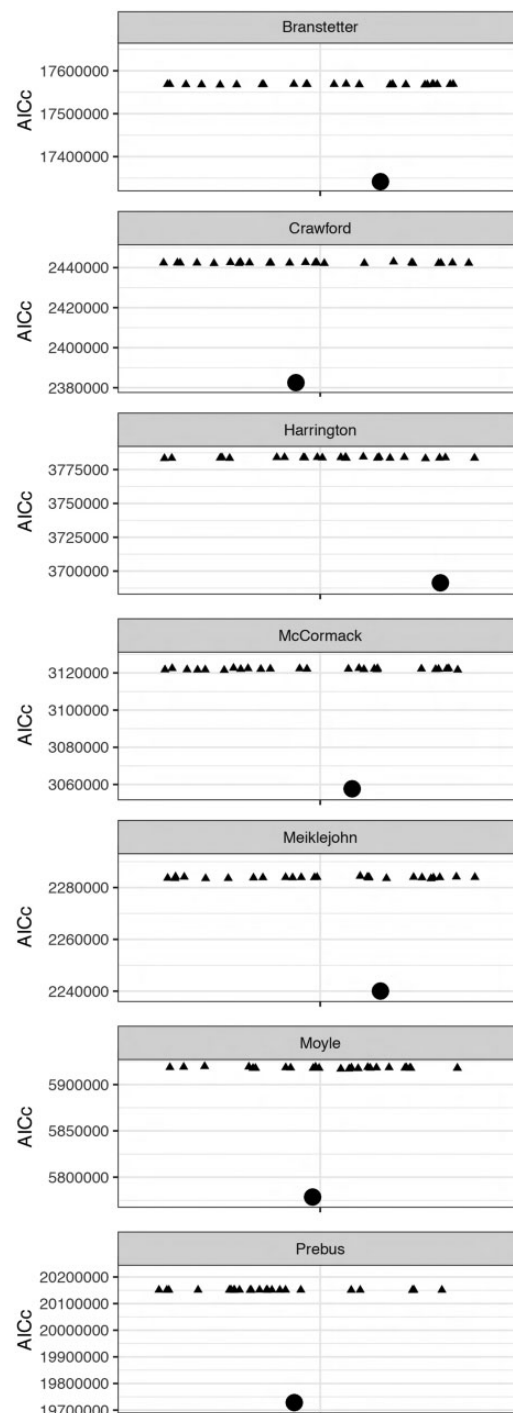


FIG. 6. SWSC-EN split UCEs into meaningful partitions. Comparisons between the AICc scores of empirical data sets (circle) and 25 permuted data sets (triangles) of partitioning schemes derived from the SWSC-EN and optimized in PartitionFinder 2. For each of the randomizations, the sites within each UCE were shuffled and the new alignments were used with the original data block definitions from the un-shuffled data as input to PartitionFinder 2. Each panel includes a data set and 25 AICc scores of partitioning schemes optimized in PartitionFinder 2. The AICc scores suggest that the SWSC-EN method improves model-fit and parameter estimates not because it searches a larger space of possible partitioning schemes, but primarily because splitting each UCE into three data blocks is uncovering biological differences in rates and/or patterns of molecular evolution within each UCE.

Table 4. Measurements of TL (the Sum of All Branch Lengths in Substitutions Per Site) and PD of Each Topology Compared to the SWSC-EN Topology for Trees Estimated with the Seven Different Partitioning Schemes Evaluated in this Study.

