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Skmer: assembly-free and alignment-free sample identification using genome skims

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Abstract
The ability to inexpensively describe taxonomic diversity is critical in this era of rapid climate and biodiversity changes. The recent genome-skimming approach extends current barcoding practices beyond short markers by applying low-pass sequencing and recovering whole organelle genomes computationally. This approach discards the nuclear DNA, which constitutes the vast majority of the data. In contrast, we suggest using all unassembled reads. We introduce an assembly-free and alignment-free tool, Skmer, to compute genomic distances between the query and reference genome skims. Skmer shows excellent accuracy in estimating distances and identifying the closest match in reference datasets.

Keywords: Assembly-free, Alignment-free, DNA Barcoding, Genome skimming, DNA reference database, Second generation sequencing

Background
The ability to quickly and inexpensively study the taxonomic diversity in an environment is critical in this era of rapid climate and biodiversity changes. The current molecular technique of choice is (meta)barcoding [1–3]. Traditional (meta)barcoding is based on DNA sequencing of taxonomically informative and group-specific marker genes (e.g., mitochondrial COI [1, 4] and 12S/16S [5, 6] for animals, plastid genes like matK for plants [7], and ITS [8] for fungi) that are variable enough for taxonomic identification, but have flanking regions that are sufficiently conserved to allow for PCR amplification using universal primers. Barcoding is used for taxonomic identification of single-species samples. In the case of metabarcoding, the goal is to deconstruct the taxonomic composition of a mixed sample consisting of multiple species [3]. Beyond the barcoding application, the barcoding marker genes have also been used to delimitate species [9] and to infer phylogenies [10, 11].

The accuracy of (meta)barcoding depends on the coverage of the reference database and the method used to search queries against it [3]. To increase coverage, reference databases with millions of barcodes have been generated (e.g., Barcode of Life Data System, BOLD, for COI [12]). Computational methods for finding the closest match in a reference dataset (e.g., TaxI [13]), and for placement of a query into existing marker trees [14–16] have been developed. However, the traditional approach to (meta)barcoding, despite its success, has some drawbacks. PCR for marker gene amplification requires relatively high-quality DNA and thus cannot be applied to samples in which the DNA is heavily fragmented. Moreover, since barcode markers are relatively short regions, their phylogenetic signal and identification resolution can be limited [17]. For example, in a recent study, 896 out of 4,174 wasp species could not be distinguished from each other using COI barcodes [18].

While low costs have kept PCR-based pipelines attractive, decreasing costs of shotgun sequencing have now made it possible to shotgun sequence 1–2 Gb of total DNA per reference specimen sample for as low as $80 [19], even after including sample preparation and labor costs. This has lead researchers to propose an alternate method that uses low-pass sequencing to generate genome skims [19, 20], and subsequently identifies plastid or mitochondrial marker genes or assembles the organelle genome. Reconstructing plastid and mtDNA genomes from low-pass shotgun data is possible because...
organellar DNA tends to be heavily overrepresented in shotgun sequencing data; for example, 10.4% of all reads from the Apocynaceae family of flowering plants were from the chloroplast in one genome-skimming study [20]. Large reference databases based on genome-skimming techniques are under construction by projects such as PhyloAlps [21], NorBol [22], and DNAmark [23].

Most current applications of genome skimming to species identification require organellar genome assembly, a task that requires relatively time-consuming manual curation steps to ensure that assembly errors are avoided [24]. This approach discards a vast proportion of the non-target data, reducing the discriminatory power. For these reasons, the DNAmark project [23] is considering alternative methods, where, instead of only relying on organellar markers, one could use the entire set of reads generated in a genome skim as the identifier of a species. This approach poses an interesting methodological question: can the unassembled data be used to taxonomically profile reference and query samples in a similar manner to conventional barcoding, but using all available genomic information and saving us from the labor-intensive task of mitochondria/plastid genome assembly? In this paper, we introduce a new assembly-free method to directly use low-coverage genome skims of both reference and query samples. By avoiding the assembly step, our approach also reduces the amount of data processing needed for expanding the reference database.

We treat genome skims simply as low-coverage “bags of reads,” both for a collection of reference species and for query samples. The problem is to find the reference genome skim that matches the query; if an exact match is not found, we seek the closest available match. A more advanced problem, not directly addressed here, is placing the query in a phylogeny of reference species. An even more difficult challenge, also not addressed here, is decomposing a query genome skim that contains DNA from several different taxa into its constituent species.

Central to solving these problems is the ability to estimate a distance between two genome skims for low and varied coverage using assembly-free and alignment-free approaches. Alignment-free sequence comparison has been widely studied [25–30], including for phylogenetic reconstruction [25, 31–44]. Most existing methods, such as Kр [28], spaced words [44], and kmacs [45], compute evolutionary distances using the length distribution of matched substrings or the count of certain words and thus require assembled genomes to produce accurate results. These methods will not work with high accuracy when both the query and the reference are a set of reads and not assembled contigs. Other methods, such as andi [41] and FSWM [43], use micro-alignments to compute distances.

Even though it may be possible to extend the idea of using micro-alignments to the assembly-free case, both andi and FSWM software currently require assemblies as input. However, several assembly-free methods also exist. Co-phylot [39] makes micro-alignments and calculates distances to reconstruct phylogenetic trees; Mash [46] computes the Jaccard index and an evolutionary distance using the k-mers; Simka [47] computes several distance measures based on the whole k-mer content of reads. However, these methods all assume high enough coverage, ensuring that most of the genome is covered. These levels of coverage are currently not economically feasible for building up large reference databases or for obtaining many query samples. Among existing methods, AAF [33] is the only one that aims to work even at lower coverage. AAF first infers a phylogeny and then corrects its branch lengths to reflect a given estimate of the coverage.

Here, we show that high levels of coverage are not necessary. We focus on a distance measure defined as the proportion of mismatches between the global alignment of two genomes. The mismatch rate, called genomic distance hereafter, is useful for species identification because it reflects the evolutionary divergence between two species. We introduce a new method, Skmer, for accurately computing the genomic distance even from low-coverage genome skims. In extensive test, we show that Skmer dramatically improves estimates of genomic distance based on genome skims and accurately places genome-skim queries on to a reference collection. This assembly-free approach can therefore be considered a viable complement to currently available DNA barcoding and genome-skimming tools.

