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We extend the work of Robinson and Turner to use hypothesis testing with persistent homology to test for measurable differences in shape between the spaces of three or more groups. We conduct a large-scale simulation study to validate our proposed extension, considering various combinations of groups, sample sizes and measurement errors. For each such combination, the percentage of p-values below an α -level of 0.05 is provided. Additionally, we apply our method to a cardiocography data set and find statistically significant evidence of measurable differences in shape between the spaces corresponding to normal, suspect and pathologic health status groups.

1. Introduction

Consider a data set, obtained via random sampling, where each data point is a vector of m quantitative variables and one categorical variable with s levels. Ideally, several of the quantitative variables are real-valued. According to the levels of the categorical variable, we will group the data points into s not necessarily distinct collections of points in \mathbb{R}^m , referred to as point clouds. For each group, we can view the corresponding point cloud as a representative subset of a space which consists of all such points in \mathbb{R}^m with the respective level of the categorical variable. Of interest is whether or not these s spaces have measurably different shapes? But what does shape even mean if m is large?

Topology, in particular algebraic topology, is an area of mathematics that can be used to qualitatively measure the shape of a point cloud. For a given point cloud, we construct an infinite family of simplicial complexes that vary according to a real-valued distance parameter. Each complex in the family is an object that inherits a shape from the point cloud and the topological tool known as homology can be used

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to detect this shape. Since any single complex within the infinite family corresponds to a choice of parameter value, we might ask which parameter value “best” captures the shape of the point cloud? Persistent homology is a study of the homological features that persist over long intervals of the distance parameter, thus sidestepping the search for a best choice parameter value. Hence, persistent homology can be used to determine if point clouds have different shape. While persistent homology allows comparisons of shapes across point clouds obtained from a sample of data points, can any resulting differences then be generalized to the corresponding spaces at large? The answer is yes, but as random sampling unavoidably introduces variability, a method is needed which can distinguish true differences in shape between the spaces from artificial differences in shape between the point clouds obtained via random sampling of data points. Statistical hypothesis testing is an inferential method often implemented to assess whether or not randomly sampled data provide sufficient evidence of a difference, with respect to some characteristic, between two or more populations (or, as we have been calling them, spaces). K. Turner and A. Robinson [2013] conducted such an assessment on $s = 2$ spaces using a specific type of hypothesis testing procedure known as a permutation test, where the characteristic of interest is shape, as measured via persistent homology. As this procedure requires multiple point clouds from both spaces, in practice the two point clouds obtained from the random sample of data points are further partitioned, via subsampling, into multiple “smaller”, or less dense, point clouds. The assessment is then conducted using the persistent homology of these subsampled point clouds within the procedure. We extend this procedure to three or more spaces, $s \geq 3$.

The remainder of the paper is organized as follows. In Section 2 we provide definitions of the Vietoris–Rips complex of a point cloud, homology groups, persistent homology and persistence diagrams. In Section 3 we describe the permutation test of Robinson and Turner. In Section 4 we propose an extension of the permutation test for three or more spaces. In Section 5 we present the results of a large-scale simulation study, incorporating various measurement errors and sample sizes, that validate our proposed extension. Finally, in Section 6 we apply our extension to a cardiocography data set and find significant evidence of differences in shape, as measured by persistent homology, between the spaces corresponding to normal, suspect and pathologic health groups.¹

2. Persistent homology

Before defining the persistent homology of a point cloud, we associate to the point cloud a nested family of abstract simplicial complexes. A thorough explanation of

¹ Throughout we use “difference in shape” to mean shape as measured by persistent homology in a specified dimension.

simplicial complexes and abstract simplicial complexes is given in [Edelsbrunner and Harer 2010; Munkres 1984]. Here we motivate the definition of an abstract simplicial complex with a brief geometric introduction to simplicial complexes, followed by the definition of the Vietoris–Rips complex, which is the abstract simplicial complex used herein.

Geometrically, a 0-simplex is a point, a 1-simplex is a line segment, a 2-simplex is a triangular subset of a plane, a 3-simplex is a solid tetrahedron, and an n -simplex is the n -dimensional analogue of these convex sets. Observe that the boundary of an n -simplex, σ , is a collection of $(n-1)$ -simplices; these boundary simplices are called faces of σ . A *simplicial complex* is a collection of simplices in \mathbb{R}^d that satisfy certain subset and intersection properties specifying how simplices can be put together to create a larger structure. More precisely, a simplicial complex is a finite collection of simplices, K , such that (1) if $\sigma \in K$ and ρ is a face of σ then $\rho \in K$, and (2) given any two simplices $\sigma_1, \sigma_2 \in K$, either $\sigma_1 \cap \sigma_2$ is the empty set or a face of both σ_1 and σ_2 . More generally, and without relying on geometry, an *abstract simplicial complex* is a finite collection of sets, A , such that if $\alpha \in A$ and $\beta \subseteq \alpha$, then $\beta \in A$. It is well known that a finite abstract simplicial complex can be geometrically realized as a simplicial complex in \mathbb{R}^N for N sufficiently large.

2.1. The Vietoris–Rips complex. The *Vietoris–Rips complex*, denoted $\text{VR}(D, r)$, is an abstract simplicial complex associated to a point cloud D for a fixed radius value $r > 0$. The elements of D form the 0-simplices or vertex set of $\text{VR}(D, r)$. A simplex of $\text{VR}(D, r)$ is a finite subset α of D such that the diameter of α is less than r . A simplex $\alpha \subseteq D$ with k -elements is called a $(k-1)$ -simplex of D . Thus, a 1-simplex corresponds to a two element set (viewed geometrically as the endpoints of a line segment), a 2-simplex corresponds to a three element set (viewed as the vertices of a triangle), and so on. Observe that if α is a k -simplex, then every subset of α is a simplex of D as the diameter of a subset of α can be no larger than the diameter of α . Hence the Vietoris–Rips complex satisfies the definition of an abstract simplicial complex. For readers that are new to topological data analysis, an example Vietoris–Rips complex is given in the Appendix.

We note that Vietoris–Rips complexes for increasing radius values are always a nested family of simplicial complexes associated to D ; that is, the complexes satisfy

$$\text{VR}(D, r_1) \subseteq \text{VR}(D, r_2) \quad \text{whenever } r_1 \leq r_2.$$

This nested feature of the complexes along with the functorial nature of homology are what give rise the concept of persistence to be defined below.

Although the Vietoris–Rips complex is relatively straightforward to define and calculate, it can be computationally expensive when used with large point clouds. There are economical alternatives to the Vietoris–Rips complex, such as the lazy

witness complex introduced in [de Silva and Carlsson 2004]. Persistent homology can be applied using any nested family of complexes indexed by some parameter.

2.2. Homology. The homology of a simplicial complex K is an algebraic measurement of how the n -simplices are attached to the $(n-1)$ -simplices within K . Below we define some technical machinery (chains, boundary maps, and cycles) used to define homology groups.

The p -chains of a simplicial complex K , denoted $C_p(K)$, is the group of formal linear combinations of the p -simplices of K with coefficients from \mathbb{Z}_2 . (More general definitions of homology with ring coefficients can be found in the standard algebraic topology texts [Edelsbrunner and Harer 2010; Hatcher 2002].) Since \mathbb{Z}_2 is a field, the p -chains of K are \mathbb{Z}_2 -vector spaces with basis the p -simplices of K .

The *boundary map*, denoted δ_p , identifies each p -chain with its boundary, a $(p-1)$ -chain. Each boundary map, $\delta_p : C_p \rightarrow C_{p-1}$, is a homomorphism and in the case of \mathbb{Z}_2 coefficients, as considered here, these maps are linear transformations.