Partition strategy	TL	PD
Branstetter		
Single	1.081	89.5
UCESP	1.038	514.2
UCE	0.982	91.7
PF-UCE	1.052	80.7
SWSC-GC	1.006	112.2
SWSC-MU	1.005	170.5
SWSC-EN ^a	1.000	0.0
Crawford		
Single	1.289	0.0
UCESP	1.252	0.0
UCE	1.213	0.0
PF-UCE	1.250	0.0
SWSC-GC	0.983	0.0
SWSC-MU	0.807	0.0
SWSC-EN ^a	1.000	0.0
Harrington		
Single	1.141	0.0
UCESP	1.174	0.0
UCE	0.837	0.0
PF-UCE	0.958	0.0
SWSC-GC	0.903	0.0
SWSC-MU	1.028	0.0
SWSC-EN ^a	1.000	0.0
McCormack		
Single	1.540	24.6
UCESP	1.437	28.4
UCE	1.388	25.5
PF-UCE	1.467	0.0
SWSC-GC	1.088	12.7
SWSC-MU	0.961	24.8
SWSC-EN ^a	1.000	0.0
Meiklejohn		
Single	1.580	0.0
UCESP	1.506	0.0
UCE	1.523	0.0
PF-UCE	1.543	0.0
SWSC-GC	1.234	0.0
SWSC-MU	1.389	0.0
SWSC-EN ^a	1.000	0.0
Moyle		
Single	0.973	117.5
UCESP	0.971	85.4
UCE	0.969	20.2
PF-UCE	0.970	112.8
SWSC-GC	0.957	99.8
SWSC-MU	0.941	27.1
SWSC-EN ^a	1.000	0.0
Prebus		
Single	1.021	0.0
UCESP	1.010	23.7
UCE	1.010	0.0
PF-UCE	1.021	11.9

(continued)

consistently much worse (i.e., higher) AICc scores than the un-shuffled alignments (fig. 6). This suggests that the improvement in AICc score seen in the SWSC-EN method is not due solely to having an increased number of starting subsets.

Table 4. Continued

Partition strategy	TL	PD
SWSC-GC	0.980	11.9
SWSC-MU	0.754	16.2
SWSC-EN ^a	1.000	0.0

NOTE.—To aid comparison of tree lengths, the tree lengths of all trees for a given data set are scaled such that the tree length of the tree estimated under the SWSC-EN partitioning scheme is equal to 1.0. Larger PD indicate larger differences between tree topologies, but they cannot be directly compared between data sets.

^aThe tree topology estimated under the partitioning scheme obtained by the SWSC-EN was used as reference. (1) Single: treating all sites as belonging to a single subset; (2) UCE: one subset for each UCE alignment; (3) PF-UCE: each UCE was defined as a data block; (4) SWSC-GC: data blocks defined by the SWSC-GC algorithm; (5) SWSC-EN: data blocks defined by the SWSC-EN algorithm; (6) SWSC-MU: data blocks defined by the SWSC-MU algorithm; and (7) UCESP: data blocks defined by the UCESP algorithm. In (3)–(7), the final partitioning scheme was optimized in PartitionFinder 2.

Phylogenetic Inference

We estimated ML tree topologies for each of the seven data sets (table 1) under each of the seven partitioning schemes used in this study (i.e., SWSC-EN, SWSC-GC, SWSC-MU, UCESP, Single, UCE, PF-UCE). The resulting trees are presented in the [Supplementary Material](#) online. For each data set, we calculated the tree lengths of every tree, and the Path Difference (PD) between the SWSC-EN topology and all of the other topologies inferred under the other six partitioning schemes (table 4).

Overall, tree lengths were similar among all seven partitioning schemes, and the SWSC-EN tree length showed no clear pattern of difference to the tree lengths from other partitioning schemes (table 4, fig. 7). The PDs show that the choice of partitioning scheme influenced the inferred phylogenetic tree in four out of seven data sets (table 4). Visual examination of the differences (see [Supplementary Material](#) online) reveals that most of the observed topological differences are associated with nodes with low bootstrap support (< 70, see [Supplementary Material](#) online), suggesting that the choice of partitioning scheme rarely influenced the resulting tree topology in ways that would affect biological inference. However, in at least one case, the SWSC-EN partitioning scheme led to a strongly-supported and biologically important change in the tree topology when compared to trees estimated with traditional approaches to partitioning. In this case, the phylogenetic position of the Sclerogibbidae, a family of wasps analyzed in the study of [Branstetter et al. \(2017\)](#), was sensitive to the partitioning scheme used. The original study [Branstetter et al. \(2017\)](#) and two of our analyses using standard partitioning schemes (i.e., a single partition, or one partition for each UCE) place the Sclerogibbidae together with a clade comprising the Embolemidae + Dryinidae, with the three families forming the sister group to the remaining non-chrysidoid lineages in the phylogeny [i.e., ((Sclerogibbidae, (Embolemidae, Dryinidae)), remainder)]. Bootstrap support for the former grouping varies considerably depending on the details of the analysis both in the original study (59–90% support) and in our analyses (52% support using a single partition, and 100% support using a partition for each UCE).