**Results**

**Skmer**

We decomposed reads into fixed-length oligomers (denoted k-mers with length k), a technique used by many existing alignment-free methods [41, 48]. Recall that the *Jaccard index* J is a similarity measure between any two sets (e.g., k-mer collections) defined as the size of their intersection divided by the size of their union. Ondov et al. describe a tool, Mash [46], in which (a) J is estimated efficiently using a hashing procedure and (b) J is used to estimate the genomic distance between two genomes. Mash, however, assumes sufficiently high coverage. Unfortunately, J, in addition to the true distance, is impacted by coverage, sequencing error, and genome length. Skmer accounts for the impact of these factors on J.

Skmer has two stages (Fig. 1): first, we use k-mer frequency profiles (computed using JellyFish [49]) to estimate the amount of sequencing error and the coverage (neither of which is known) using a novel method. Let $M_i$ be the number of k-mers observed i times in the genome skim. Let $h = \arg\max_{i \geq 2} M_i$. Then, defining $\xi = \frac{M_h}{M_h + (k+1)}$,
we derive (see “Estimating sequencing coverage and error rate” section):

$$\lambda = \frac{M_1}{M_h} \frac{\xi h}{h!} e^{-\xi} + \xi (1 - e^{-\xi})$$

(1)

$$\epsilon = 1 - \left(\frac{\xi}{\lambda}\right)^{1/k}$$

(2)

where $\lambda$ and $\epsilon$ are our estimates of the $k$-mer coverage and the sequencing error rate, respectively.

In stage two, we use the hashing technique of Mash to compute $J$. Finally, given these estimates, we compute the genomic distance using

$$D = 1 - \left(\frac{2(\zeta_1 L_1 + \zeta_2 L_2)f}{\eta_1 \eta_2 (L_1 + L_2)(1 + f)}\right)^{1/k}$$

(3)

where for $i \in \{1, 2\}$, $\eta_i = 1 - e^{-\lambda_i (1 - \epsilon_i)^k}$ and $\zeta_i = \eta_i + \lambda_i (1 - (1 - \epsilon_i)^k)$ (for high coverage, we define $\zeta_i$ and $\eta_i$ differently; see “Sequencing error” section for details), and $L_i$ is the estimated genome length.

We used a series of experiments to study the accuracy of Skmer compared to existing methods with respect to (i) the error in computed distances, (ii) the ability to find the closest match to a query sequence in a reference dataset of genome skims, and (iii) phylogenetic inference. We compared the performance against Mash and AAF [33]. AAF is a method that uses $k$-mers to estimate phylogenetic distances among a set of at least four sequences. We conclude by comparing Skmer against the results of using COI barcodes from available barcode databases.

**Distance accuracy for pairs of genome skims**

We first compare the accuracy of Mash and Skmer in estimating distances between two genome skims. Since AAF outputs a phylogenetic tree and so requires at least four species, we cannot include it in our first set of analyses on pairs of genomes.

**Simulated genomes with controlled distance**

Starting from the highly repetitive genome assembly of the wasp species *Cotesia vestalis*, we simulated new genomes with controlled true distance $d$ by randomly adding SNPs, and then we simulated genome skims by randomly subsampling reads and adding error (see “Genomic datasets” section). On these simulated genomes, distances are computed with high accuracy by Mash when coverage is high (Fig. 2), except where the true distance is also high (i.e., 0.2). However, the accuracy of Mash quickly degrades when the coverage is reduced to $4 \times$ or less. In contrast, even when the coverage is reduced to $\frac{1}{8} \times$, Skmer has high accuracy. For example, with the true distance set to 0.05, Mash estimates the distance as 0.081 with $1 \times$ coverage (an overestimation by 62%) while Skmer corrects the
Comparing the accuracy of Mash and Skmer on simulated genomes. Genome skims are simulated using ART with read length $\ell = 100$. Substitutions applied to the assembly of *C. vestalis* at six different rates ($x$-axis), and genome skims simulated at varying coverage range from $\frac{1}{8}$ to $16\times$. The estimated distance ($y$-axis) by Mash (left) and Skmer (right) is plotted versus the real distances for each coverage level (color). The mean (dots) and standard error (lines) of distances are shown (10 repeats). True distance is shown in red. See Additional file 1: Figure S1 for a scaled representation.

Fig. 2 Comparing the accuracy of Mash and Skmer on simulated genomes. Genome skims are simulated using ART with read length $\ell = 100$. Substitutions applied to the assembly of *C. vestalis* at six different rates ($x$-axis), and genome skims simulated at varying coverage range from $\frac{1}{8}$ to $16\times$. The estimated distance ($y$-axis) by Mash (left) and Skmer (right) is plotted versus the real distances for each coverage level (color). The mean (dots) and standard error (lines) of distances are shown (10 repeats). True distance is shown in red. See Additional file 1: Figure S1 for a scaled representation.

distance to 0.045 (an underestimation by 10%). Note that applying Mash* (Mash without the unnecessary approximation $(1 - D)^k \approx e^{-kD}$ used by default in Mash) to the complete assemblies generally generates very accurate results, as expected, but even given the full assembly, Mash* still has a small but noticeable error when $d = 0.2$. Note that results are extremely consistent across our ten different runs of subsampling (Fig. 2). We repeated the simulation with a lower range of coverage ($\frac{1}{64}$ to $1\times$). Interestingly, even with very low coverage, the absolute distance error is small in many cases (Additional file 1: Figure S2); however, for $d \geq 0.1$, Skmer estimates start to degrade below $\frac{1}{8}\times$ coverage.

Repeating the process with the *Drosophila melanogaster* genome as the base genome also produces similar results (Additional file 1: Figure S3). The only condition where Skmer has an absolute error larger than 0.01 is with coverage below $1\times$ and $d = 0.2$ (Fig. 2). However, we note that for $d = 0.001$, the relative error is not small with low coverage (Additional file 1: Figure S4b) indicating that distinguishing very small distances (perhaps below species level) requires high coverage. Estimating the right order of magnitude when the true distance is 0.001 seems to require $2\times$ coverage (preferably $8\times$) while $1\times$ coverage is sufficient to distinguish distances at or above 0.01 (Additional file 1: Figure S4).

**Pairs of insect and bird genomes**

We now test methods on several pairs of insect and avian genomes, subsampled to create genome skims. Note that unlike the simulated datasets, here, genomes can undergo all types of genetic variations and complex rearrangements, and thus, do not have the same length. We carefully selected several pairs of genomes to cover a wide range of mutation distance and genome length.