Notice that $\delta_p \circ \delta_{p+1}$ is the zero map as the boundary of a boundary is empty. This fundamental property of chain complexes ensures that the image of δ_{p+1} is a normal subgroup of the kernel of δ_p . The collective sequence of boundary maps and chains, as shown below, is called a *chain complex*:

$$\dots \xrightarrow{\delta_n} C_n(K) \xrightarrow{\delta_{n-1}} \dots \xrightarrow{\delta_2} C_1(K) \xrightarrow{\delta_1} C_0(K) \xrightarrow{\delta_0} 0.$$

Homology groups are defined using both the kernel and image of each boundary map. The kernel of δ_p is the set of all p -chains whose boundary is empty. The elements of the kernel of δ_p are called p -cycles of K . The image of δ_{p+1} is the set of p -chains that are boundaries of a $(p+1)$ -chain. The p -th homology group of K , denoted $H_p(K; \mathbb{Z}_2)$, is defined as the quotient group $\ker(\delta_p) / \text{im}(\delta_{p+1})$.

As the parameter $r > 0$ increases, the Vietoris–Rips complex includes more simplices, thus the homology of the complex changes. The functorial property of homology and the inclusion map $i : \text{VR}(D, r_1) \rightarrow \text{VR}(D, r_2)$ whenever $r_1 \leq r_2$, give rise to induced maps between the homology of the complexes

$$i_* : H_*(\text{VR}(X, r_1); \mathbb{Z}_2) \rightarrow H_*(\text{VR}(X, r_2); \mathbb{Z}_2).$$

A nontrivial homology class $\alpha \in H_*(\text{VR}(X, r_1); \mathbb{Z}_2)$ is said to be born at radius r_b if r_b is the least radius value for which $H_*(\text{VR}(X, r_b); \mathbb{Z}_2)$ contains an element mapping onto α under the map

$$H_*(\text{VR}(X, r_b); \mathbb{Z}_2) \rightarrow H_*(\text{VR}(X, r_1); \mathbb{Z}_2).$$

The homology class α is said to die at radius value r_d provided that r_d is the least radius value for which the class α maps to zero in the mapping

$$H_*(\text{VR}(X, r_1); \mathbb{Z}_2) \rightarrow H_*(\text{VR}(X, r_d); \mathbb{Z}_2).$$

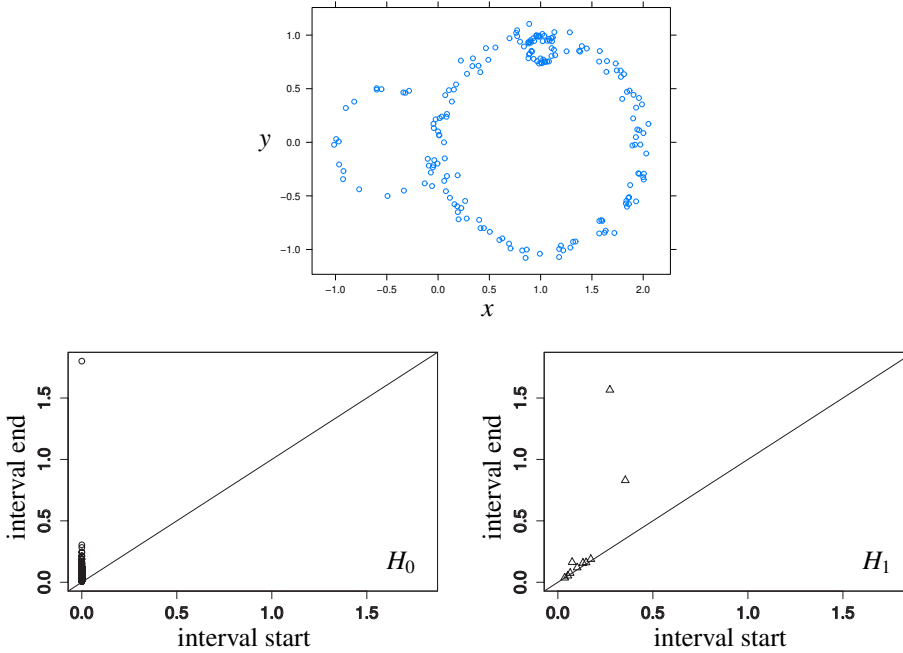


Figure 1. An example data set (top) and the corresponding persistence diagrams for the homological dimensions 0 and 1.

The topological feature that α represents is then said to have a birth and death “time” corresponding to the radius values r_b and r_d . We say that the class α persists over the interval $[r_b, r_d]$. Persistent homology of a data set D is a cataloguing of the homological classes of the abstract simplicial complexes $\text{VR}(D, r)$ that persist for large intervals of radius values, r .

For a fixed k , the *persistence diagram for $H_k(\text{VR}(X, *); \mathbb{Z}_2)$* is a plot of points (r_b, r_d) for each nonzero class $\alpha \in H_k(\text{VR}(X, *); \mathbb{Z}_2)$.

Figure 1 contains an example data set that includes several 1-dimensional homological features of varying size and the corresponding persistence diagrams in dimensions 0 and 1.

Within the persistence diagram in Figure 1, we see two lone triangles at the points $p_1 = (0.35, 0.8)$ and $p_2 = (0.3, 1.55)$. The point p_2 , with the early birth time, is the 1-dimensional homology class representing the larger circular feature on the right. The earlier birth time is due to the closer scattering of the data points about the larger circle. The point p_1 , with the earlier death time, is the 1-dimensional homology class representing circle of smaller radius on the left. The early death time is due to the smaller radius of this circular feature. The persistence diagram in Figure 1 also contains several triangles near the diagonal which represent classes that only persist for a short while, and it includes a triangle at the point $(0.1, 0.15)$

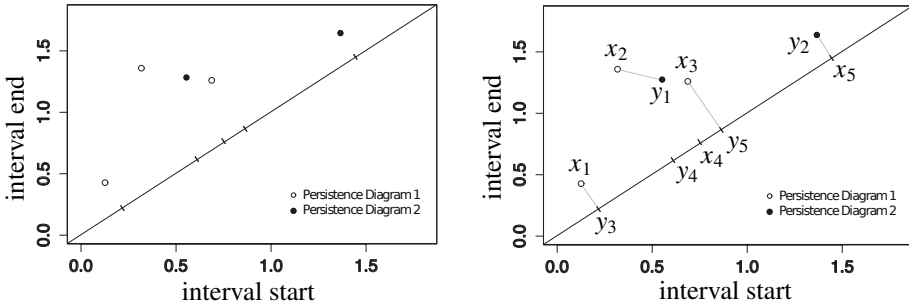


Figure 2. On the left, two superimposed persistence diagrams of the same homological dimension. On the right, the points $\{x_1, \dots, x_5\}$, $\{y_1, \dots, y_5\}$ and line segments indicating the optimal bijection. The diagram distance is the sum of the lengths of the line segments $\overline{x_1 y_3} + \overline{x_2 y_1} + \overline{x_3 y_5} + \overline{x_5 y_2}$. The segment $\overline{x_4 y_4}$ is not included as it is a segment between diagonal points.

representing the 1-dimensional homology class resulting from the tiny circle of points at the top of the larger circle. Notice that the 0-dimensional homology classes, which are plotted as small circles in the persistence diagram, all have birth time $r = 0$ as a result of each data point representing a unique 0-dimensional class at $r = 0$. As r increases, the complex consists of fewer connected components until it is one connected component. The 0-dimensional persistence class plotted at the point $(0, 0.35)$ represents the joining of the last two components into a single component. In other words, for $r \geq 0.35$ the simplicial complex $VR(X, r)$ is one connected component. The 0-dimensional class plotted at $(0, 2)$ is merely the result of using a maximum radius value of $r = 2$ in the persistent homology calculation. This class indicates that the complex $VR(X, 2)$ is one connected component.

