

Heuristics based on Adjacency Graph Packing for DCJ Distance Considering Intergenic Regions

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Abstract. *In this work, we explore heuristics for the Adjacency Graph Packing problem, which can be applied to the Double Cut and Join (DCJ) Distance Problem. The DCJ is a rearrangement operation and the distance problem considering it is a well established method for genome comparison. Our heuristics will use the structure called adjacency graph adapted to include information about intergenic regions, multiple copies of genes in the genomes, and multiple circular or linear chromosomes. The only required property from the genomes is that it must be possible to turn one into the other with DCJ operations. We propose one greedy heuristic and one heuristic based on Genetic Algorithms. Our experimental tests in artificial genomes show that the use of heuristics is capable of finding good results that are superior to a simpler random strategy.*

1. Introduction

In biology, it is often important to have metrics to compare genomes of different individuals. Such metrics can be used to infer evolutionary distance for the construction of phylogenetic trees. These metrics can also help in the identification of ortholog genes (genes separated by speciation).

Many metrics for genome comparison were proposed over time, including the well established rearrangement distance [Fertin et al. 2009]. The rearrangement distance is a measure of the number of rearrangement operations, large scale mutations affecting the order and orientation of the genetic material, needed to transform one genome into another. One of the most studied rearrangement operations is the Double Cut and Join (DCJ). The DCJ operation consists of cutting a genome in two points and joining the resulting parts.

Initially, the DCJ distance was studied in genomes with a single copy of each gene, and the orientation of the genes was taken into account [Yancopoulos et al. 2005]. In that scenario, there is an exact polynomial time algorithm to compute the distance in linear time [Bergeron et al. 2006]. A generalized version of the DCJ Distance Problem considers any two genomes as long as they have the same set of genes. In this case, the DCJ Distance Problem is NP-hard [Shao et al. 2015] and some proposed solutions

for it include an integer linear programming formulation [Shao et al. 2015] and an $O(k)$ -approximation algorithm [Rubert et al. 2017], where k is the maximum number of copies of a gene in the genomes.

In more recent works, a new approach was proposed to include information about intergenic regions [Fertin et al. 2017, Brito et al. 2020, Oliveira et al. 2021, Oliveira et al. 2024]. The usual representation in these works considers the number of nucleotides between genes, which is called the size of the intergenic region between these genes. With this representation, the DCJ Distance Problem is already NP-hard, even if the genomes have a single copy of each gene and there is a $4/3$ -approximation algorithm for it [Fertin et al. 2017].

The main structure proposed to deal with the DCJ Distance Problem is the adjacency graph. This graph was initially proposed for the problem without gene repetition [Bergeron et al. 2006] and is capable of representing multiple chromosomes, that can be linear or circular. This structure was later adapted to deal with multiple gene copies [Shao et al. 2015, Siqueira et al. 2021b] or intergenic regions [Fertin et al. 2017]. However, as far as we know, there is still no work combining multiple genes and intergenic regions for the DCJ Distance Problem.

This work proposes heuristics based on the adjacency graph for the DCJ Distance Problem considering genomes with repeated genes and taking into account intergenic regions and gene orientation. The heuristics are capable of dealing with genomes containing multiple circular or linear chromosomes. We assume that the genomes have the same set of genes. The next section introduces some definitions and formally states the problems. In Section 3, we describe the developed heuristics. In Section 4, we describe some experimental tests, and we conclude the paper in Section 5.

2. Definitions

We represent a genome as a set of chromosomes. Each *chromosome* \mathcal{C} is encoded by a pair (S, \check{S}) , composed of a string S of size $|S|$, representing the genes, and a list of non-negative integers \check{S} of size $|\check{S}|$, representing the intergenic regions. Each character of S has an associated sign $+$ or $-$ to represent the orientation of the correspondent gene. We use the term *label* to the symbol used to represent a character, disregarding the sign. Genes that are considered equal will be represented by characters with the same label. Our representation will allow for both linear and circular chromosomes. Figure 1 shows two genomes.

In linear chromosomes, we apply a process called *capping* for the representation. In this process, we insert a character $+\#$ at the beginning and at the end of S . We call these characters *caps*. They do not correspond to real genes, but to simplify our definitions they are treated as genes at the beginning and end of the chromosome. Considering this process, if \mathcal{C} is linear, then $|S| = |\check{S}| + 1$. If \mathcal{C} is circular, then $|S| = |\check{S}|$.

The i -th character of S , denoted by S_i , represents the i -th gene of the chromosome, if it is linear. If the chromosome is circular, we list the genes by cutting it at some point and assume that S_1 and $S_{|S|}$ are adjacent. The i -th integer of \check{S} , denoted by \check{S}_i , represents the size of the intergenic region between S_i and S_{i+1} . In circular chromosomes, the integer $\check{S}_{|\check{S}|}$ represents the size of the intergenic region between $S_{|S|}$ and S_1 .