Here, the true genomic distance is not known, but we use the distance estimated by Mash* on the full assemblies as the true distance $d$. For all pairs of insect and avian genomes (Fig. 3), Mash has high error for coverage below $8\times$ while Skmer successfully corrects the estimated distance and obtains values extremely close to the results of running Mash* on the full assembly. For example, the distance between *Anopheles stephensi* with length of $\sim196$ Mbp and *Anopheles maculatus* with length of $\sim132$ Mbp is estimated to be 0.104 based on the full assembly and 0.102 (2% underestimation) with only $\frac{1}{2}\times$ coverage using Skmer, while Mash would estimate the distance to be 0.163 ($\sim$57% overestimation).

**Distance accuracy for all pairs genome skims**

We now turn to datasets with sets of genome skims, evaluating the accuracy of all pairs of distances. Here, since we have at least four sequences in each test, in addition to Mash, we also compare our results with AAF.
Fig. 3 Comparing the accuracy of Mash and Skmer on pairs of insects (a) and birds (b) genomes. Genome skims are simulated at coverage 1 × to 8 × (shades of blue). The estimated distance (y-axis) is plotted for Mash (left) and Skmer (right) for each pair of species (x-axis). The results of Mash* run on assemblies, which is taken as the ground truth, is shown in red. Mash overestimates at lower coverages. Skmer estimates are closer to the ground truth and are less sensitive to the coverage. See also Additional file 1: Figure S5.
Fixed sequencing effort

So far, our experiments have controlled for the coverage by subsampling varying amounts of sequence data, proportional to the genome length. In our genome-skimming application, coverage will not be fixed. Often, the amount of sequence data obtained for each species will be relatively similar. As a result, genomes of different length end up being sequenced with different coverage depth proportional to the inverse of their length. We therefore performed a study where all species are subsampled to produce 100 Mb of sequence data in total resulting in varying levels of coverage (based on the genome length, Additional file 1: Table S5). The error in the distance estimated by Mash relative to the ground truth can be quite large (higher than 300% in the worst case) while Skmer consistently makes accurate estimates close to the true distance even at the lowest amount of coverage (Figs. 4 and 5, and Additional file 1: FigureS9). Repeating the analysis with 0.5 Gb or 1 Gb total sequence data produced similar patterns, but as expected, increasing the sequencing effort reduces the error for all methods (Additional file 1: Figures S6-S8).

Before error correction, AAF has error levels that are comparable to Mash (Figs. 4, 5). The correction applied by AAF, similar to Skmer, reduces the negative impact of low coverage but not to the same extent. Thus, Skmer has less error compared to corrected AAF (with 100 Mb sequence and across all datasets, the mean error of Skmer is 3.13% and AAF-corrected is 22.7%). For example, in the Drosophila dataset, the worst-case error of AAF between any two pairs of genome skims is 31%, whereas the error never exceeds 8% for Skmer. Note that when computing the error of AAF, we use the result of running AAF on full assemblies as the ground truth.

To quantify the impact of distance estimates on downstream analyses, we used FastME [50] to infer phylogenetic trees using distances computed by Mash and Skmer on genome skims and with correction using the JC69 model [51]. AAF by default generates trees as part of its output. We compare these trees to those computed by Mash/AAF run on the full assemblies (taken as the ground truth) using the weighted Roubinson-Foulds (WRF) distance [52] (Table 1). WRF is the sum of branch length differences between the two trees (using zero length for missing branches), and we normalized WRF by the sum of branch lengths of both trees. In all three datasets, Skmer distances lead to trees with lower WRF distance to the ground truth compared to Mash and AAF/uncorrected. AAF correction reduces WRF compared to uncorrected AAF; however, Skmer trees have two to 14 times less error compared to the corrected AAF, except in one case where AAF/corrected has 1.05% error and Skmer has 1.19% (Table 1). Increasing the size of skims to 0.5 Gb and 1 Gb helps all methods to produce more accurate trees.

Heterogeneous sequencing effort

In addition to changes in the genomic length, the sequencing effort per species may also vary across sequencing protocols, experiments, and research labs, and so a database of reference genome skims may consist of samples with heterogeneous sequencing efforts. To capture this, for each species, we choose its total sequencing effort from three possible values 0.1 Gb, 0.5 Gb, and 1 Gb, uniformly at random, and estimate all pairs of distances within each dataset as before (Fig. 6 and Additional file 1: Figure S9). Similar to the case of fixed sequencing effort, Skmer mitigates large relative error in the distances estimated by Mash and produces more accurate results than both Mash and AAF (Table 2, Fig. 6, and Additional file 1: Figure S9). For example, comparing to the case of fixed 100-Mb genome skims of the Drosophila dataset, the worst-case error of AAF is increased to 70%, while using Skmer it remains almost the same (8%). Comparing trees inferred from distances estimated by various methods also confirms the higher accuracy of Skmer (Table 1). For instance, on the Anopheles dataset, Skmer has only 0.58% WRF distance to the reference tree whereas Mash and AAF-corrected trees have 14.75% and 8.45% WRF distance.

Genome skims from real reads

Running time

So far, all of our tests used simulated reads. When analyzing real genome skims, there are additional complications such as extraneous DNA (real or artifactual) and the over-representation of organelle genome. We next tested Skmer using real reads. We created 100-Mb skims of 14 Drosophila genomes by subsampling short-read data produced in a recent Drosophila genome assembly study [53]. Before running Skmer or Mash, we filtered reads that (even partially) aligned to 12 Drosophila-associated microbial genomes as reported in previous studies [54–56] (see Additional file 1: Table S1), to the human genome, or to the mitochondrial genome of respective Drosophila species. We then estimated all pairs of distances as before and computed the error relative to the distances computed from the assemblies (Fig. 7). Consistent with the results that, we obtained on the simulated skims, Skmer has less error compared to Mash. The average error of Mash on this dataset is 43.48% (±2.29%) with maximum error of 217%. Skmer, on the other hand, has an average error of 4.21% (±0.35%) and its maximum error is 22.2%.

Skmer and Mash have comparable running time, while AAF is much slower. In the experiment with heterogeneous sequencing effort, the total running time (using 24 CPU cores) to compute distances based on genome skims for all \( \binom{47}{2} \) pairs of birds using Mash, Skmer, and AAF was roughly 8, 33, and 460 min, respectively.
a

![Diagram showing distance error with fixed 100 Mb sequence per genome for Anopheles and Drosophila. Each genome is skinned with 100 Mb sequence and distances are computed using Mash, Skmer, and AAF. True distance used in calculating the error is computed by applying each method (AAF and Mash) to the full genome assemblies. The heatmaps on the left show the error of Mash (upper triangle) and Skmer (lower triangle), and the heatmaps on the right are for AAF before correction (upper) and after correction (lower).]

b

![Diagram showing distance error with fixed 100 Mb sequence per genome for Anopheles and Drosophila. Each genome is skinned with 100 Mb sequence and distances are computed using Mash, Skmer, and AAF. True distance used in calculating the error is computed by applying each method (AAF and Mash) to the full genome assemblies. The heatmaps on the left show the error of Mash (upper triangle) and Skmer (lower triangle), and the heatmaps on the right are for AAF before correction (upper) and after correction (lower).]