The discussion above defines a persistence diagram for a data set using the Vietoris–Rips complex. There are, however, several other routes that lead to the creation of a persistence diagram. The omnibus test described below can be applied to a collection of persistence diagrams obtained by any means.

2.3. A metric on persistence diagrams. We follow Robinson and Turner in selecting the metric on persistence diagrams that is analogous to the L^2 norm in the space of functions on a discrete space. Given two persistence diagrams X and Y , let $x_1, x_2, \dots, x_n \in X$ be a listing of the off-diagonal points of X and $y_1, y_2, \dots, y_m \in Y$ be the off-diagonal points of Y . Select points x_{n+1}, \dots, x_{n+m} and y_{m+1}, \dots, y_{m+n} along the diagonal so that x_{n+k} is the point closest (in Euclidean distance) to y_k and vice versa. Let $X' = \{x_1, \dots, x_{n+m}\}$ and $Y' = \{y_1, \dots, y_{n+m}\}$. We consider the set of all bijections $\phi : X' \rightarrow Y'$ such that (1) the off-diagonal point x_k is paired either with an off-diagonal point of Y or with y_{m+k} and (2) the diagonal point x_l

is paired either with y_{l-n} or with one of the diagonal points in Y' . For a specific bijection ϕ , if both x_k and y_j are diagonal points, the *cost* of assigning x_k to y_j , denoted $C(x_k, y_j)$, is 0, else the cost is the Euclidean distance between x_k and y_j .

Define $d(X, Y)$, the *distance between the persistence diagrams X and Y* , by

$$d(X, Y) = \left(\inf_{\phi: X' \rightarrow Y'} \sum_{x \in X'} C(x, \phi(x)) \right)^{\frac{1}{2}}.$$

A bijection between X and Y is called optimal if it achieves the infimum. The Hungarian algorithm [Kuhn 1955; Munkres 1957], also known as Munkres' assignment algorithm, presents a method for obtaining an optimal bijection in polynomial time. Figure 2 gives an example of two simple persistence diagrams and the bijection exhibiting their diagram distance.

3. Hypothesis testing and topological data analysis

When persistent homology is applied to point clouds obtained from a random sample of points from various spaces, an element of variability is unavoidably introduced. Point clouds obtained from different samples of the same space, if somewhat representative, are expected to have “small” differences in their respective persistence diagrams, while point clouds obtained from samples of different spaces are expected to have comparatively “large” differences in their persistence diagrams. However, when the true shape-related features of two spaces are unknown, and all that is available are the point clouds obtained from samples of each of these spaces, what qualifies as a small or large difference is unclear. A tool is needed which can determine whether or not the shapes of the underlying spaces are measurably different. Statistical hypothesis testing is a method that can be implemented in these situations to decide if there is sufficient evidence to classify the shapes of the spaces as measurably different. A thorough development of statistical hypothesis testing is available in many standard sources, including [Casella and Berger 2002; DeGroot and Schervish 2012].

3.1. Hypothesis testing via the joint loss function. Consider two spaces in \mathbb{R}^m , arbitrarily labeled X_1 and X_2 , suspected of having measurably different shapes. Suppose n_1 point clouds are available from X_1 and n_2 point clouds are available from X_2 , with their corresponding persistence diagrams in a fixed dimension denoted respectively by $X_{1,1}, X_{1,2}, \dots, X_{1,n_1}$ and $X_{2,1}, X_{2,2}, \dots, X_{2,n_2}$. Further suppose that each of these $n_1 + n_2$ point clouds was obtained via random sampling from either X_1 or X_2 . Note that in practice, for each space, a single point cloud will usually be obtained via a random sample of X_i and then partitioned, via subsampling, into n_i smaller, or less dense, point clouds. Within the statistical hypothesis testing paradigm, the null hypothesis asserts that the shapes of X_1 and X_2 are not measurably

different, while the alternative hypothesis asserts the opposite. The corresponding test statistic, proposed by Robinson and Turner [2013], is the *joint loss function*

$$\sigma_{\chi_2}^2 = \sum_{m=1}^2 \frac{1}{2n_m(n_m - 1)} \sum_{i=1}^{n_m} \sum_{j=1}^{n_m} d(X_{m,i}, X_{m,j})^2,$$

where $d(\cdot, \cdot)$ is the persistence diagram distance metric described in Section 2.3.

The joint loss function is ultimately an aggregate measure of within-group variation. More specifically, $\sigma_{\chi_2}^2$ adds the variation in the $\binom{n_1}{2}$ persistence diagram distances from X_1 and the variation in the $\binom{n_2}{2}$ persistence diagram distances from X_2 . Unfortunately, the sampling distribution of $\sigma_{\chi_2}^2$ is nontrivial to determine and is currently unknown, which renders the “standard” (i.e., distribution-based) hypothesis testing paradigm impossible. To circumvent this, Robinson and Turner propose implementing a permutation test, which in this context is free of any distributional assumptions. A thorough development of permutation tests, and the often corresponding approximate permutation test p-values, is available in [Higgins 2004; Ramsey and Schafer 2013].

To perform the permutation test, we assume that the null hypothesis is true, i.e., X_1 and X_2 are not measurably different in shape. Such an assumption effectively means that the observed labeling of the point clouds to either space X_1 or X_2 is just one of $\binom{n_1+n_2}{n_1}$ possible assignments, all of which are arbitrary and equally likely. For each of these possible assignments, the value of $\sigma_{\chi_2}^2$ is then computed. Collectively, these values yield the *permutation distribution* for $\sigma_{\chi_2}^2$, which is analogous to a sampling distribution in the standard hypothesis testing paradigm. Finally, analogous to a standard hypothesis testing p-value, the *permutation test p-value* is obtained by calculating the proportion of values in the permutation distribution which are less than or equal to the observed value of the joint loss function. In practice, the number of possible assignments may be unreasonably large, in which case the above procedure is subtly altered to produce an *approximate permutation test p-value*. In particular, rather than using the $\binom{n_1+n_2}{n_1}$ possible assignments of the $n_1 + n_2$ point clouds to the two spaces, numerous (e.g., 1000) randomly selected permutations (i.e., “shuffles”) of the $n_1 + n_2$ point clouds are instead used where after each shuffle the first n_1 point clouds are labeled as “belonging” to space X_1 and the remaining n_2 point clouds are labeled as “belonging” to space X_2 .

If the null hypothesis of the permutation test is actually false, then we would expect the permutation test p-value to be small since the observed labeling of point clouds would be the only assignment that did not mix point clouds from both spaces. When a permutation test p-value is less than the α -level, an a priori established threshold (e.g., 0.05), the observed value of $\sigma_{\chi_2}^2$ is considered smaller than what can reasonably be explained by chance assignment of the point clouds to spaces X_1

and X_2 . The null hypothesis would then be rejected and X_1 and X_2 classified as having measurably different shape.

It is important to note that if the point clouds were not obtained via random sampling of X_1 and X_2 , then a permutation test only allows us to draw conclusions with respect to the point clouds. For instance, if the permutation test p-value is less than our threshold, then we can conclude that the shapes of the point clouds from X_1 and X_2 are measurably different; however, this conclusion cannot be generalized to X_1 and X_2 at large. As limited as such a conclusion may be, it is still informative to know that such differences exist among the point clouds, particularly when $m > 3$ and the corresponding point clouds cannot be visualized.