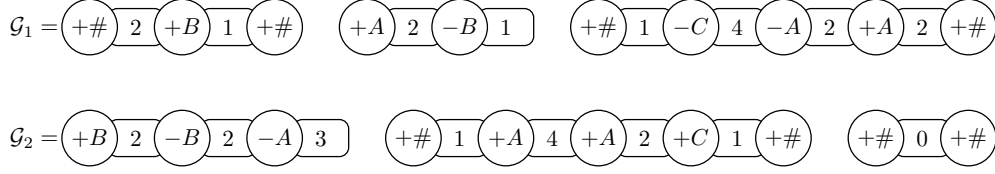


Figure 1. Two balanced genomes \mathcal{G}_1 and \mathcal{G}_2 . The chromosomes of \mathcal{G}_1 are represented by the following pairs of strings and list of integers $([+\#, +B, +\#], [2, 1])$, $([+A, -B], [2, 1])$, and $([+\#, -C, -A, +A, +\#], [1, 4, 2, 2])$. The chromosomes of \mathcal{G}_2 are represented by the following pairs of strings and list of integers $([+B, -B, -A], [2, 2, 3])$, $([+\#, +A, +A, +C, +\#], [1, 4, 2, 1])$, and $([+\#, +\#], [0])$. The linear chromosomes are capped and one chromosome was included in \mathcal{G}_2 to ensure the same number of chromosomes in both genomes.

Two genomes are called *balanced* if all labels, except $\#$, appear in the same number of genes, and the sum of the intergenic region sizes is the same in both genomes. In this work we only consider balanced genomes. Additionally, until the genomes have the same number of chromosomes, we add in the genome with fewer chromosomes linear chromosomes with two caps and an intergenic region of size 0 between them. With this addition of chromosomes the genomes remain balanced and have the same number of characters with the label $\#$.

Given two genomes \mathcal{G}_1 and \mathcal{G}_2 , the *adjacency graph* $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$ is composed of the vertex set V , edge set E (separated in two subsets E_r and E_d), and weight function $w : E_r \rightarrow \mathbb{N}$. For each character S_i in a string S from a chromosome of \mathcal{G}_1 or \mathcal{G}_2 , we have two vertices $v_{S_i}^t$ and $v_{S_i}^h$ in V , if S_i is not a cap, or we have one vertex v_{S_i} in V , if S_i is a cap. To simplify our examples, we will use the labels to represent the vertices (adding h and t accordingly). The edges in E_r are called *reality edges* and connect vertices of two consecutive characters S_i and S_{i+1} as follows:

- $v_{S_i}^h$ is connected with $v_{S_{i+1}}^t$, if both have sign $+$.
- $v_{S_i}^t$ is connected with $v_{S_{i+1}}^h$, if both have sign $-$.
- $v_{S_i}^h$ is connected with $v_{S_{i+1}}^h$, if S_i has sign $+$ and S_{i+1} has sign $-$.
- $v_{S_i}^t$ is connected with $v_{S_{i+1}}^t$, if S_i has sign $-$ and S_{i+1} has sign $+$.
- If S_i or S_{i+1} is a cap, just consider the above cases without the h or t in the vertex correspondent to this cap.

The edges in E_d are called *desire edges* and connect the vertices from a character S_i of a string S in a chromosome of \mathcal{G}_1 to the vertices of each character R_j of a string R of \mathcal{G}_2 with the same label as S_i . If S_i is a cap, v_{S_i} is connected with v_{R_j} . Otherwise, $v_{S_i}^t$ is connected with $v_{R_j}^t$ and $v_{S_i}^h$ is connected with $v_{R_j}^h$.

Two vertices $v_{S_i}^t$ and $v_{S_i}^h$ from the same character S_i are called *twin vertices* and two desire edges connecting $v_{S_i}^t$ with $v_{R_j}^t$ and $v_{S_i}^h$ with $v_{R_j}^h$ are called *twin edges*.

The weight function w is used to include intergenic region information in the graph. For each reality edge e connecting a vertex of S_i with a vertex of S_{i+1} , we have $w(e) = \check{S}_i$. Figure 2 shows an adjacency graph created from the genomes in Figure 1.

The *Double Cut and Join (DCJ)* operation can be described using the adjacency graph. Consider four vertices v_1, v_2, v_3 and v_4 , correspondent to characters of \mathcal{G}_1 , and

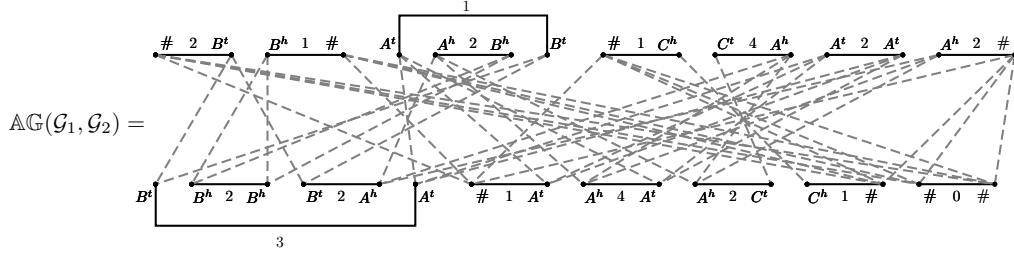


Figure 2. The adjacency graph from the genomes in Figure 1. The reality edges are shown as continuous lines, whose labels represent the weights, and the desire edges are shown as dashed lines.