Fig. 4 Distance error with fixed 100 Mb sequence per genome for a 22 Anopheles and b 21 Drosophila. Each genome is skinned with 100 Mb sequence and distances are computed using Mash, Skmer, and AAF. True distance used in calculating the error is computed by applying each method (AAF and Mash) to the full genome assemblies. The heatmaps on the left show the error of Mash (upper triangle) and Skmer (lower triangle), and the heatmaps on the right are for AAF before correction (upper) and after correction (lower).

**Leave-out search against a reference database of genome skims**

We now study the effectiveness of using genomic distance to search a database of genome skims to find the closest match to a query genome skim. Given a query genome skim and a reference dataset of genomes, we can order the reference genomes based on their distance to
Fig. 5 Distance error with fixed 100 Mb sequence per genome for the avian dataset. The errors of Mash and AAF for the two eagle species (H. albicilla and H. leucocephalus) were extremely large (Mash: \(\approx 4000\%\), AAF > 3000\% error), dominating the color spectrum; we excluded H. albicilla to help readability; for the eagles, Skmer’s estimate is 0.00244 (\(\sim 9\%\) error).
Phylogeny reconstruction and comparison to organelle markers

5.64 Birds 0.1G 37.03%
2.05 Drosophila 0.1G 23.87%

As the last experiment, we estimated phylogenetic trees as described in “Leave-out” section. Figure 8 shows the effectiveness of the search as the distance of the closest available dataset, finding the match is relatively easy. To study the ranking. When the query genome is available in the reference library, the query can be provided to the user as a result. The results can be provided to the user as a perfect match to the best available genome.

On all three datasets, Skmer consistently and often substantially outperforms Mash and AAF in terms of finding the best remaining match, except the Drosophila dataset where Mash and Skmer have comparable rank error, while both are better than AAF (Fig. 8). Even in that case, on average, the distance of the best match found by Skmer is closer to the distance of the true best match compared to the best hit found by Mash. Moreover, the mean rank error of Skmer is smaller than Mash (Additional file 1: Figure S10) if we exclude only one species Drosophila willistoni (which is at distance 0.1565 ≤ d ≤ 0.1622 from other species). It is also notable that over the avian dataset, Skmer has mean rank error less than 0.5 for all range of distances, while Mash and AAF can be off by more than 2.5 on average. These results demonstrate that correcting the distance not only impacts our understanding of the absolute distance, but also impacts results of searching a reference library.

**Phylogeny reconstruction and comparison to organelle markers**

As the last experiment, we estimated phylogenetic trees for Anopheles and Drosophila datasets after transforming the genomic distances estimated by Skmer to Jukes-Cantor (JC) distances [51]. For each dataset, we also built another tree based on available COI barcodes, using an identical method. We compare the results against a reference tree obtained from Open Tree of Life [57]. We restricted the results to species for which COI barcodes were available (Fig. 9ab).

For the Anopheles species, Skmer produces a tree that is almost identical to the reference tree (with only one branch difference out of nine), while COI tree differs from the reference in seven branches. Similarly, for the Drosophila species, Skmer differs from the reference in three branches (with small local changes) out of 13 total branches in the reference tree, whereas COI tree is very inconsistent with the reference tree (seven branches are different). We also built maximum-likelihood trees from COI barcodes (Additional file 1: Figure S11), but the number of incorrect branches did not reduce. Comparing the distribution of all pairwise genomic distances obtained from genome skims and barcodes (Fig. 9c), Skmer has larger distances and fewer pairs with zero or close to zero distance, indicating that Skmer has a higher resolution in differentiating between samples. For example, four species of the Anopheles genus A. coluzzii, A. gambiae, A. arabiensis, and A. melas have very small pairwise distances based on COI barcodes, while using Skmer, the estimated distances are in the range 0.02–0.04 for these species.

### Discussion

We showed that Skmer can compute the genomic distance between a pair of species from genome skims with very low coverage (at or even below 1 ×), with much better accuracy than the main two alternatives, Mash and AAF. We also showed that the distances computed by Skmer can accurately place a voucher genome skim within a reference database of genome skims, and can be used to infer the phylogenetic tree with reasonable accuracy. While Skmer is not the first k-mer-based approach for distance estimation or phylogenetic reconstruction, as we showed, the alternatives have low accuracy given low-coverage data. We compare with Mash because it is used within Skmer and is one of the most widely used alignment-free methods. However, we note that authors of Mash do not claim it can handle low coverage, and so our results are not a criticism of their approach. Besides the methods we discussed, many other alignment-free sequence comparison and phylogeny reconstruction algorithms exist [25, 28, 29, 31, 32, 34–43]. However, these methods take as input assembled (but unaligned) sequences, and thus, are not applicable in an assembly-free pipeline. In other words, their goal is to avoid the alignment step and not the assembly step.

Compared to using COI markers, currently used in practice, we showed that using all k-mers, including those

| Dataset     | Sequencing effort | Mash (corrected) | Skmer (corrected) | AAF (uncorrected) | AAF (corrected) |
|-------------|-------------------|------------------|-------------------|-------------------|-----------------|
| Anopheles   | 0.1 G             | 23.19%           | 1.07%             | 19.92%            | 6.30%           |
|             | 0.5 G             | 12.84%           | 0.45%             | 9.74%             | 4.9%            |
|             | 1 G               | 8.92%            | 0.37%             | 9.59%             | 3.3%            |
|             | Mixed             | 14.75%           | 0.58%             | 8.46%             | 8.45%           |
| Drosophila  | 0.1 G             | 23.87%           | 2.05%             | 20.29%            | 5.85%           |
|             | 0.5 G             | 13.33%           | 0.72%             | 10.37%            | 5.25%           |
|             | 1 G               | 7.11%            | 0.58%             | 10.84%            | 2.2%            |
|             | Mixed             | 16.58%           | 1.11%             | 11.36%            | 10.87%          |
| Birds       | 0.1 G             | 37.03%           | 5.64%             | 31.81%            | 21.13%          |
|             | 0.5 G             | 25.16%           | 1.91%             | 20.88%            | 6.86%           |
|             | 1 G               | 19.42%           | 1.19%             | 15.54%            | 1.05%           |
|             | Mixed             | 28.14%           | 3.08%             | 18.15%            | 7.57%           |