4. Extending hypothesis testing to three or more groups

While the methods of Section 3 are useful for determining whether or not two spaces are measurably different in a particular homological dimension, many practical applications involve more than two spaces. The cardiotocography data set considered in Section 6 is one such example. Given $s \geq 3$ spaces, suppose we have n_1 point clouds, obtained via random sampling, from space X_1 , n_2 point clouds from space X_2 , \dots , and n_s point clouds from space X_s . Analogous to before, note that in practice, for each space, a single point cloud will usually be obtained via a random sample of X_i and then partitioned, via subsampling, into n_i smaller, or less dense, point clouds. In this section we extend the methods of Section 3 to obtain a hypothesis testing procedure which can determine whether or not sufficient evidence of measurable differences in shape exists between the s spaces.

4.1. Hypotheses and justification. To conduct such an inquiry, we follow through with the suggestion of Robinson and Turner and use an approach analogous to a standard one-way ANOVA procedure in which there are potentially two stages of hypothesis testing. An omnibus (i.e., “global”) test is conducted at the first stage and if this test produces significant results, a number of post hoc (i.e., “local”) tests are performed at the second stage to identify the source(s) of the global significance. A thorough development of the one-way ANOVA procedure is available in [Casella and Berger 2002; DeGroot and Schervish 2012; Ramsey and Schafer 2013]. As with the joint loss function in Section 3, the sampling distribution of the test statistic corresponding to the omnibus test, which is presented below in Section 4.2, is nontrivial to determine and currently unknown. Hence, we again use a permutation test to carry out the omnibus test, which we will henceforth refer to as the *omnibus permutation test*. The logic behind and mechanics of this test are developed below in Section 4.2.

The null hypothesis for the omnibus permutation test asserts that the shapes of X_1, X_2, \dots, X_s are not measurably different, while the alternative hypothesis asserts that the shapes of at least two of the s spaces are measurably different. If

we fail to reject the null hypothesis of this omnibus permutation test, then we are done. However, if we reject the null hypothesis, then we know that at least two of the s spaces have shapes that are measurably different, though we do not yet know which spaces. Hence, up to $\binom{s}{2}$ post hoc tests are performed, one for each possible pairing of two of the s spaces. For each post hoc test, the null hypothesis asserts that the shapes of the two spaces are not measurably different, while the alternative hypothesis asserts that the shapes are measurably different. Thus, each post hoc test can be conducted via the methods described in Section 3.

Before describing the test statistic and corresponding details for the omnibus permutation test, note that the primary purpose of the test pertains to management of the familywise type I error rate. A type I error is the general term used to identify a hypothesis test decision in which the null hypothesis is incorrectly rejected. For any single hypothesis test, the pre-established α -level is the probability of making a type I error. When multiple post hoc tests are performed, the familywise type I error rate refers to the probability of incorrectly rejecting at least one of the corresponding null hypotheses. Many methods exist for bounding the familywise type I error rate associated with multiple pairwise post hoc tests (e.g., Bonferroni), but such methods invariably require different and smaller α -levels for each individual post hoc test. Hence, an insignificant omnibus permutation test result prevents the analyst from unnecessarily performing post hoc tests and needlessly managing the familywise type I error rate. Stated another way, if the null hypothesis of the omnibus permutation test is true, then all of the null hypotheses of the various post hoc tests are also true, and thus do not need to be performed, which eliminates any need to manage the familywise type I error rate. However, if an omnibus permutation test in which the null hypothesis is ultimately true is not performed, then $\binom{s}{2}$ post hoc tests are unnecessarily performed and the familywise type I error rate must needlessly be managed.

4.2. Omnibus permutation test specifics. Suppose, possibly after subsampling, that n_1 point clouds are available from X_1 , n_2 point clouds from X_2 , \dots , and n_s point clouds from X_s , with their corresponding persistence diagrams in a fixed dimension denoted respectively by $X_{1,1}, X_{1,2}, \dots, X_{1,n_1}$, $X_{2,1}, X_{2,2}, \dots, X_{2,n_2}$, and $X_{s,1}, X_{s,2}, \dots, X_{s,n_s}$. Analogous to the test statistic for the two-space permutation test presented in Section 3, the test statistic for the omnibus permutation test, for three or more spaces, is a function of the diagram distances for all $\binom{n_1}{2}$ pairings of persistence diagrams from X_1 , all $\binom{n_2}{2}$ pairings of persistence diagrams from X_2 , \dots , and all $\binom{n_s}{2}$ pairings of persistence diagrams from X_s . In particular, the *omnibus joint loss function* is defined as

$$\sigma_{X_s}^2 = \sum_{m=1}^s \frac{1}{2n_m(n_m - 1)} \sum_{i=1}^{n_m} \sum_{j=1}^{n_m} d(X_{m,i}, X_{m,j})^2,$$

where $d(\cdot, \cdot)$ is the persistence diagram distance metric described in Section 2.3. Analogous to $\sigma_{\chi_2}^2$, the function $\sigma_{\chi_s}^2$ is ultimately an aggregate measure of variability since the omnibus joint loss function adds the within-group variation of persistence diagram distances from each of the s spaces. As previously mentioned, the sampling distribution of $\sigma_{\chi_s}^2$ is nontrivial to determine and currently unknown; hence, we turn to the omnibus permutation test.

The logic behind and the mechanics of this omnibus permutation test are analogous to the two-space permutation test described in Section 3. We assume that the null hypothesis is true, which effectively means that the observed assignment of the point clouds to the s spaces is just one of $\prod_{i=1}^{s-1} \binom{\sum_{j=i}^s n_j}{n_i}$ possible assignments, all of which are arbitrary and equally likely. For each of these possible assignments, the value of $\sigma_{\chi_s}^2$ is then computed. Collectively, these values yield the permutation distribution for $\sigma_{\chi_s}^2$. Finally, the permutation test p-value is then obtained by calculating the proportion of values in the permutation distribution which are less than or equal to the observed value of $\sigma_{\chi_s}^2$. As in the two-space scenario of Section 3, in practice the number of possible assignments may be unreasonably large, in which case the above procedure is analogously altered to produce an *approximate permutation test p-value*. In particular, rather than using the $\prod_{i=1}^{s-1} \binom{\sum_{j=i}^s n_j}{n_i}$ possible assignments of the $n_1 + n_2 + \dots + n_s$ point clouds to the s spaces, numerous (e.g., 1000) randomly selected permutations (i.e., “shuffles”) of the $n_1 + n_2 + \dots + n_s$ point clouds are instead used where after each shuffle the first n_1 point clouds are labeled as “belonging” to space X_1 , the next n_2 point clouds are labeled as “belonging” to space X_2 , ..., and the remaining n_s point clouds are labeled as “belonging” to space X_s .

Analogous to the two-space scenario of Section 3, if the null hypothesis of this omnibus permutation test is actually false, then we would expect the permutation test p-value to be small since the observed labeling of point clouds would be the only assignment that did not mix point clouds across the s spaces. The permutation test p-value is then compared to the α -level (e.g., 0.05). If the permutation test p-value is smaller than this threshold, then the observed value of $\sigma_{\chi_s}^2$ is considered smaller than what can reasonably be explained by chance assignment of the point clouds to the s spaces. The null hypothesis would then be rejected and at least two of the s spaces are declared as having measurably different shape. To then identify the source(s) of this difference, i.e., to determine which spaces have measurably different shape, a requisite number of post hoc tests are conducted via the two-space methods of Section 3.