two integers x and y , such that there is a reality edge e_1 connecting v_1 with v_2 and a reality edge e_2 connecting v_3 with v_4 . Besides, we require that $0 \leq x \leq w(e_1)$ and $0 \leq y \leq w(e_2)$. The $dcj(v_1, v_2, v_3, v_4, x, y)$ operation removes the edges e_1 and e_2 and adds new edges connecting v_1 with v_3 and connecting v_2 with v_4 . The new edges have weights $x + w(e_2) - y$ and $y + w(e_1) - x$, respectively. This operation can be interpreted as cutting the genome \mathcal{G}_1 in two intergenic regions and joining the resulting parts. The points where the genome is cut can be joined in different ways, depending on the order in which the vertices are passed as arguments to dcj . The representation of the genome has the signs of the reversed parts flipped, except caps that always have a positive sign. Figure 3 shows two examples of the DCJ operation.

The *DCJ Distance Problem* consists of finding the minimum number of DCJ operations to transform a genome \mathcal{G}_1 in another genome \mathcal{G}_2 .

An *alternating cycle* is a cycle that alternates between reality and desire edges. An alternating cycle is *balanced* if the sum of weights from the reality edges of one of the genomes is equal to the sum of weight from the reality edges of the other genome. An *alternating cycle packing* of the adjacency graph is a set of alternating cycles that do not share edges, contain all vertices, and every desire edge that has a twin edge in a cycle is also in a cycle.

An alternating cycle packing corresponds to an assignment of the genes of \mathcal{G}_1 to the genes of \mathcal{G}_2 by following the desire edges. With this assignment we can treat the genomes as if they do not have replicated genes. In that case, there is a known 4/3-approximation algorithm for the DCJ Distance Problem [Fertin et al. 2017]. So we are interested in finding a cycle packing that tries to produce an assignment that leads to the shortest DCJ distance. As the DCJ Distance Problem is NP-hard, we will use the 4/3-approximation algorithm instead of the exact distance.

3. Proposed Heuristics

We are going to adapt the heuristics proposed for a similar cycle packing problem [Siqueira et al. 2021b], where there are no intergenic regions and each genome has a single linear chromosome. Our heuristics are based on the idea of producing multiple cycle packing and selecting the best one by some criterion. This criterion can be the DCJ distance approximation, but to increase the efficiency of the algorithms we use only the number of balanced cycles in the packing as the selection criteria. The number r of packing to be produced is the same for all heuristics.