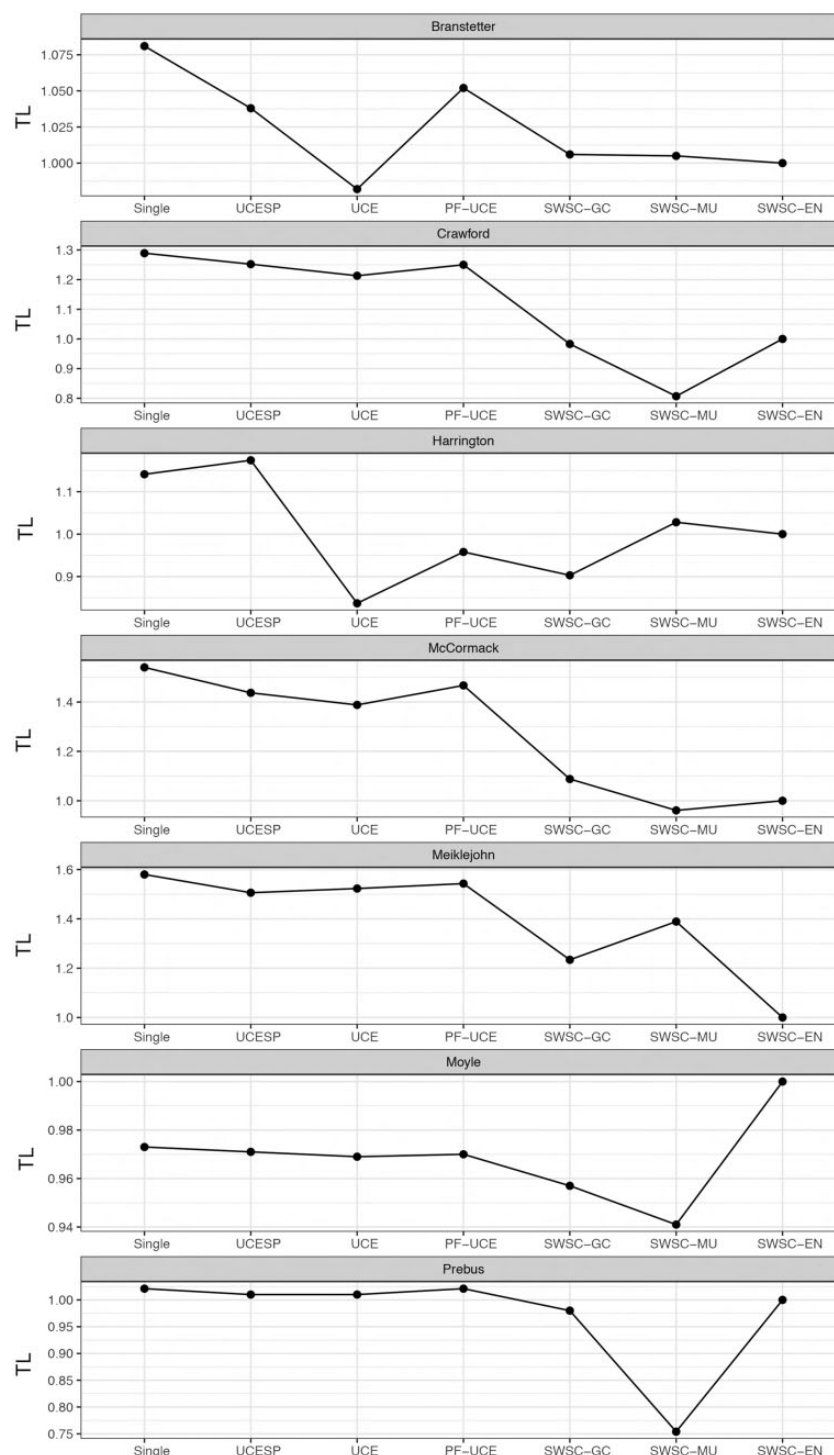


Fig. 7. Tree lengths (TL) of phylogenetic trees estimated under seven partitioning schemes and across seven data sets. To aid comparison of tree lengths, the tree lengths of all trees for a given data set are scaled such that the tree length of the tree estimated under the SWSC-EN partitioning scheme is equal to 1.0. There is no clear trend between TL values and partitioning schemes across the data sets. (1) Single: treating all sites as belonging to a single subset; (2) UCE: one subset for each UCE alignment; (3) PF-UCE: each UCE was defined as a data block; (4) SWSC-GC: data blocks defined by the SWSC-GC algorithm; (5) SWSC-EN: data blocks defined by the SWSC-EN algorithm; (6) SWSC-MU: data blocks defined by the SWSC-MU algorithm; and (7) UCESP: data blocks defined by the UCESP algorithm. In (3)–(7), the final partitioning scheme was optimized in PartitionFinder 2.

In contrast, the SWSC-EN partitioning scheme places the Sclerogibbidae as sister to the remaining nonchrysidoid lineages, with the other two families as sister to this grouping [i.e., ((Embolemyidae, Dryinidae), (Sclerogibbidae, remainder))],

with 100% bootstrap support for both groupings. Notably, the same grouping was recovered with 100% bootstrap support in the original study when they explicitly accounted for variation among gene trees when estimating the species tree

(Branstetter et al. 2017). It would be premature to suggest that any of these analyses are definitive, but it may be prudent to lean towards preferring the relationships supported by the model that best fits the data, which in the case of our analyses is the SWSC-EN partitioning scheme. These results show that on some occasions, the new methods we propose here may contribute to clarifying phylogenetic relationships.

Discussion