5. Simulation study

To confirm the two-space permutation test introduced by Robinson and Turner [2013] and to validate our proposed generalization for three or more spaces, we conducted a large-scale simulation study. Throughout the study, shape was measured

via 1-dimensional persistent homology. Three different scenarios were considered and all three consisted of three spaces ($s = 3$). For each scenario, a trial consisted of obtaining 20 point clouds, via random sampling of points, from each of the three spaces and then calculating the approximate omnibus permutation test p-value. While the 20 point clouds from a particular space were ultimately drawn independently, they can be viewed as 20 disjoint subsamples of one larger, i.e., more dense, point cloud obtained via random sampling of points of the space. All approximate omnibus permutation test p-values were based on 100,000 randomly selected permutations of the 60 collective point clouds. In particular, for each permutation, the 60 point clouds were shuffled and then labeled such that the first 20 were in the first space, the next 20 were in the second space, and the final 20 were in the third space. In the third and final scenario, each of the three possible post hoc tests were additionally performed using the two-space permutation test described in Section 3. The corresponding approximate two-space permutation test p-values were based on 100,000 randomly selected permutations of the 40 collective point clouds. In particular, for each permutation, the 40 point clouds were shuffled and then labeled such that the first 20 were in the first space and the final 20 were in the second space. A total of 100 trials were performed for each scenario and the percentage of these 100 trials that produced approximate (omnibus/two-space) permutation test p-values less than or equal to 0.05 was calculated.

5.1. *Unbalanced unit circles.* For the first scenario, each of the three spaces was the unit circle; hence, the omnibus permutation test null hypothesis that there is no measurable difference in shape between the three spaces is ultimately true. The number of sampled points making up a point cloud from each space, however, was not the same (i.e., the sample sizes are unbalanced). Each point cloud in the first space consisted of a random sample of size 18, whereas each point cloud in the second space consisted of a random sample of size 36 and each point cloud in the third space consisted of a random sample of size 54. For all three spaces, samples were obtained without allowing for measurement error; i.e., all sampled points were on their respective unit circle. Counterintuitively, 100% of the 100 trials performed produced approximate omnibus permutation test p-values less than or equal to 0.05. In fact, 100% of the trials produced approximate omnibus permutation test p-values less than 0.01. Thus, in every trial the null hypothesis would be rejected at the 5% level and we would conclude that the shapes of at least two of the three spaces are measurably different.

While such results may appear to suggest that the omnibus permutation test is ineffective, ultimately these results are an expected consequence of allowing different (i.e., unbalanced) sample sizes across point clouds. Relative to a point cloud obtained from a random sample of size 18 from the unit circle, a point cloud

obtained from a random sample of size 54 is likely to produce a persistence diagram (corresponding to homology dimension 1) containing a point that is measurably further from the diagonal. This point in the persistence diagram is expected as the circular feature within the point cloud will be born sooner and thus persist for a longer time interval. Hence, in order for the hypothesis testing methods described in Sections 3 and 4 to detect truly measurable differences in shape between the various spaces, every point cloud, both within a space and across spaces, must consist of the same number of randomly sampled data points. We will henceforth refer to this procedural necessity as balanced sampling. In practice, balanced sampling will usually be implemented at the subsampling level when the sampled data points of a space are partitioned, via subsampling, into multiple point clouds; this is demonstrated using the cardiocography data set considered in Section 6.

5.2. *Balanced samples from circles with varying radius.* For the second scenario, the three spaces were circles with radii of 1, $\frac{1}{2}$ and $\frac{1}{3}$ units. Notice that these three spaces are topologically equivalent, though geometrically different, and there is in fact a measurable difference in shape among the three spaces as measured by persistent homology in dimension 1. Hence, the null hypothesis for the corresponding omnibus permutation test is ultimately false. Point clouds for each of the three circles consisted of random samples of size 24. As in the unbalanced unit circles scenario, all samples were obtained without allowing for measurement error; i.e., all sampled points were on their respective circle. Of the 100 trials performed, 100% of them produced approximate omnibus permutation test p-values less than or equal to 0.05. In fact, as in the unbalanced unit circles scenario, 100% of the trials produced approximate omnibus permutation test p-values less than 0.01. Hence, in every trial the null hypothesis would be rejected at the 5% level and we would conclude that the shapes of at least two of the three spaces are measurably different.

As the three spaces of this second scenario are all topologically equivalent, these results suggest that the omnibus permutation test is capable of recognizing when purely geometrical differences exist between the spaces. Stated another way, this second scenario suggests that the hypothesis testing methods described in Sections 3 and 4 are not scale invariant. This is not a surprising result. More specifically, as seen in the example data of Figure 1, a point cloud obtained from a sample of points from the circle with radius $\frac{1}{3}$ will result in birth and death times for comparatively smaller radii values than a point cloud obtained from a sample of points from the unit circle. This is an artifact of the distances between neighboring points in the point cloud from the circle with radius $\frac{1}{3}$ typically being smaller than those from the unit circle. While in practice it will usually be difficult to determine whether a significant hypothesis test is a result of topological or geometrical differences between the various spaces, it is informative nonetheless to find evidence of any measurable difference in shape.

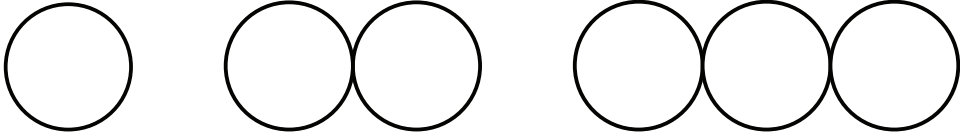


Figure 3. The three spaces of the first case of the balanced wedges simulation scenario. On the left is the unit circle, in the middle is the wedge of two unit circles, and on the right is the wedge of three unit circles.

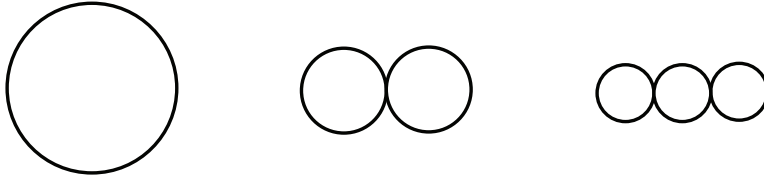


Figure 4. The three spaces of the second case of the balanced wedges simulation scenario. On the left is the unit circle, in the middle is the wedge of two circles of radius $\frac{1}{2}$, and on the right is the wedge of three circles of radius $\frac{1}{3}$.

5.3. *Balanced wedges.* The third and final scenario consisted of three distinct, but related cases in which only balanced sample sizes were considered. In the first case, the three spaces were the unit circle, the 2-wedge consisting of two unit circles, and the 3-wedge consisting of three unit circles. Hence, in this first case, the radius of every component circle is 1. An image of these three spaces is given in Figure 3. In the second case, the three spaces were the unit circle consisting of one circle of radius 1, the 2-wedge consisting of two circles of radius $\frac{1}{2}$, and the 3-wedge consisting of three circles of radius $\frac{1}{3}$. Hence, in this second case, the radii of the component circles within a space sum to 1. An image of these three spaces is given in Figure 4. In the third and final case, the three spaces were the unit circle, the unit circle with a single chord traversing the interior of the circle, and the unit circle with two nonintersecting chords traversing the interior of the circle. Hence, in this third case, the area of each of the three spaces is π units. An image of these three spaces is given in Figure 5. Observe that across these three scenarios the representations of the three spaces are topologically equivalent, but geometrically different. We consider all three scenarios since persistence diagrams are unavoidably influenced by such differences.

Within each of the three cases, the null hypothesis of the omnibus permutation test is ultimately false. In other words, there are measurable differences in shape between the three spaces. The point clouds for each of the three spaces, in all

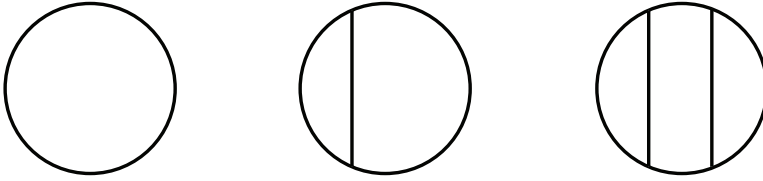


Figure 5. The three spaces of the third case of the balanced wedges simulation scenario. On the left is the unit circle, in the middle is the unit circle with a single chord, and on the right is the unit circle with two nonintersecting chords.

three cases, were obtained from random samples of the same size (i.e., balanced samples). Ten different sample sizes were considered: 6, 12, 18, 24, 30, 36, 42, 48, 54, and 60. Figure 6 provides examples of point clouds obtained from random samples of sizes 12 and 60, respectively, from each of the three spaces for the second case.