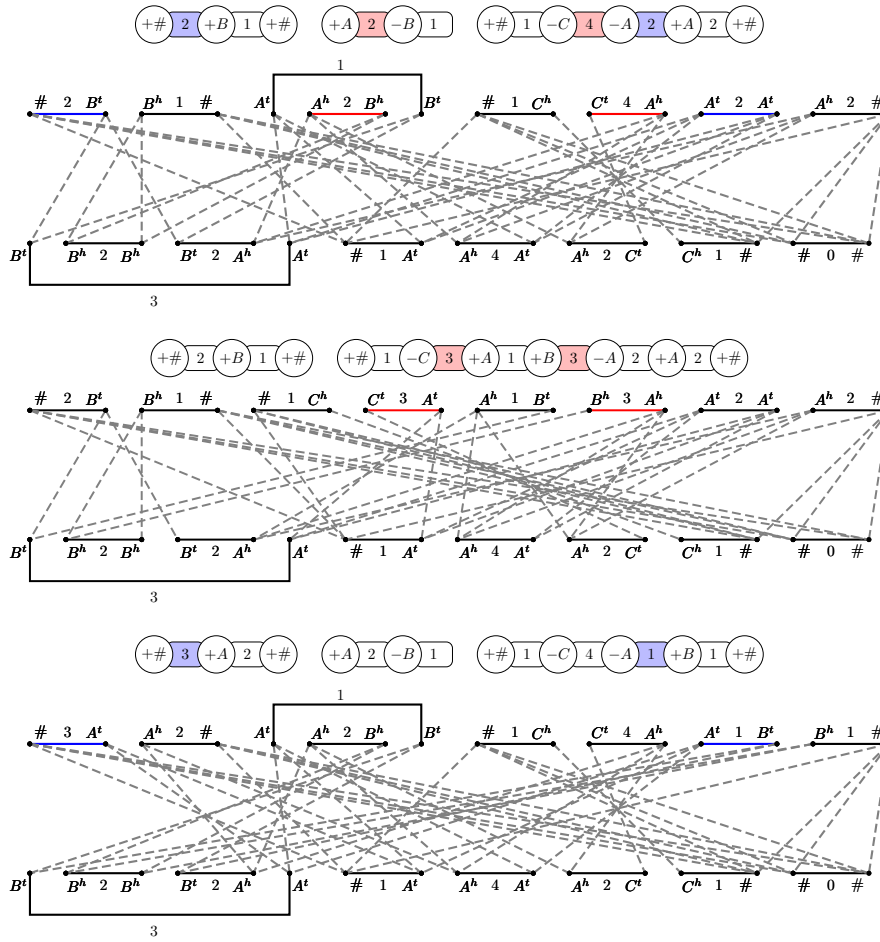


Figure 3. Two applications of the DCJ operation in the genome \mathcal{G}_1 of Figure 1 and in the graph of Figure 2. The first operation is $dcj(A^h, B^h, C^t, A^h, 1, 2)$, the cut and formed intergenic regions and edges are marked in red. The second operation is $dcj(\#, B^t, A^t, A^t, 1, 0)$, the cut and formed intergenic regions and edges are marked in blue.

The first heuristic, called Simple Random (SR), uses a simple random generation to generate the r packings. At each step, we select a random desire edge that is not the only desire edge incident to its vertices, and remove all other desire edges incident to its vertices. If that edge has a twin we do the same for it.

The second heuristic, called Greedy BFS (GBFS), uses a greedy approach by selecting a random vertex and applying a breadth-first search to find the shortest alternating cycle containing that vertex. For every edge selected for the cycle remove all other desire edges incident to its vertices and, if it has a twin, we remove all other desire edges incident to the vertices of this twin. We repeat this process until all vertices are in a cycle. During the breadth-first searches we must keep track of the edges that have to be removed to ensure that the cycle respects the restriction that every edge must have its twin in a cycle as well. Algorithm 1 shows a pseudo-code of this heuristic, and Figure 4 shows the production of one cycle packing.

The last heuristic uses a Genetic Algorithm (GA) approach [Mitchell 2008] to produce the set of packings. This approach is inspired by evolution and consists of producing

Algorithm 1: Greedy BFS

Data: A adjacency graph $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$ and the number of cycle packings r
Result: A cycle packing for $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$

- 1 $\mathbf{M} \leftarrow \emptyset$
- 2 **while** $|\mathbf{M}| < r$ **do**
- 3 $\mathcal{P} \leftarrow \emptyset$
- 4 **while** \mathcal{P} does not cover all vertices of $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$ **do**
- 5 $v \leftarrow$ vertex of $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$ not belonging to any cycle of \mathbf{M}
- 6 $C \leftarrow$ cycle resulting from a breadth-first search in $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$,
 starting with v
- 7 Remove the necessary vertices from $\mathbb{A}\mathbb{G}(\mathcal{G}_1, \mathcal{G}_2)$
- 8 $\mathcal{P} \leftarrow \mathcal{P} \cup C$
- 9 $\mathbf{M} \leftarrow \mathbf{M} \cup \{\mathcal{P}\}$
- 10 **return** Cycle packing belonging to \mathbf{M} with the largest number of balanced cycles

an initial set of individuals, called population, and then applying crossovers and mutations to generate new populations. The individuals are evaluated by a fitness function, which, in our case, is the number of balanced cycles in the packing. The algorithm stops when a fixed number of individuals are generated.

Besides the total number of individuals r , our genetic algorithm is specified by two parameters: the size of the population p ($0 \leq p \leq r$) and the mutation rate m ($0 \leq m \leq 1$). An initial population is generated by the GBFS heuristic. Until a total of r individuals are produced, at each iteration the algorithm produces a new population of p new individuals by applying the following steps:

- **Selection:** At this step, the algorithm selects, with repetition, $2p$ individuals to take part in the crossovers. The selection of each individual is by tournament, in which two individuals are randomly chosen and the one with the highest fitness is selected.
- **Crossover:** The $2p$ selected individuals are paired, and the algorithm applies the crossover operation in each pair to generate a new individual. Given two cycle packings \mathcal{P} and \mathcal{P}' of an adjacency graph, a *crossover* of \mathcal{P} and \mathcal{P}' is a new cycle packing created by the following procedure. Let L and L' be two randomly ordered lists of the cycles from \mathcal{P} and \mathcal{P}' , respectively. Starting with an empty set \mathcal{P}'' and with the original adjacency graph, while \mathcal{P}'' is not a cycle packing and there are cycles in L or L' , remove a cycle from L or L' (with a 50% probability to remove from each list, or 100% probability to remove from one list if the other is empty), and add it to \mathcal{P}'' if all its edges are still available. As in the GBFS heuristic, remove the necessary edges from the graph to ensure the restriction about the twin edges. If both lists L and L' are empty and \mathcal{P}'' is not yet a cycle packing, use breadth-first searches, as in the GBFS heuristic, to complete the packing. Figure 5 shows an example of crossover.
- **Mutation:** After the new individuals are generated, a mutation is applied to each one in order to increase the diversity of the population. In the mutation of a cycle