Model selection is an important part of phylogenetic inference (Sullivan and Joyce 2005; Shapiro et al. 2006; Lanfear et al. 2014). In this study, we focused on selecting partitioned models of molecular evolution for data set comprised of UCEs. We first showed that patterns of molecular evolution within UCEs vary in predictable ways (fig. 1). Importantly, this variation is likely to violate at least one key assumption of most commonly-used models of molecular evolution: that base composition within a UCE can be adequately modeled as a single vector of base frequencies. This suggests that model violation might limit the accuracy of inferences made from UCE data sets, as long as each UCE is assumed to have evolved under a single Markov model.

We proposed four new approaches to partitioning UCEs, and showed that an approach based on site entropies to divide each UCE into three data blocks dramatically improved the fit of the models to the data when compared to three commonly-used approaches to partitioning in UCE studies (fig. 5). Phylogenetic trees estimated using partitioning schemes from this new method (which we call the SWSC-EN method) showed some differences in tree length and tree topology when compared to trees estimated using standard approaches (table 4, fig. 7). These differences tended to be relatively minor, but since it is hard to predict a priori the effect of model choice on tree inference, and since the models estimated with the SWSC-EN method have much improved AICc scores, it seems prudent to employ these new models where appropriate. Indeed, in a single case we observed a strongly-supported change in a topology when comparing the tree estimated using the SWSC-EN method to trees estimated using more traditional methods such as treating all data as a single data block, or assigning each UCE to its own data block. The SWSC-EN method we describe here can be implemented for any UCE data set using the scripts we provide on Github at <https://github.com/Tagliacollo/PFinderUCE-SWSC-EN> (doi: 10.5281/zenodo.1028743).