For each of these ten sample sizes, three distinct measurement errors were considered: 0 (i.e., no error), $\frac{1}{3}$, and $\frac{2}{3}$ units. For example, in the 2-wedge of the first case, measurement error was incorporated in the following manner. A random sample of points was obtained separately from each of the two unit circles of the 2-wedge. Each point on the circles was obtained by randomly selecting the angle of the point from a uniform distribution $U(0, 2\pi)$. Each point was then assigned a radius value of 1 and converted to Cartesian coordinates. Finally, for

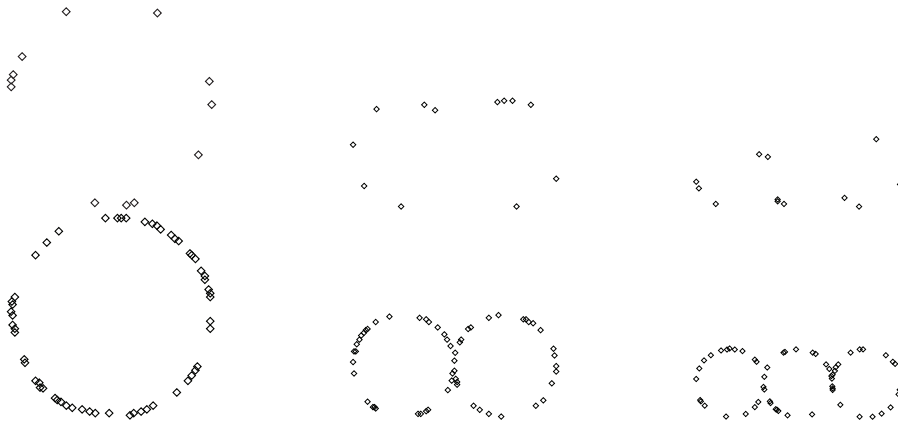


Figure 6. Point clouds obtained from random samples of each of the three spaces of the second case of the balanced wedges simulation scenario. The first row contains point clouds obtained from random samples of size 12. The second row contains point clouds obtained from random samples of size 60.

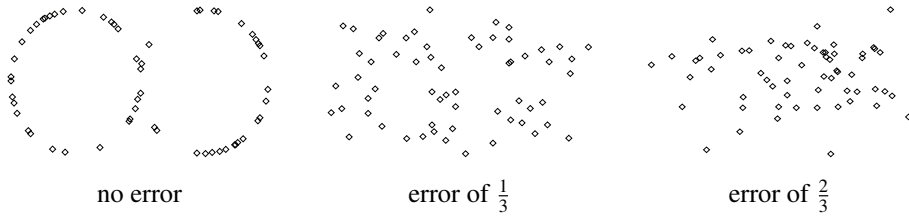


Figure 7. Point clouds obtained from random samples of size 60, under various measurement errors, from the 2-wedge of the first case of the balanced wedges simulation scenario.

each point, two errors were randomly sampled from a normal distribution $\mathcal{N}(0, \sigma)$, where σ is the specified measurement error (e.g., $\frac{1}{3}$), and respectively added to the Cartesian coordinates of the point. For each of the three measurement errors, Figure 7 exemplifies a point cloud obtained from a sample of size 60 from the 2-wedge. From these images it is clear that as the measurement error increases, the extent to which the point cloud resembles the 2-wedge dramatically decreases. Measurement errors for the other spaces of the second case, as well as for the other cases of the third scenario, was analogously incorporated.

For each of the 30 combinations of sample size and measurement error, the percentage of the 100 trials producing an approximate omnibus permutation test p-value less than or equal to 0.05 for case one is given in Table 1. Two trends are readily apparent from these results. First, as sample size increases for a fixed measurement error, the percentage of significant omnibus permutation test results almost uniformly increases. This is intuitive and desirable since we would expect measurable differences in shape between the three spaces to become more easily identifiable as sample size increases. Second, as measurement error increases for a fixed sample size, the percentage of significant omnibus permutation test results almost uniformly decreases. This too is intuitive and desirable since we would expect measurable differences in shape between the three spaces to become less easily identifiable as measurement error increases. Given these trends and the fact that there are so many entries in the table at or near 100%, these results suggest that the proposed omnibus permutation test successfully identified measurable differences in shape between at least two of these three spaces. The results for the second and third cases, depicted in Figures 4 and 5, are analogous to those above for the first case and, therefore, are omitted.

As the omnibus permutation test successfully identified measurable differences in shape between at least two of the three spaces, in all three cases, each of the three possible post hoc tests were then conducted. For each such post hoc test, the null hypothesis asserts that there is no measurable difference in shape between the two spaces, while the alternative hypothesis asserts the opposite. Hence, in all

sample size	noise		
	0	$\frac{1}{3}$	$\frac{2}{3}$
6	6%	9%	1%
12	95%	57%	18%
18	100%	65%	41%
24	100%	96%	41%
30	100%	100%	85%
36	100%	100%	98%
42	100%	100%	100%
48	100%	100%	100%
54	100%	100%	100%
60	100%	100%	100%

Table 1. Balanced unit wedges — results of omnibus permutation tests. For each combination of sample size and measurement error, the percentage of approximate omnibus permutation test p-values (out of 100) yielding a value less than or equal to 0.05 is given. The three spaces are the unit circle, the 2-wedge and the 3-wedge.

three tests, for all three cases, the null hypothesis is ultimately false. As the results across the three cases were ultimately analogous, only the results for the first case are discussed below. In particular, for each of the 30 combinations of sample size and measurement error, the percentage of the 100 trials producing an approximate post hoc test p-value less than or equal to 0.05 is given in Table 2 for the circle versus the 2-wedge, in Table 3 for the circle versus the 3-wedge, and in Table 4 for the 2-wedge versus the 3-wedge.

The two trends that were apparent in the corresponding omnibus permutation tests for this simulation scenario are also readily apparent in all three of these post hoc tests. Specifically, as sample size increases for a fixed measurement error, the percentage of significant post hoc tests tends to increase. Similarly, as measurement error increases for a fixed sample size, the percentage of significant post hoc tests tends to decrease. A cell-by-cell comparison of the percentages among the three post hoc tests, however, reveals an additional interesting trend. The percentages for the post hoc test between the circle and the 3-wedge are almost uniformly larger than or equal to the corresponding percentages between the circle and the 2-wedge, which are in turn almost uniformly larger than or equal to the corresponding percentages between the 2-wedge and the 3-wedge. This too is mostly intuitive and desirable since, among the three spaces, the unit circle and the three wedge are the most different with respect to shape. We are uncertain why the post hoc test appears more adept at recognizing measurable differences

sample size	noise		
	0	$\frac{1}{3}$	$\frac{2}{3}$
6	2%	5%	2%
12	90%	29%	13%
18	99%	40%	15%
24	100%	83%	28%
30	100%	97%	49%
36	100%	100%	64%
42	100%	100%	80%
48	100%	100%	82%
54	100%	100%	92%
60	100%	100%	97%

Table 2. Balanced wedges, first case—results of unit circle vs. 2-wedge post hoc tests. For each combination of sample size and measurement error, the percentage of approximate two-space permutation test p-values (out of 100) yielding a value less than or equal to 0.05 is given.