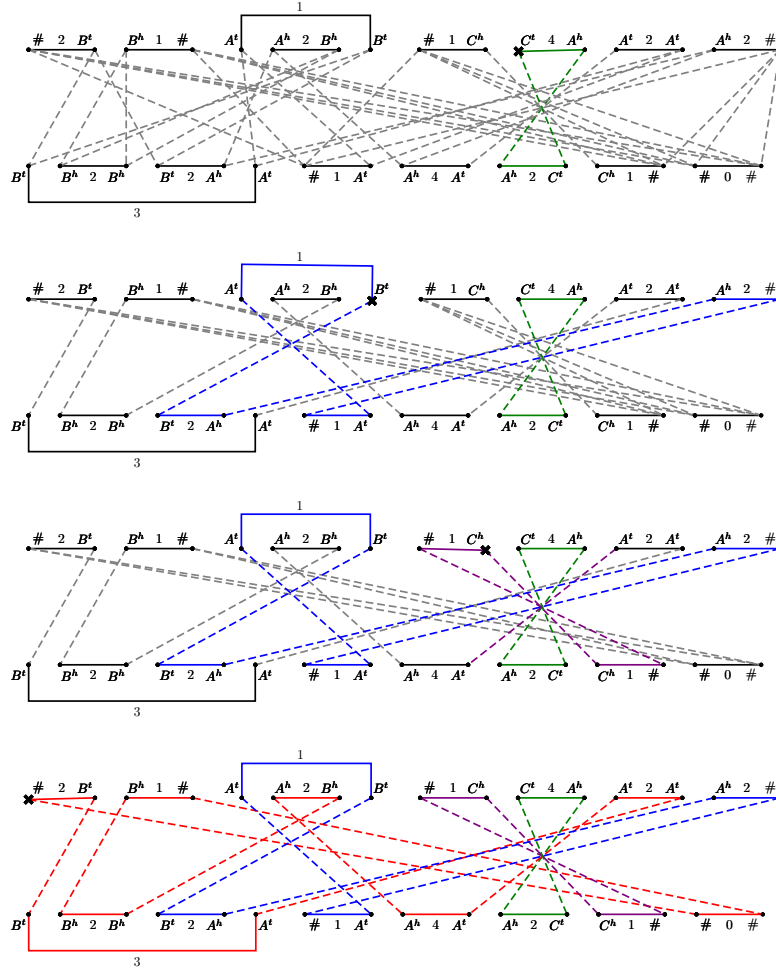


Figure 4. Example of the construction of one packing from the GBFS heuristic applied to the adjacency graph from Figure 2. The cycles obtained on each breadth-first search are marked with an \times indicating the starting vertex of the search. The resulting packing has two balanced cycles.

pack \mathcal{P}'' , we remove each cycle of \mathcal{P}'' with probability m and create a new packing from the remaining cycles of \mathcal{P}'' using breadth-first searches.

- **Elitism:** As we are going to replace the old population with the new one, we cannot guarantee the quality of the new population. So, to ensure that at least the best individual is kept, we replace the individual with the lowest fitness from the new population with the individual with the highest fitness from the old population, if the old individual has a higher fitness than the new one.

Algorithm 2 shows a pseudo-code of the GA heuristic.

4. Experiments

The instances used in our experiments and the implementation of the heuristics (in C++) are available at a public repository¹. The following tests were conducted on a computer

¹<https://github.com/compbio group/Heuristics-based-on-Adjacency-Graph-Packing-for-DCJ-Distance-Considering-Intergenic-Regions>

Algorithm 2: Genetic Algorithm

Data: An adjacency graph $\mathbb{AG}(\mathcal{G}_1, \mathcal{G}_2)$, the number of cycle packings r , size of the population p , and mutation rate m

Result: An adjacency packing for $\mathbb{AG}(\mathcal{G}_1, \mathcal{G}_2)$

- 1 $\mathbf{P}_0 \leftarrow p$ cycle packings generated by the GBFS heuristic
- 2 $\mathbf{M} \leftarrow \mathbf{P}_0$
- 3 $i \leftarrow 0$
- 4 **while** $|\mathbf{M}| < r$ **do**
- 5 $\mathbf{S} \leftarrow$ Sequence of p selected pairs of individuals from \mathbf{P}_i
- 6 $\mathbf{P}'_{i+1} \leftarrow$ Set of p new individuals created by crossover of individuals from \mathbf{S}
- 7 $\mathbf{P}''_{i+1} \leftarrow$ Set of p individuals obtained by mutation of individuals from \mathbf{P}'_{i+1}
- 8 **if** the best individual of \mathbf{P}_i is better than the worse individual of \mathbf{P}''_{i+1} **then**
- 9 $\mathbf{P}_{i+1} \leftarrow$ Set \mathbf{P}''_{i+1} with its worse individual replaced with the best individual of \mathbf{P}_i
- 10 **else**
- 11 $\mathbf{P}_{i+1} \leftarrow \mathbf{P}''_{i+1}$
- 12 $\mathbf{M} \leftarrow \mathbf{M} \cup \{\mathbf{P}_{i+1}\}$
- 13 $i \leftarrow i + 1$
- 14 **return** Cycle packing belonging to \mathbf{M} with the largest number of balanced cycles

equipped with an “Intel(R) Xeon(R) CPU E5-2470 v2” with 40 cores at 2.40GHz, 32GB of RAM, and 9GB of swap space.