Phylogenetics is no longer limited by our ability to sequence sufficient loci from a sample (Delsuc et al. 2005; Posada 2016). Rather, the current challenges have shifted to finding appropriate ways to model the vast quantities of data that we have (Kumar et al. 2012). We hope that the new methods we propose here will help improve the accuracy of phylogenetic inferences made from UCEs. However, we note that our approach has many limitations. For example, our observations (fig. 1) suggest that rates of evolution and

GC content vary in quite predictable ways within each UCE. Dividing each UCE up into three data blocks provides a crude but pragmatic way to model this variation within the current phylogenetic workflow, but it might be more appropriate to model this variation directly. For example, the patterns of rates of evolution and GC content within UCE could be modeled with four additional parameters added to a standard model from the GTR family: a minimum rate and a maximum GC content that applied to the center of the UCE, and a maximum rate and a minimum GC content that applied to the two ends of the UCE. Such an approach would likely provide a better description of the observed patterns of variation within each UCE, and would require fewer parameters than dividing each UCE up into three separate data blocks.

Conclusion

Partitioning remains a popular method for accounting for variation in rates of molecular evolution; however, it relies on the assumptions that predefined groups of sites have evolved under the same process. We present a new method that improves phylogenomic estimates of species relationships by improving partitioning schemes of UCE alignments. This method can be implemented using scripts available from GitHub at <https://github.com/Tagliacollo/PFinderUCE-SWSC-EN> (doi: 10.5281/zenodo.1028743). All of the methods we describe here can be implemented using scripts available at <https://github.com/Tagliacollo/PartitionUCE> (doi: 10.5281/zenodo.1027526).

Materials and Methods

Empirical Data Sets

We evaluated the performance of the two new partitioning methods described here using seven empirical UCE data sets available from the Dryad Digital Repository. The data sets range from 515 to 2,098 UCEs, from 10 to 187 taxa, and from 183,747 to 1,561,581 nucleotide sites (table 1). Non-UCE markers were excluded from one data set (Meiklejohn et al. 2016). The modified version is available from GitHub along with all of the other data sets we used for this study at <https://github.com/Tagliacollo/PartitionUCE> (doi: 10.5281/zenodo.1027526).

Visualizing Patterns of Molecular Evolution in UCEs

We measured three properties of molecular evolution of each site in each UCE of the seven empirical UCE data sets used in this study (table 1): the EN of a site, the GC content of a site; and the MU of a site. We calculate the entropy (H) of each site using the following formula:

$$H_i = - \left(\sum_{j=1}^4 p_j \log_2 p_j \right), \quad (1)$$

where $j = 1, 2, 3$, and 4 corresponds to nucleotide A, C, G, and T, and p_j corresponds to the proportion of a nucleotide j at site i . The H_i varies between 0 and 2, where 0 indicates

invariant sites and 2 indicates sites with an equal frequency of all four nucleotides.

We calculate the GC content as the ratio of the sum of the counts of G and C to the sum of the counts of all four nucleotides, ignoring gaps and ambiguous nucleotides. Finally, we calculate the multinomial likelihood (M) using the following formula:

$$M_i = (p_j + \dots + p_4)^N = \sum_{b_j + \dots + b_4 = N} \left(\frac{N!}{b_j! \dots b_4!} \right) \prod_{j=1}^4 p_j^{b_j}, \quad (2)$$

where

$$\left(\frac{N!}{b_j! \dots b_4!} \right) = \frac{N!}{b_j! \dots b_4!}, \quad (3)$$

where N corresponds to the number of species, $j = 1, 2, 3$, and 4 corresponds to nucleotides A, C, G, and T, respectively, p_j represents the proportion of nucleotide j at site i , and b_j represents the counts of nucleotides in the alignment of the UCE.

New Methods for Automated Partitioning of UCEs

In what follows, we use the word ‘data block’ to refer to a user- or algorithmically defined set of sites that are assumed to have evolved in similar ways. We refer to a ‘subset’ as a group of one or more data blocks, and we refer to a ‘partitioning scheme’ as a group of subsets in which each site in the alignment is included only once.

The new partitioning methods, the SWSC and UCESP, were developed specifically for phylogenomic studies of UCE markers. The SWSC uses a sliding-window approach to divide each UCE into three data blocks, which can then be combined across UCEs using standard algorithms in PartitionFinder 2 (Lanfear et al. 2016). The UCESP takes a different approach, and groups nucleotide sites sharing similar locations in the alignments into partitions. We implemented both methods using Python scripts that are available on GitHub at <https://github.com/Tagliacollo/PartitionUCE> (doi: 10.5281/zenodo.1027526).