For each method, we show normalized weighted RF distance (%) of trees inferred from genome-skim distances to trees inferred from full-assembly distances. Italics: the lowest error.
from the nuclear genome, improves the phylogenetic accuracy. These improvements are resulting from distances that have a larger range and more resolution compared to COI. Also, the increased resolution should not be surprising given that the entire genome is much larger than any single locus, reducing the variance in estimates of the distance. Beyond the question of resolution, gene trees and species trees need not match [58], a fact

**Fig. 6** Distance error with heterogeneous sequencing effort for Anopheles and Drosophila. Species have random amount of sequence chosen uniformly among 0.1 Gb, 0.5 Gb, and 1 Gb. See Additional file 1: Figure S9 for birds.
Table 2 Comparing the average error of Mash, Skmer, and AAF in estimating distances over three datasets with heterogeneous sequencing effort

| Dataset | Mash (uncorrected) | Skmer (uncorrected) | AAF (corrected) | AAF (corrected) |
|---------|--------------------|---------------------|-----------------|-----------------|
| Anopheles | 28.72% (1.10%) | 0.84% (0.03%) | 13.48% (0.56%) | 11.36% (0.44%) |
| Drosophila | 29.05% (0.59%) | 0.84% (0.04%) | 15.25% (0.38%) | 10.94% (0.33%) |
| Birds | 64.29% (0.54%) | 2.21% (0.04%) | 36.02% (0.29%) | 5.28% (0.16%) |

The standard error of the mean is provided in parentheses. Italics: the lowest error

that can further reduce the accuracy of marker genes for both species identification and phylogeny reconstruction. By using the entire genome, Skmer ensures that an average distance across the genome is computed, reducing the sensitivity to gene tree/species tree discordances. Moreover, a recent result shows that the JC-transformed genomic distance is a statistically consistent estimator of the species distances despite gene tree discordance due to incomplete lineage sorting [59], further encouraging our use of the genomic distance as a measure of the evolutionary divergence.

We showed that genomic distances as small as 0.01 can be estimated accurately from genome skims with 1× or lower coverage. What does a distance of 0.01 mean? The answer will depend on the organisms of interest. For example, two eagle species of the same genus (Haliaeetus albicilla and Haliaeetus leucocephalus) have $D \approx 0.003$ but two Anopheles species of the same species complex (A. gambiae and A. coluzzii) have $D \approx 0.018$. Broadly speaking, for eukaryotes, detecting distances in the $10^{-2}$ order is often enough to distinguish between species (Additional file 1: Figure S12). On the other hand, to differentiate individuals in a population, or very similar species, we may need to reliably estimate distances of the order $10^{-3}$. Detection at these lower levels seems to require $> 1 \times$ coverage using Skmer (Additional file 1: Figure S4b) but future work should study the exact level of sequencing required for accurate ordering of species at distances in the order of $10^{-3}$ or less. Moreover, the question of the minimum coverage required may avail itself to information-theoretical bounds and near-optimal solutions, similar to those established for the assembly problem [60, 61].

Although most of our tests were performed on genome skims simulated from assemblies, we also tested Skmer on genome skims simulated by subsampling previous whole-genome sequencing experiments. Several complications have to be addressed in real applications. The actual coverage of real genome skims may not be uniform

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Fig. 7 Comparing the error of Mash and Skmer on a dataset of 14 Drosophila genome skims. Each SRA is subsampled to 100 Mb and then filtered to remove contamination. True distances are computed from the assemblies.
The mean rank and distance error of the best remaining match in leave-out experiments. The distance of the closest genome in the reference to a query is varied from 0.01 to 0.1 (x-axis). The rank and distance errors (y-axis) of the best match to a query, are computed by comparing the order given by each method with the order obtained by applying Mash* to the full assemblies (ground truth). For each dataset, the experiment is repeated by taking each species as the query, and then the errors are averaged. Three methods, Mash, Skmer, and AAF, are compared on a the Anopheles dataset, b the Drosophila dataset, and c the avian dataset.
and randomly distributed and they can have an overrepresentation of mitochondrial or plastid sequence. More importantly, other sources of DNA originating from, for example, parasites, diet, fungi, commensals, bacteria, and human contamination may all be present in the sample and may cause a bias in the estimation of distances. In our test, we simply searched all reads in a genome skim against a few bacterial genomes and the human reference genome; this simple scheme filtered out up to ~10% of reads (for *D. virilis*). These filtering strategies were
sufficient to produce reliable distance estimates in the case of Drosophila genomes. We recommend that before using Skmer, such database searches should be used to find and eliminate bacterial or fungal contamination (using BLAST [62] or perhaps metagenomic tools such as Kraken [63]), as well as removing contaminant reads with human origin (using for example Bowtie2 [64]). However, in future, it will be beneficial to develop better methods for finding extraneous reads without reliance on known sources.

A related direction of future work is to explore whether Skmer can be extended to environmental DNA analyses, i.e., queries consisting of genome skims of multi-taxa samples. While Skmer is presented here in a general setting, its best use is for eukaryotic organisms, where the notion of species is better established and species can be separated with reasonable effort. We tested Skmer on birds and insects, but we predict it will work equally well for plants, a prediction that we plan to test in future work.

Throughout our experiments, we used Mash* run on the assemblies to compute the ground truth. Given the true alignment of the two genomes, we can compute the true genomic distance as the proportion of mismatches among *aligned* orthologous positions (i.e., ignoring gaps). To ensure that Mash* closely approximates true distances, we used simulated genomes of Rat and Mouse from the Mammalian dataset of the Alignathon competition [65]. This simulation uses Evolver [66] and includes many forms of mutation, including indels, rearrangements, duplications, and losses. On this dataset, the true distance based on the known true alignment is 0.145 and Mash* estimated the distance as 0.143, which is a very good approximation. In contrast, FastANI [67], an alignment-free sequence mapping tool for estimating average nucleotide identity, computes the distance as 0.189. If we count gaps as non-matching positions in the definition of distance, then the true distance would be 0.287, which also does not match FastANI. Presumably, FastANI, which relies on alignment of short blocks, counts short gaps (with *some* definition of short) as mismatch but excludes larger ones. Thus, on real data, Mash* is the best available option to approximate the true distance. Finally, note that, for real genomes, we chose not to use estimated whole genome alignments (WGA) to compute the ground truth because WGA is a difficult problem, and WGAs that are available are not necessarily accurate. We get inconsistent estimates of distance when we use pairwise or multiple WGAs. For example, between *D. melanogaster* and *D. yakuba*, the distance changes from 0.10 when using the multiple WGA [68], to 0.21 if we use the pairwise WGAs [69] from the UCSC genome browser [70], which is the state of the art.