We created a database of simulated genomes to test our heuristics. We produced 25000 pair of genomes separated into groups of 1000. Each group is defined by the number ℓ of labels used and the number o of DCJ operations applied. We created each pair by the following process (each random selection is uniformly distributed):

- We randomly produced 200 characters picking the label from a set with ℓ symbols and choosing the signs randomly.
- We randomly chose 201 integers in the interval $[0, 100]$ to represent the intergenic regions.
- We produced a genome \mathcal{G} containing a single linear chromosome with the chosen characters and intergenic regions.
- We created the first genome of the pair by applying $o/2$ DCJ operations in \mathcal{G} .
- We created the second genome of the pair by applying $o/2$ DCJ operations in \mathcal{G} .

For our tests all heuristics produced a set of $r = 1000$ alternating cycle packings. For the GA heuristic, we chose the parameters $m = 0.2$ and $p = 100$.

In Table 1 and Table 2 we can see the results of our experiments. For each heuristic and each group of instances, we show, in Table 1, the average and standard deviation of the distance computed from the best cycle packing found and, in Table 2, the average and standard deviation of the execution time (in seconds) of the algorithms. In both tables the

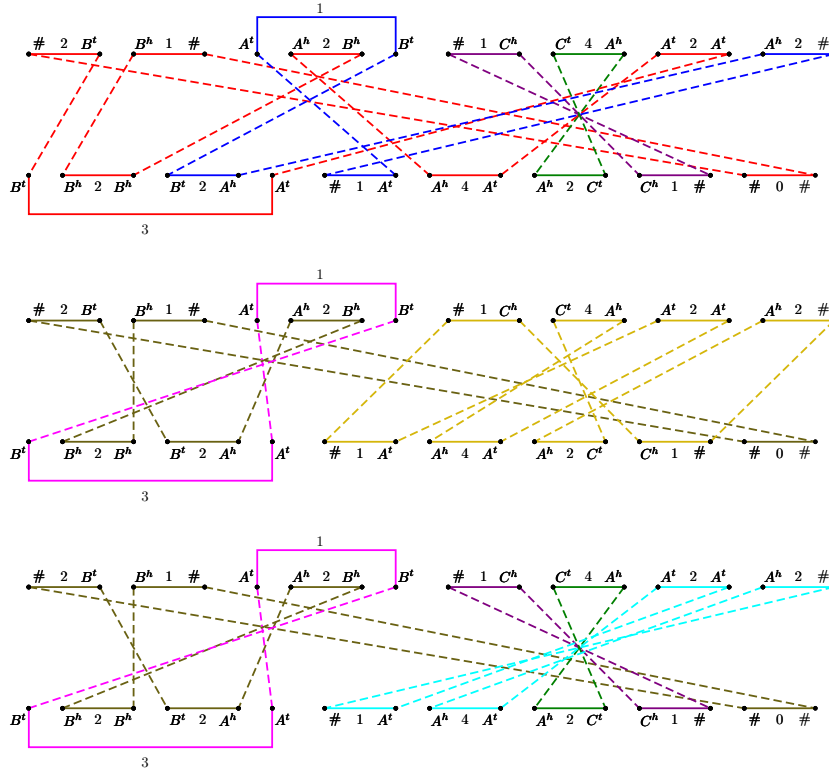


Figure 5. Example of the crossover of two cycle packing of the graph in Figure 2. Two cycles are selected from the first packing, two from the second packing, and one new cycle is inserted by a breadth-first search to complete the packing.

standard deviation is shown as a percentage of the mean.

We can see that GA found the shortest distance on average, followed by GBFS and SR. That indicates that using a more sophisticated strategy to produce the packings results in a better result. The largest difference between the average distances found by GA and by GBFS is 54.44 for $o = 40$ and $\ell = 40$, while the shortest such difference is 6.29 for $o = 10$ and $\ell = 50$. Considering the standard deviation, the distinction between the distances found by GA and by GBFS is clearer with larger values of o .

Additionally, the running time of GA was shorter than the running time of the other heuristics. This is a consequence of the fact that the GA heuristic does not produce all cycle packings from scratch, but produces most packings from crossovers and mutations.

5. Conclusion

We described and tested three heuristics for Adjacency Graph Packing that can be used to estimate DCJ distances. One of the heuristics is based on a simple random strategy (SR), another uses a greedy strategy based on BFS (GBFS), and the last one is based on Genetic Algorithms (GA). These heuristics were tested on a database of simulated genomes and GA found the shorter distances on average with the shortest running time.