Both methods require the following input: a concatenated nexus alignment comprised of UCE markers and including nexus-formatted character sets (charsets) that define the location of each UCE in the alignment. The methods export PartitionFinder configuration files including data blocks defined by the individual methods. We avoid defining data blocks with fewer than 50 nucleotide sites that do not contain all four nucleotides, or have no variable sites. These conditions avoid the creation of very small data blocks, which can provide unreliable parameter estimates and thus mislead algorithms such as PartitionFinder 2 (Lanfear et al. 2016). The occurrence of all four nucleotide bases and at least one variable site in each data block was necessary to ensure that RAXML (Stamatakis et al. 2008) could analyze that data block. Example input files can be found in the ‘raw_data’ folder available on

GitHub at <https://github.com/Tagliacollo/PartitionUCE> (doi: 10.5281/zenodo.1027526).

Sliding-Window Site Characteristics (SWSC)

The SWSC is graphically summarized in figure 2, and includes the following four steps which we describe in more detail below:

- (1) Calculate a metric (i.e., EN, GC content, or MU) for each UCE site.
- (2) Define all valid ways of dividing the UCE into three contiguous groups of sites.
- (3) Calculate the total sum of squared errors (SSE) for each of the splits defined in (2).
- (4) Select the split from (3) with the smallest SSE.

Step 1 involves simply calculating a metric for each site of the UCE, as described in the New Approaches section: the EN of a site, which can serve as a rough proxy for the rate of evolution of that site; the GC content (GC); and the MU of a site, which describes the likelihood of observing a particular site pattern given the observed base frequencies of a particular UCE.

Step 2 of the SWSC algorithm is to propose all valid ways of dividing a UCE into three contiguous groups of sites, which we hereafter refer to as a split. We do this by first listing all possible splits for a single UCE. The number of such splits is large, and is defined by $\binom{N}{2}$, where N is the number of sites in the UCE, and the 2 refers to the positions of the two cuts required to split a single UCE into three contiguous parts. Given the list of all possible splits, we then reject splits that do not meet the following conditions: (1) all three sections defined by the split must contain at least 50 bases; (2) all three sections of the split must have at least one variable site; (3) all three sections of the split must contain at least one of each of the four nucleotide bases (see above). This results in a set of all valid splits for a single UCE.

Step 3 of the SWSC algorithm is to calculate the SSE for each of the splits defined in step 2. To do this, we first calculate for each valid split the SSE within each of the three contiguous groups of sites, by summing the absolute differences of the metric at each site and the mean of the metric for that group of sites (e.g., the sum of the absolute differences between the GC content of each site and the mean GC content for the window). We then sum the SSEs for the three groups of sites to obtain a total SSE for the split.

Step 4 of the SWSC algorithm is to define the best split as the one with the smallest total SSE. In cases where we have more than one split with the same minimum SSE, we choose the split that has the lowest variance in the lengths of the three contiguous groups of sites. This maximizes the sizes of each window and avoids small data blocks which could provide unreliable parameter estimates.

The outcome of the SWSC algorithm for a single UCE is a single split that defines three groups of contiguous sites, each of which is then used as a data block for input into PartitionFinder 2 (see below).

UCE Site Position (UCESP)

The UCESP is graphically summarized in [figure 3](#), and includes the following three steps, which we describe in more detail below:

- (1) Find the central nucleotide site for each UCE.
- (2) Create data blocks based on their locations relative to the central site in each UCE.
- (3) Merge small data blocks.

Step 1 of the UCESP is to define the central site of the UCE as site 0. Sites to the left of this site are labeled with negative numbers, and sites to the right of this site are labeled with positive numbers. Step 2 of the UCESP is to create one data block for each label, by combining all the sites with the same label into a single data block. For a data set with 1,000 UCEs, this will create many data blocks with 1,000 sites (one from each UCE). However, the longest UCEs in the data set will lead to the creation of much smaller data blocks, because they will have sites labeled with large numbers that are rarely found in other UCEs. Small data blocks cannot be practically analyzed; therefore, step 3 of the UCESP is to progressively merge small data blocks with their nearest neighbors until each data block meets the same criteria previously described in the step 2 of the SWSC method. This algorithm results in a single set of data blocks which can be used for downstream analyses in PartitionFinder 2.