sample size	noise		
	0	$\frac{1}{3}$	$\frac{2}{3}$
6	2%	5%	1%
12	97%	65%	30%
18	100%	85%	40%
24	100%	100%	53%
30	100%	100%	95%
36	100%	100%	100%
42	100%	100%	100%
48	100%	100%	100%
54	100%	100%	100%
60	100%	100%	100%

Table 3. Balanced wedges, first case—results of unit circle vs. 3-wedge post hoc tests. For each combination of sample size and measurement error, the percentage of approximate two-space permutation test p-values (out of 100) yielding a value less than or equal to 0.05 is given.

in shape between the circle and the 2-wedge rather than between the 2-wedge and the 3-wedge. Regardless, all three of these trends, when coupled with the volume of entries in all three tables which are at or near 100%, indicate that the proposed

sample size	noise		
	0	$\frac{1}{3}$	$\frac{2}{3}$
6	0%	1%	1%
12	4%	17%	13%
18	62%	16%	18%
24	86%	33%	14%
30	93%	42%	20%
36	87%	66%	26%
42	95%	67%	43%
48	99%	87%	65%
54	100%	93%	66%
60	100%	98%	84%

Table 4. Balanced wedges, first case—results of 2-wedge vs. 3-wedge post hoc tests. For each combination of sample size and measurement error, the percentage of approximate two-space permutation test p-values (out of 100) yielding a value less than or equal to 0.05 is given.

post hoc tests successfully identified measurable differences in shape between each of the three possible pairings of these three spaces. Such findings additionally corroborate the legitimacy of the two-space permutation test.

5.4. Summary of findings. In summary, the major findings of the simulation study are three-fold. First and foremost, these simulations demonstrate that the proposed omnibus permutation testing procedure successfully identified measurable differences in shape between at least two of the three spaces. Second, these simulations confirm that the post hoc testing component successfully identified measurable differences in shape between any two spaces; such findings corroborate the legitimacy of the two-space permutation testing procedure. Third and finally, these simulations reveal that, for any number of spaces, balanced sampling is required in obtaining the point clouds utilized in the testing procedure.

6. Applications to real data sets

We apply our methods to the cardiocography (CTG) data set that is freely available from the University of California at Irvine Machine Learning Repository at <https://archive.ics.uci.edu/ml/datasets/cardiocography>. The CTG data set includes 23 variables for each of 2126 subjects. We apply our methods on a focused subset of four quantitative variables, including fetal heart rate baseline in beats per minute, number of accelerations per second, number of uterine contractions per second,

and number of light decelerations per second. These four quantitative variables are chosen because they are seemingly independent, and we want to consider no more than four such variables. The categorical variable of interest is health status, which has three levels: normal, suspect, and pathologic. The question of interest is whether or not the 4-dimensional space created by the quantitative variables has a measurably different shape across the three health status groups. To answer this question, we use the omnibus permutation testing procedure developed in Section 4.1, measuring shape via 1-dimensional persistent homology. Before this procedure can be performed, however, multiple point clouds from the three health status groups must be obtained via balanced subsampling of the subjects.

Of the 2126 sampled subjects, 1655 are of normal health status, 295 are of suspect health status, and 176 are of pathologic health status. Hence, from the sampled data points we obtain three 4-dimensional point clouds, one consisting of 1655 subjects from the normal health status group, another consisting of 295 subjects from the suspect health status group, and one other consisting of 176 subjects from the pathologic health status group. As our methods require balanced sampling across multiple point clouds from each of the groups, we partitioned, via subsampling, each given point cloud into smaller 4-dimensional point clouds consisting of 44 subjects each. Consequently, we obtained 37 point clouds from the normal health status group, 6 point clouds from the suspect health status group, and 4 point clouds from the pathologic health status group. As neither 1655 nor 295 are divisible by 44, we simply discarded the leftover 27 normal health status subjects and the 31 suspect health status subjects.

The omnibus permutation test was then performed using the persistence diagrams corresponding to the 47 subsampled point clouds. The null hypothesis asserted that there were no measurable differences in shape between the three spaces corresponding to the three health status groups. The resulting approximate permutation test p-value of 0.00005 was based on 100,000 randomly sampled permutations of the 47 point clouds. In particular, for each permutation, the 47 point clouds were shuffled and then labeled such that the first 37 were in the normal health status group, the next 6 were in the suspect health status group, and the last 4 were in the pathologic health status group. Given that the p-value is so small, we reject the null hypothesis and conclude that there are measurable differences in shape between at least two of the three spaces.

To determine the source(s) of the difference, we ultimately performed three post hoc tests, one for each possible pairing of the three health status groups. For each such test, the null hypothesis asserted that there were no measurable differences in shape between the two spaces of the respective health status groups. For the normal and suspect health status groups, the approximate permutation test p-value of 0.00009 was based on 100,000 randomly sampled permutations of the 43 point

clouds. In particular, for each permutation, the 43 point clouds were shuffled and then labeled such that the first 37 were in the normal health status group and the final 6 were in the suspect health status group. For the normal and pathologic health status groups, the approximate permutation test p-value of 0.0060 was based on 100,000 randomly sampled permutations of the 41 point clouds. In particular, for each permutation, the 41 point clouds were shuffled and then labeled such that the first 37 were in the normal health status group and the final 4 were in the pathologic health status group. Finally, for the suspect and pathologic health status groups, the approximate permutation test p-value of 0.3012 was based on 100,000 randomly sampled permutations of the 10 point clouds. In particular, for each permutation, the 10 point clouds were shuffled and then labeled such that the first 6 were in the suspect health status group and the final 4 were in the pathologic health status group. Note that while (exact) permutation test p-values could have straightforwardly been obtained for the post hoc tests involving normal versus pathologic (101,270 possible assignments) and suspect versus pathologic (210 possible assignments), such p-values could not have reasonably been obtained for the post hoc test involving normal versus suspect (6,096,454 possible assignments) or for the omnibus test (1.087394×10^{12} possible assignments); therefore, for the sake of consistency, approximate permutation test p-values were obtained in all instances. Based on these results, there is significant evidence of measurable differences in shape between the spaces corresponding to the normal and suspect health status groups, and between the normal and pathologic health status groups, but insignificant evidence of such differences between the suspect and pathologic health status groups.

7. Conclusion

For multiple point clouds obtained from (sub)sampled points of three or more spaces, we propose using an omnibus permutation test on the corresponding persistence diagrams to determine whether statistically significant evidence exists of measurable differences in shape between any of the respective spaces. If such differences do exist, we then propose using a number of post hoc (i.e., two-space) permutation tests to identify the specific pairwise differences. To validate this proposed procedure, we conducted a large-scale simulation study using point clouds obtained from samples of points from three spaces. Various combinations of spaces, sample sizes and measurement errors were considered in the simulation study and for each combination the percentage of p-values below an α -level of 0.05 was provided. The results of the simulation study clearly suggest that the procedure works, but additionally reveal that the method is neither scale invariant nor insensitive to imbalanced sample sizes across point clouds. Finally, we applied our omnibus testing procedure to a cardiocography data set and found statistically significant

evidence of measurable differences in shape between the spaces corresponding to normal, suspect and pathologic health status groups.

While the proposed omnibus testing procedure is applicable in any homological dimension, the simulation study and CTG application presented in this paper focus exclusively on homological dimension 1. Hence, to validate the effectiveness of the method in other homological dimensions, and to assess the consistency of the method across various dimensions, additional simulation studies can be performed.