The connection between genomic distance and phylogenetic distance depends on mutation processes considered. If only substitutions are allowed and assuming the Jukes-Cantor model, the phylogenetic distance is $-\frac{3}{4} \ln (1 - \frac{3}{4}d)$; note this transformation is monotonic and does not change rankings of matches to a query search. Assuming a more complex model such as GTR [71], genomic distance is not enough to estimate the phylogenetic distance. However, we have devised a simple procedure to estimate GTR distances using the log-det approach [72] by repeated applications of Skmer to perturbed reads (Additional file 1: Appendix B). The GTR distances can rank matches to a query differently from the genomic distance; the accuracy of the two distances should be compared in future work.

Insertions, deletions, duplications, and losses can all lead to differences between genomes, thereby reducing the Jaccard index and increasing the genomic distance. They also impact genomic length. Interestingly, in our experiments, Skmer run with the true coverage is less accurate than with estimated coverage (Additional file 1: Figure S13). We speculate that on genomes with repeats, by overestimating coverage, our method gives an estimate of the "effective" coverage, reducing the impact of repeats on the Jaccard index. Nevertheless, with these complex mutations, the correct definitions of the evolutionary distance and genomic distance are not straightforward, nor is it clear how the Jaccard index should be translated to the genomic distance. Here, we used a heuristic approach that simply averaged the length of the two genomes, leaving these broader questions about the best definition of genomic distance in the presence of large structural variations to future work.

Conclusions

Skmer is an assembly-free and alignment-free tool for estimating the distance between two genome skims. It can estimate a wide range of distances with high accuracy from low-coverage and mixed-coverage genome skims with no prior knowledge of the coverage or the sequencing error. Our paper shows that the idea of genome-wide sample identification using genome skims has merit and should be pursued in the future.

Methods

Consider an idealized model where two genomes are the outcome of a random process that copies a genome and introduces mutations at each position with fixed probability $d$. Moreover, substitutions are the only allowed mutation. In this case, the per-nucleotide hamming distance $D$ between the two genomes is a random variable (r.v.) with expected value $d$. We would like to estimate $d$. While this is a simplified model, we will test the method on real pairs of genomes that differ due to complex mutational processes (also, see Additional file 1: Appendix B for extensions). We start with known results connecting the Jaccard index and the hamming distance and then show how these results can be generalized to low-coverage genome skims.
Throughout, we present our results succinctly and present derivations and more careful justifications in Additional file 1: Appendix A of the supplementary material.

**Jaccard index versus genomic distance**

The Jaccard index of subsets $A_1$ and $A_2$ is defined as

$$J = \frac{|A_1 \cap A_2|}{|A_1 \cup A_2|} = \frac{|A_1 \cap A_2|}{|A_1| + |A_2| - |A_1 \cap A_2|}. \quad (4)$$

Let $W$ be the number of shared $k$-mers between the two genomes. Note that $J = \frac{W}{W + W} = \frac{2}{2+1} = \frac{2}{3}$, where $L$ is the genome length. Assuming random genomes and no repeats, perhaps justifiably [73], the probability that a changed $k$-mer exists elsewhere in the genome is vanishingly small for sufficiently large $k$. Thus, we assume a $k$-mer is in the shared $k$-mers set only if no mutation falls on it, an event that has probability $(1 - d)^k$. Thus, we can model $W$ as a binomial with probability $(1 - d)^k$ and $L$ trials. As Ondov et al. [46] pointed out, we can estimate

$$D = 1 - \left( \frac{2J}{J + 1} \right)^{\frac{1}{2}} \quad (5)$$

and they further approximate $D$ as $\frac{1}{2} \ln \left( \frac{J - 1}{J} \right)$. To be able to estimate large distances, we avoid the unnecessary approximation and use Eq. 5 directly. We skim each genome to obtain $k$-mer sets $A_1, A_2$ and estimate $J$ using Eq. 4, which can be computed efficiently using a hashing technique used by Mash [46]. Note that, however, Eq. 5 assumes a high coverage of the genome so that each $k$-mer is sampled at least once with very high probability. This assumption is violated for genome skims in consequential ways. As a simple example, suppose the coverage is low enough that a $k$-mer is sampled with probability 0.5. Then, even for identical genomes, we estimate $J$ as $\frac{1}{2}$, resulting in a distance estimate of $D \approx 0.032$ for $k = 21$.

**Extending to genome skims with known low coverage and error**

We now show how Eq. 5 can be refined to handle genome skims despite low and uneven coverage, sequencing error, and varying genome lengths. We first assume that coverage and error are known and later show how to compute these.

**Low coverage**

When the genome is not fully covered, three sources of randomness are at work: mutations and sampling of $k$-mers from each of the two genomes. Each genome of length $L$ is sequenced independently using randomly distributed short reads of length $\ell$ at coverages $c_1$ and $c_2$ to produce two genome skims. Under the simplifying assumption that genomes are not repetitive, we choose $k$ to be large enough so that each $k$-mer is unique with high probability. Therefore, the number of distinct $k$-mers in each genome is $L - k \approx L$. The probability of covering each $k$-mer can be approximated as $\eta_i = 1 - e^{-\lambda_i}$ where $\lambda_i = c_i (1 - k/\ell)$. Modeling the sampling of $k$-mers as independent Bernoulli trials, $|A_i|$ becomes binomially distributed with parameters $\eta_i$ and $L$. By independence, $W = |A_1 \cap A_2|$ also becomes binomially distributed with parameters $\eta_1 \eta_2 (1 - d)^k$ and $L$. Moreover, $U = |A_1 \cup A_2|$ can also be modeled approximately as a Gaussian with mean $(\eta_1 + \eta_2 - \eta_1 \eta_2 (1 - d)^k) L$. Treating $\eta_1$ and $\eta_2$ as known and dividing $\frac{W}{U}$ by $\frac{U}{T}$ gives us:

$$J = \frac{W}{U} = \frac{\eta_1 \eta_2 (1 - D)^k}{\eta_1 + \eta_2 - \eta_1 \eta_2 (1 - D)^k};$$

thus,

$$D = 1 - \left( \frac{\eta_1 + \eta_2}{\eta_1 \eta_2} \right)^{\frac{1}{2}}. \quad (6)$$