This work can be further extended to consider genomes with distinct sets of genes, as was done for the original heuristics developed for unichromosomal genomes with-

Table 1. Average and standard deviation for the DCJ distances resulting from the heuristics.

o	ℓ	GA		GBFS		SR	
10	10	145.03	± 15.21%	172.02	± 10.86%	204.44	± 11.14%
10	20	59.03	± 51.18%	108.45	± 35.98%	148.47	± 33.12%
10	30	25.60	± 66.17%	51.23	± 64.45%	87.31	± 63.51%
10	40	16.15	± 63.07%	27.74	± 75.39%	54.80	± 86.22%
10	50	12.01	± 44.94%	18.30	± 73.85%	35.55	± 99.68%
20	10	163.85	± 9.80%	186.75	± 6.25%	214.11	± 5.70%
20	20	103.75	± 28.18%	152.03	± 14.79%	182.64	± 13.58%
20	30	59.35	± 38.88%	109.22	± 30.38%	146.76	± 26.82%
20	40	37.29	± 43.65%	75.06	± 43.67%	109.62	± 41.23%
20	50	28.58	± 37.72%	52.02	± 49.17%	81.83	± 52.58%
30	10	175.16	± 7.61%	194.90	± 5.45%	217.66	± 4.02%
30	20	135.22	± 17.37%	172.33	± 8.77%	196.27	± 7.93%
30	30	96.39	± 27.27%	147.38	± 14.37%	172.66	± 13.77%
30	40	68.36	± 34.96%	120.39	± 22.65%	149.94	± 19.43%
30	50	51.12	± 32.35%	93.15	± 30.95%	123.60	± 29.17%
40	10	185.00	± 5.75%	201.67	± 4.20%	219.28	± 3.38%
40	20	156.60	± 10.76%	184.72	± 6.37%	203.64	± 5.53%
40	30	119.66	± 18.99%	165.50	± 9.60%	186.53	± 8.91%
40	40	92.53	± 24.36%	146.97	± 13.69%	169.63	± 12.04%
40	50	75.16	± 26.47%	127.49	± 17.35%	151.83	± 16.10%
50	10	191.97	± 4.72%	205.41	± 3.74%	221.23	± 2.86%
50	20	168.78	± 7.91%	192.35	± 5.03%	208.59	± 4.08%
50	30	144.03	± 12.30%	178.56	± 7.17%	195.46	± 6.28%
50	40	117.53	± 18.38%	163.73	± 9.80%	182.52	± 8.61%
50	50	98.91	± 21.72%	147.36	± 12.51%	168.67	± 11.39%

out intergenic regions [Siqueira et al. 2024]. There are known algorithms for rearrangement distances with intergenic regions and distinct sets of genes, but without gene replicas [Alexandrino et al. 2021a, Alexandrino et al. 2021b]. It is also relevant to explore other approaches to the problem, including exact and approximation algorithms. The use of String Partition problems with intergenic regions can be one approach to develop approximations [Siqueira et al. 2021a, Siqueira et al. 2022].

Another path for future works is the application of the proposed algorithms to compare real genomes. These tests can include the integration of the algorithms in bioinformatic tools used in the construction of phylogenetic trees or detection of orthologous genes.

Acknowledgement

This work was supported by the São Paulo Research Foundation, FAPESP (grant 2021/13824-8).

Table 2. Average and standard deviation for the running times of the heuristics (in seconds).

o	ℓ	GA		GBFS		SR	
10	10	36.78	± 4.11%	213.56	± 26.05%	40.78	± 14.53%
10	20	25.73	± 7.41%	136.13	± 41.69%	40.61	± 13.82%
10	30	16.10	± 6.94%	67.38	± 37.00%	38.25	± 13.56%
10	40	11.88	± 5.58%	37.31	± 28.46%	37.52	± 12.81%
10	50	10.31	± 4.05%	22.75	± 19.60%	34.95	± 11.60%
20	10	37.91	± 3.46%	225.08	± 23.60%	39.70	± 14.25%
20	20	32.76	± 6.35%	172.17	± 35.19%	39.64	± 14.05%
20	30	26.04	± 8.09%	119.97	± 44.08%	39.62	± 13.48%
20	40	19.13	± 8.04%	82.09	± 43.96%	38.43	± 12.97%
20	50	15.39	± 7.88%	56.28	± 38.76%	36.28	± 11.96%
30	10	38.07	± 3.45%	232.65	± 24.05%	41.24	± 14.62%
30	20	37.30	± 5.73%	195.65	± 29.77%	40.57	± 14.26%
30	30	33.24	± 7.82%	164.24	± 40.45%	40.24	± 13.33%
30	40	29.02	± 9.58%	132.96	± 47.19%	39.22	± 13.39%
30	50	24.13	± 9.82%	102.31	± 49.57%	37.05	± 12.37%
40	10	39.03	± 3.38%	240.62	± 21.66%	41.05	± 14.24%
40	20	39.48	± 4.90%	202.19	± 29.20%	40.69	± 14.11%
40	30	38.01	± 6.84%	180.91	± 36.32%	40.47	± 14.18%
40	40	35.15	± 8.67%	160.33	± 45.16%	40.09	± 13.15%
40	50	31.70	± 9.77%	139.39	± 47.48%	38.39	± 12.49%
50	10	39.62	± 3.54%	243.66	± 20.86%	41.44	± 14.56%
50	20	40.99	± 4.98%	211.95	± 28.17%	39.90	± 13.85%
50	30	41.46	± 6.53%	194.58	± 36.40%	40.08	± 13.76%
50	40	39.86	± 8.21%	180.37	± 43.45%	40.55	± 13.47%
50	50	38.77	± 9.86%	161.15	± 46.69%	39.22	± 13.08%