Appendix

For readers that are less familiar with simplicial complexes, homology, and persistence diagrams, we include here examples of each for a small accessible example. Consider the set, D , of five points in the plane as pictured in Figures 8 and 9. Each point in D is a 0-simplex, each line segment drawn between points is a 1-simplex, and each shaded triangle a 2-simplex. As the parameter r increases beyond $r = 4$ the Vietoris–Rips complex will contain additional 2-simplices, a 3-simplex at $r = 4.9$, and eventually a 4-simplex when $2r$ is equal to the diameter of D . Note that the abstract simplicial complex $\text{VR}(D, 4.9)$ in Figure 9 cannot be geometrically realized in \mathbb{R}^2 since it contains pairs of 2-simplices whose intersection is not a face of either simplex.

The complex $\text{VR}(D, 4)$, on the left in Figure 9, is labeled with an ordering assigned to its 0, 1, and 2-simplices: the five 0-simplices, v_1, v_2, v_3, v_4, v_5 ; six 1-simplices $e_1, e_2, e_3, e_4, e_5, e_6$; and one 2-simplex f_1 .

With respect to this notation, the boundary of a chain is relatively easy to calculate. For example, $\delta_1(e_6 + e_1 + e_2) = v_5 + v_3$ and $\delta_2(f_1) = e_2 + e_3 + e_4$. More precisely, the chain complex of $\text{VR}(D, 4)$ is

$$0 \longrightarrow \mathbb{Z}_2 \xrightarrow{\delta_2} (\mathbb{Z}_2)^6 \xrightarrow{\delta_1} (\mathbb{Z}_2)^5 \xrightarrow{\delta_0} 0,$$

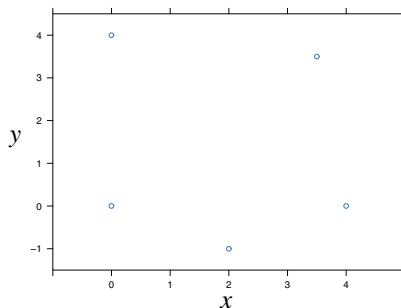


Figure 8. Five data points in the plane.

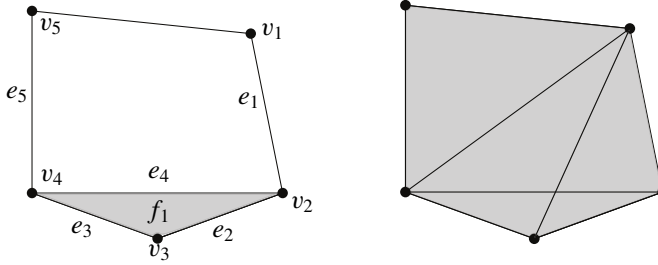


Figure 9. Representations of the abstract simplicial complexes $\text{VR}(D, 4)$ and $\text{VR}(D, 4.9)$ for the five point data set D .

with boundary maps given in matrix form by

$$\delta_2 = \begin{bmatrix} 0 \\ 1 \\ 1 \\ 1 \\ 0 \\ 0 \end{bmatrix}, \quad \delta_1 = \begin{bmatrix} 1 & 0 & 0 & 0 & 0 & 1 \\ 1 & 1 & 0 & 1 & 0 & 0 \\ 0 & 1 & 1 & 0 & 0 & 0 \\ 0 & 0 & 1 & 1 & 1 & 0 \\ 0 & 0 & 0 & 0 & 1 & 1 \end{bmatrix}, \quad \text{and} \quad \delta_0 = [0 \ 0 \ 0 \ 0 \ 0].$$

Intuitively, the p -th homology group measures equivalence classes of p -cycles of K that are not “filled” by $(p+1)$ -chains. In homological dimension $p = 1$ for the complex $\text{VR}(D, 4)$, an example of a 1-cycle that is not the boundary of a 2-cycle is $e_1 + e_2 + e_3 + e_5 + e_6$. Hence this 1-cycle is in a nonzero equivalence class of $H_1(\text{VR}(D, 4); \mathbb{Z}_2)$. The 1-cycle $e_2 + e_3 + e_4$, however, is the boundary of the 2-cycle f_1 (this 1-cycle is “filled” by f_1), so this 1-cycle is equivalent to zero in the homology group. Hence, in dimension $p = 1$, the homology of $\text{VR}(D, 4)$ is measuring the circular hole that is seen in the complex.

To complete the homology calculation for the simplicial complex $\text{VR}(D, 4)$, we see that the kernel of δ_0 is $(\mathbb{Z}_2)^5$ and the rank of δ_1 is 4. Thus $H_0(\text{VR}(D, 4); \mathbb{Z}_2) \cong \mathbb{Z}_2$. Similarly, the nullity of δ_1 is 2 and the image of δ_2 is 1-dimensional. This implies that $H_1(\text{VR}(D, 4); \mathbb{Z}_2) \cong \mathbb{Z}_2$. We have $H_2(\text{VR}(D, 4); \mathbb{Z}_2) \cong 0$, since the kernel of δ_2 is 0. Because the complex contains no simplices in higher dimensions, $H_p(\text{VR}(D, 4); \mathbb{Z}_2) = 0$ for all $p > 2$.

The calculation $H_0(\text{VR}(D, 4); \mathbb{Z}_2) = \mathbb{Z}_2$ measures that $\text{VR}(D, 4)$ is a connected complex. The nontrivial group $H_1(\text{VR}(D, 4); \mathbb{Z}_2) = \mathbb{Z}_2$ measures the existence of a 1-dimensional cycle that is not the boundary of a 2-simplex, namely $e_1 + e_2 + e_3 + e_5 + e_6$.

For the complex $\text{VR}(D, 4.9)$, on the right in Figure 9, the homology groups are $H_0(\text{VR}(D, 4.9)) = \mathbb{Z}_2$ and $H_p(\text{VR}(D, 4.9)) = 0$ for all $p \geq 1$. In this example, the

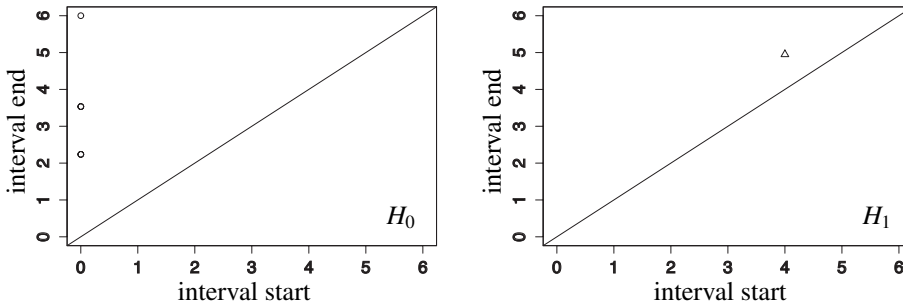


Figure 10. The persistence diagrams corresponding to the five-point data set in Figure 8 in the homological dimensions 0 and 1.

first homology group disappeared, or died, as r increases from 4 to 4.9 as a result of the additional 2-simplicies that span the 1-cycle $e_1 + e_2 + e_3 + e_5 + e_6$.

The persistence diagrams in Figure 10 display the H_0 and H_1 persistence diagrams for the five-point data set D first seen in Figure 8. Note that all points in a persistence diagram are plotted above the line $y = x$, as a persistent homology class must be born before it can die.

In homological dimension 1 (the H_1 diagram), the small triangle plotted at the point $(4, 4.9)$ indicates that the five-point data set contains a 1-dimensional homology class that is born at radius 4 and dies at radius 4.9. In homological dimension 0 (the H_0 diagram), the circles plotted at the points $(0, 2.236)$ and $(0, 3.54)$ represent the connection of data points by 1-simplices at $r = 2.236$ and at $r = 3.54$ resulting in the death of a connected component when it is joined with another connected component by a 1-simplex. For $r > 3.54$ the five points are path connected via 1-simplices; thus this connected complex gives rise to a single 0-dimensional persistent homology class. This single class is plotted at $(0, 6)$ as a result of considering only r -values in the range $0 \leq r \leq 6$.

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
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