**Sequencing error**

Each error reduces the number of shared $k$-mers and increases the total number of observed $k$-mers, and thus can also change the Jaccard index. Let $e_i$ denote the base-miscall rate for genome $i$. For large $k$ and small $e_i$, the probability that an erroneous $k$-mer produces a non-novel $k$-mer is negligible. The probability that a $k$-mer is covered by at least one read, without any error, is approximately

$$\eta_i = 1 - e^{-\lambda_i (1 - e_i) k}. \quad (7)$$

Adding up the number of error-free and erroneous $k$-mers, the total number of $k$-mers observed from both genomes can again be approximately modeled as a Gaussian with mean $\tilde{\xi}_L$ for

$$\tilde{\xi}_i = \eta_i + \lambda_i (1 - (1 - e_i) k). \quad (8)$$

Just as before, we can simply estimate $D$ by solving for it in

$$J = \frac{\eta_1 \eta_2 (1 - D)^k}{\tilde{\xi}_1 + \tilde{\xi}_2 - \eta_1 \eta_2 (1 - D)^k}. \quad (9)$$

When the coverage is sufficiently high, each $k$-mer will be covered by multiple reads with high probability, and low-abundance $k$-mers can be safely considered as erroneous. Mash has an option to filter out $k$-mers with abundances less than some threshold $m$ to remove $k$-mers that are likely to be erroneous. In this case,

$$\tilde{\xi}_i = \eta_i = 1 - \sum_{t=0}^{m-1} \frac{(\lambda_i (1 - e_i) k)^t}{t!} e^{-\lambda_i (1 - e_i) k} \quad (9)$$

assuming all erroneous $k$-mers are removed. For instance, filtering single-copy $k$-mers (i.e., $m = 2$) gives us:

$$\tilde{\xi}_i = \eta_i = 1 - e^{-\lambda_i (1 - e_i) k} - \lambda_i (1 - e_i) k e^{-\lambda_i (1 - e_i) k}$$

and the Jaccard index follows the same equation as (8). Since this filtering approach only works for high coverage,
we filter low-coverage k-mers only when our estimated coverage is higher than a threshold (described below). Note that the genome skims compared may use different filtering schemes, yet Eq. 8 holds regardless.

**Differing genome lengths**

Based on a model where the genomic distance between genomes of different lengths is defined to be confined to the mutations that are falling on homologous sequences, we can drive

\[
J = \frac{\eta_1 \eta_2 \min(L_1, L_2)(1 - D)^k}{\xi_1 L_1 + \xi_2 L_2 - \eta_1 \eta_2 \min(L_1, L_2)(1 - D)^k}.
\]

This computation does not penalize for genome length difference. While a rigorous modeling of evolutionary distance for genomes of different length requires sophisticated models of gene gain, duplication, and loss, we take the heuristic approach used by Ondov et al. [46] and simply replace \(\min(L_1, L_2)\) with \((L_1 + L_2)/2\). This ensures that the estimated distance increases as genome lengths becomes successively more different. This leads us to our final estimate of distance given by:

\[
D = 1 - \left(\frac{2(\xi_1 L_1 + \xi_2 L_2)}{\eta_1 \eta_2 (L_1 + L_2)(1 + f)}\right)^{1/k}.
\]  

(10)

**Estimating sequencing coverage and error rate**

So far we have assumed a perfect knowledge of sequencing depth and error. However, for genome skims, the genome length is not known; thus, we need to estimate the coverage in order to apply our distance correction. We also assume a constant base error rate, and co-estimate it with the coverage.

The sequencing depth, which is the average number of reads covering a position in the genome, can be estimated from the k-mer coverage profiles. The probability distribution of the number of reads covering a k-mer is a Poisson r.v. with mean \(\lambda\), where \(\lambda\) is defined as k-mer coverage. As we look into the histogram data, it is easier to work with counts instead of probabilities. Let \(M_i\) denote the total number of k-mers of length \(k\) in the genome, and \(M_i\) count the number of k-mers covered by \(i\) reads. Thus, for \(i \geq 0\), \(E[M_i] = \frac{M_i \xi^i}{\pi} e^{-\lambda}\). For a given set of reads, we can count the number of times that each k-mer is seen, and assuming zero sequencing error, it equals the number of reads covering that k-mer. Then, we can aggregate the number of k-mers covered by \(i\) reads and find \(M_i\) for \(i \geq 1\). However, since in a genome skim, large parts of the genome may not be covered, both \(M\) and \(M_0\) are unknown. To deal with this issue, we could take the ratio of consecutive counts to get a series of estimates of \(\lambda\) as \(\tilde{\lambda}_i = \frac{M_{i+1}}{M_i} (i + 1)\) for \(i = 1, 2, \ldots\). In practice, sequencing errors change the frequency of k-mers and has to be considered when estimating the coverage. Assuming that the error is introduced at a constant rate along the reads, we can use the information in the k-mer counts to co-estimate \(\epsilon\) and \(\lambda\). Like before, we assume that the k-mer length \(k\) is large enough that any error will introduce a novel k-mer, so the count of all erroneous k-mers is added to the count of single-copy k-mers. Moreover, for k-mers with more than one copy, the number of times that each k-mer is seen equals the number of reads covering that k-mer without any error. Formally, let \(\hat{M}_i\) denote the count of k-mers seen \(i\) times in the presence of error, and \(\rho = (1 - \epsilon)^k\) denote the probability of error-free k-mer.

\[
\mathbb{E}[\hat{M}_i] = \begin{cases} \sum_{j \geq 1} M_j \frac{\rho^j}{\rho} e^{-\lambda(j + 1)} & i \geq 2 \\ \sum_{j \geq 1} M_j \frac{\rho^j}{\rho} e^{-\lambda(j + 1)} & i = 1 \\ M_j \frac{\rho^j}{\rho} e^{-\lambda} & i \geq 2 \\ M (\xi e^{-\xi} + \lambda - \xi) & i = 1 \end{cases}
\]

(11)

where \(\xi = \lambda \rho\) is the average number of error-free reads covering a k-mer. A family of estimates for \(\xi\) is obtained by taking the ratio of consecutive counts of error-free k-mers as \(\tilde{\xi}_i = \frac{\hat{M}_{i+1}}{\hat{M}_i} (i + 1)\) for \(i \geq 2\). Then, using an estimate of \(\hat{\xi}\) and the count of single-copy k-mers, we get a series of estimates of \(\lambda\) for \(i \geq 2\) as

\[
\tilde{\xi}_i = \frac{\hat{M}_1}{\hat{M}_i} \frac{\xi^i}{i!} e^{-\tilde{\xi}} + \tilde{\xi} \left(1 - e^{-\tilde{\xi}}\right).
\]

(12)

Moreover, we can estimate the error rate from the estimates of \(\lambda\) and \(\tilde{\xi}\) as

\[
\tilde{\epsilon} = 1 - \left(\frac{\tilde{\xi}}{\tilde{\lambda}}\right)^{1/k}.
\]

(13)

While any of these \(\tilde{\xi}_i\) and \(\tilde{\lambda}_i\) can be used in principle, the empirical performance can be affected by the choice; in our tool, we use heuristic rules (described below) that seek to use large \(M_i\) values.