References

- Alexandrino, A. O., Brito, K. L., Oliveira, A. R., Dias, U., and Dias, Z. (2021a). Reversal Distance on Genomes with Different Gene Content and Intergenic Regions Information. In *Proceedings of the 8th International Conference on Algorithms for Computational Biology (AlCoB'2021)*, volume 12715, pages 121–133. Springer International Publishing.
- Alexandrino, A. O., Oliveira, A. R., Dias, U., and Dias, Z. (2021b). Incorporating Intergenic Regions into Reversal and Transposition Distances with Indels. *Journal of Bioinformatics and Computational Biology*, 19(06):2140011.
- Bergeron, A., Mixtacki, J., and Stoye, J. (2006). A Unifying View of Genome Rearrangements. In *International Workshop on Algorithms in Bioinformatics*, pages 163–173. Springer.

- Brito, K. L., Jean, G., Fertin, G., Oliveira, A. R., Dias, U., and Dias, Z. (2020). Sorting by Genome Rearrangements on both Gene Order and Intergenic Sizes. *Journal of Computational Biology*, 27(2):156–174.
- Fertin, G., Jean, G., and Tannier, E. (2017). Algorithms for Computing the Double Cut and Join Distance on both Gene Order and Intergenic Sizes. *Algorithms for Molecular Biology*, 12(1):16.
- Fertin, G., Labarre, A., Rusu, I., Tannier, É., and Vialette, S. (2009). *Combinatorics of Genome Rearrangements*. Computational Molecular Biology. The MIT Press, London, England.
- Mitchell, M. (2008). *Introduction to Genetic Algorithms*. Springer Berlin Heidelberg, Cambridge, MA, USA.
- Oliveira, A. R., Brito, K. L., Alexandrino, A. O., Siqueira, G., Dias, U., and Dias, Z. (2024). Rearrangement distance problems: An updated survey. *ACM Computing Surveys*, 56(8).
- Oliveira, A. R., Jean, G., Fertin, G., Brito, K. L., Bulteau, L., Dias, U., and Dias, Z. (2021). Sorting Signed Permutations by Intergenic Reversals. *IEEE/ACM Transactions on Computational Biology and Bioinformatics*, 18(6):2870–2876.
- Rubert, D. P., Feijão, P., Braga, M. D. V., Stoye, J., and Martinez, F. H. V. (2017). Approximating the DCJ Distance of Balanced Genomes in Linear Time. *Algorithms for Molecular Biology*, 12(1):1–13.
- Shao, M., Lin, Y., and Moret, B. M. (2015). An exact algorithm to compute the double-cut-and-join distance for genomes with duplicate genes. *Journal of Computational Biology*, 22(5):425–435.
- Siqueira, G., Alexandrino, A. O., and Dias, Z. (2022). Signed Rearrangement Distances Considering Repeated Genes and Intergenic Regions. *Proceedings of 14th International Conference on Bioinformatics and Computational Biology (BICoB'2022)*, 83:31–42.
- Siqueira, G., Alexandrino, A. O., Oliveira, A. R., and Dias, Z. (2021a). Approximation Algorithm for Rearrangement Distances Considering Repeated Genes and Intergenic Regions. *Algorithms for Molecular Biology*, 16(1):1–23.
- Siqueira, G., Oliveira, A. R., Alexandrino, A. O., and Dias, Z. (2021b). Heuristics for Cycle Packing of Adjacency Graphs for Genomes with Repeated Genes. In *Proceedings of the 14th Brazilian Symposium on Bioinformatics (BSB'2021)*, pages 93–105. Springer International Publishing.
- Siqueira, G., Oliveira, A. R., Alexandrino, A. O., Jean, G., Fertin, G., and Dias, Z. (2024). Assignment of orthologous genes in unbalanced genomes using cycle packing of adjacency graphs. *Journal of Heuristics*.
- Yancopoulos, S., Attie, O., and Friedberg, R. (2005). Efficient Sorting of Genomic Permutations by Translocation, Inversion and Block Interchange. *Bioinformatics*, 21(16):3340–3346.