Partitioning Schemes: PartitionFinder

We used PartitionFinder 2 ([Lanfear et al. 2016](#)) to estimate the optimal partitioning schemes and models of molecular evolution for seven different approaches to partitioning each of the seven data sets analyzed here ([table 1](#)): four of the partitioning approaches are described here (i.e., SWSC-EN, SWSC-GC, SWSC-MU, UCESP), and we compare these to three partitioning approaches that are widely used in phylogenomic studies of UCEs (treating the entire alignment as one partition; assigning one partition to each UCE; and assigning one partition to each UCE and optimizing this scheme in PartitionFinder). Given an optimal partitioning scheme for each method, we then define the best method for each data set as the method that results in the partitioning scheme with the best (i.e., lowest) AICc score. We used PartitionFinder 2 with the following settings: start with a tree estimated through maximum parsimony, linked branch lengths, and GTR + G model of nucleotide evolution, and the relaxed clustering algorithm with default settings. For each data set, we calculated the AICc scores of the following partitioning schemes: (1) Single: treating all sites as belonging to a single subset; (2) UCE: one subset for each UCE alignment; (3) PF-UCE: each UCE was defined as a data block; (4) SWSC-GC: data blocks defined by the SWSC-GC algorithm; (5) SWSC-EN: data blocks defined by the SWSC-EN algorithm; (6) SWSC-MU: data blocks defined by the SWSC-MU algorithm; and (7) UCESP: data blocks defined by the UCESP algorithm. In

(3)–(7), the final partitioning scheme was optimized in PartitionFinder 2.

Checking Reliability of SWSC-EN

To evaluate whether the performance of the SWSC-EN method results from simply from splitting the UCEs into smaller subsets, we compared the observed AICc score for each of the seven empirical data sets to 25 AICc scores calculated from the same data sets in which the sites of each UCE were shuffled. Shuffling the sites of each UCE serves to hold the number and size of initial data blocks input to PartitionFinder 2 constant, but within each UCE it randomizes the sites that are assigned to each of the three data blocks. For each of the 25 randomizations for each of the seven empirical data sets (175 analyses in total) we (1) shuffled the sites within each UCE; (2) used the shuffled alignments with the original data block definitions from the un-shuffled data as input to PartitionFinder 2; (3) optimized the partitioning as described above. The scripts and permuted data sets are available from GitHub at <https://github.com/Tagliacollo/PartitionUCE> (doi: 10.5281/zenodo.1027526).

Phylogenomic Inferences: Branch Order and Branch Lengths

We used IQ-Tree ([Nguyen et al. 2015](#)) to infer phylogenetic trees for each of the seven empirical data sets ([table 1](#)), under each of the seven partitioning schemes. The aim was to evaluate whether more appropriate partitioning schemes for UCE data sets leads to changes in branch order and/or branch lengths. The IQ-Tree runs used default parameters, including linked branch lengths among partitions (option `-spp`) and 100 nonparametric bootstrap replicates to investigate node support. We compared the branching orders of topologies through visual inspection of the trees, and by calculating the PD between pairs of topologies. The PD is a measure of the difference between two topologies, which is similar in principle to the more commonly-used Robinson–Foulds distance, but has more attractive statistical properties such as not giving maximal differences to pairs of topologies that differ by the placement of a single taxon ([Steel and Penny 1993](#)). We compared branch lengths between topologies by plotting the pairs of topologies facing each other, and by calculating and comparing the total tree lengths (i.e., the sum of all branch lengths) of topologies estimated from the same data using different models. The latter approach may reveal any systematic differences of the methods to over- or underestimate branch lengths relative to each other.

Supplementary Material

Supplementary data are available at *Molecular Biology and Evolution* online.

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