**Skmer: implementation**

Skmer takes as input two or more genome skims. It uses JellyFish [49] to compute \(M_i\) values, which are then used in estimating \(\lambda\) and \(\epsilon\) based on Eqs. 12 and 13, by setting \(\xi = \tilde{\xi}_h\) and \(\lambda = \tilde{\lambda}_h\), where \(h = \arg\max_{i \geq 2} \tilde{M}_i\). Then, Mash is used to estimate the Jaccard index, with \(k = 31\) (selected empirically; Additional file 1: Figure S14) and sketch size \(10^7\). Finally, we use Eq. 10 to compute the hamming distance with \(\eta\) and \(\zeta\) values computed using Eqs. 6, 7 if \(c < 5\) or else using Eq. 9. The genome length \(L\) is estimated as the total sequence length divided by the coverage \(c\).
Experimental setup

Method settings

For Skmer, we use the default parameters described above. For Mash, similar to Skmer, we used \( k = 31 \) (selected empirically; Additional file 1: Figure S14) and sketch size \( 10^7 \). As Mash handles errors by removing low copy \( k \)-mers, we set the minimum cardinality for \( k \)-mers to be included as \( \lfloor \frac{5}{k} \rfloor + 1 \) with our estimate of \( c \).

AFF has an algorithm to correct hamming distances for low coverage, but the correction relies on adjusting the length of tip branches in a distance-based inferred phylogeny. As such, it cannot run on a pair of genomes and requires at least four genomes. Also, AAF leaves coverage estimation to the user with some guidelines, which we fully follow (Additional file 1: Appendix C).

For building phylogenetic trees, we transformed Skmer distances using the JC69 [51] model and used FastME [50] to construct the distance-based trees via BIONJ [74] method.

Genomic datasets

We used an assembly of *Cotesia vestalis* (GenBank accession: GCA_000956155.1) as well as three sets of publicly available assembled genomes (Additional file 1: Tables S2-S4) and used ART [75] to simulate genome skims of read length \( \ell = 100 \) with default sequencing error profile, controlling for the sequencing depth (coverage) (Additional file 1: Appendix C). Specifically, the data included 21 *Drosophila* genomes (flies) and 22 genomes from the *Anopheles* genus (mosquitoes) obtained from InsectBase [76], and 47 avian species from the Avian Phylogenomic Project [77, 78].

For the experiment on real genome skims, high-coverage SRA’s of 14 *Drosophila* species were obtained from NCBI database under project number PRJNA427774 [79] and then subsampled to 100 Mb. Assemblies used to compute true distances for these 14 *Drosophila* species were obtained from the Drosophila project [80]. We used the tool fastp [81] for filtering low-quality reads and adapter removal. We also used Megablast [82] to search against a database of bacterial and mitochondrial genomes and remove contaminant reads. We used Bowtie2 [64] with the highest sensitivity to remove the reads aligning (even partially) to the human reference genome.

To simulate genomes with controlled genomic distance, we introduced random mutations. As a challenging case, we took the highly repetitive assembly of the wasp species *Cotesia vestalis*, and mutated it artificially; we only applied single nucleotide mutations distributed uniformly at random across the genome. We repeated the study on the simpler case of the fly species *D. melanogaster*. We generate genome skims using ART with \( \ell = 100 \), default error profile of Illumina sequencer, and varying coverage between \( \frac{1}{64} \times \) and \( 16 \times \). For simulated genomes, we repeated the subsampling 10 times and reported the mean and standard error.

In order to compare with DNA barcoding method, we downloaded available COI barcodes for the *Drosophila* and *Anopheles* species in the BOLD database [12]. Out of 21 *Drosophila* and 22 *Anopheles* species in our dataset, 16 *Drosophila* and 19 *Anopheles* species had one or more barcodes in BOLD. For each species, we selected a barcode, and using MUSCLE [83], aligned all barcodes within each dataset and constructed the phylogenetic tree assuming the Jukes-Cantor model. Under the same model of substitution, we transformed Skmer distances and built the Skmer tree. We used FastME [50] to construct the distance-based trees via the BIONJ [74] method. The maximum-likelihood COI trees were built using PhyML [84].

Evaluation metrics

For simulated data, the true distance is controlled and is thus known. For biological datasets, the ground truth is unknown. Instead, we use the distance measured on the full assembly by each method as its ground truth; thus, the ground truth for AAF is computed using AAF. We show both absolute error and the relative error, measured as \( \frac{|d - \hat{d}|}{d} \) where \( d \) and \( \hat{d} \) are the true and the estimated distances.

Leave-out

We used a leave-out strategy to study the accuracy of searching for a query genome in a reference set. For a query genome \( G_q \) in a set of \( n \) genomes \( \{G_1 \ldots G_n\} \), we ordered all genomes based on their distances to \( G_q \) calculated using the full assemblies, which represents the ground truth; let \( G_1 \ldots G_n \) denote the order, and \( d_1 \ldots d_n \) be the respective distances from the query (note \( G_1 = G_q \) and \( d_1 = 0 \)). For \( 0.01 \leq d \leq 0.10 \), we removed genomes \( 1 \ldots i \) from the datasets where \( i \) is the largest value such that \( d_i \leq d \), leaving us with \( G_1^{i+1} \ldots G_n^{i+1} \). We then ordered the remaining genomes by each method; let \( x_1 \ldots x_{n-i} \) be the order obtained by a method and let \( r \) be the rank of the best remaining genome according to the ground truth in the estimated order (i.e., \( x_1 = G_q^{i+r} \)). Since \( r = 1 \) implies perfect performance, and \( r > 1 \) indicates error, we measured rank error as the mean of \( r - 1 \) across all query genomes (1 \( \leq q \leq n \)). Moreover, the mean (relative) distance error is defined as the mean of \( \frac{d_{q}^{i+r} - d_{q}^{i+1}}{d_{q}^{i+1}} \) over all queries.

Additional file

Additional file 1: Supplementary material (PDF 1209 kb)
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