SPCS: A Spatial and Pattern Combined Smoothing Method for Spatial Transcriptomic Expression

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Abstract

The recently developed spatial transcriptomics (ST) technique has made it possible to view spatial transcriptional heterogeneity in a high throughput manner. It is based on highly multiplexed sequence analysis and uses barcodes to split the sequenced reads into respective tissue locations. However, this type of sequencing technique suffers from high noise and drop-out events in the data, which makes smoothing a necessary step before performing downstream analysis. Traditional smoothing methods used in the similar single cell RNA sequencing (scRNA-seq) data are one-factor methods that can only utilize associations in transcriptome space. Since they do not account for associations in the Euclidean space, i.e. tissue location distances on the ST slide, these one-factor methods cannot take full advantage of all the knowledge in ST data. In this study, we present a novel two-factor smoothing technique, Spatial and Pattern Combined Smoothing (SPCS), that employs k-nearest neighbor technique to utilize associations from transcriptome and Euclidean space from the ST data. By performing SPCS on multiple ST slides from pancreatic ductal adenocarcinoma (PDAC), dorsolateral prefrontal cortex (DLPFC) and simulated high-grade serous ovarian cancer (HGSOC) datasets, smoothed ST slides have better separability, partition accuracy, and biological interpretability than the ones smoothed by pre-existing one-factor methods. Source code of SPCS is provided in Github (<https://github.com/Usos/SPCS>).

Keywords

Spatial transcriptomics, imputation, two-factor expression smoothing, k-nearest neighbors, tissue region partition, pancreatic ductal adenocarcinoma, dorsolateral prefrontal cortex, high-grade serous ovarian cancer

58 Introduction

59 Tumors and organ tissues are highly heterogeneous with different phenotypes distributed
60 throughout the tissue [1, 2]. Until recently, this tissue heterogeneity had to be evaluated
61 using histology image analysis or using scRNA-seq. Histology imaging offers a view on
62 spatial tissue heterogeneity whereas scRNA-seq offers insight into the transcriptional
63 heterogeneity. Methods like immunohistochemistry (IHC) labelled histology slides also
64 allow researchers to view protein in tissue images [3-7]. However, IHC is only capable of
65 capturing protein expression level(s) of one or a few proteins at a time. These technologies
66 lacked the ability to view transcriptional heterogeneity spatially in a high throughput
67 manner.

68 Fortunately, revolutionary new techniques that can be broadly categorized as spatial
69 transcriptomics (ST) have recently become available that allow transcriptome profiles to
70 be measured for different locations in a tissue [8, 9]. These techniques have already been
71 profoundly useful in understanding tumor heterogeneity [10-12] and heterogeneity of brain
72 tissues [13-16]. In recent years, research on this new ST technology has been prolific --
73 in 2017 there were only five articles mentioning “spatial transcriptomics” in PubMed
74 (<https://pubmed.ncbi.nlm.nih.gov/>), but by 2020 there were a total of 70 articles, 14 times
75 as many as just three years earlier. However, ST technology is still largely being studied
76 methodologically to improve the library preparation [17], sequencing techniques, and
77 bioinformatic analysis pipelines [18] used in spatial transcriptomic analysis. In comparison,
78 scRNA-seq analysis has now largely established standard practice protocols [19, 20] such
79 that the data preprocessing and initial bioinformatic analyses are consistent between
80 experiments. Once ST techniques has established the standard practice, like single cell
81 technologies, then they will be widely available for any number of disease and normal

tissues.

The most common form of ST data to date, much like the now widely abundant scRNA-seq data, is based on highly multiplexed sequence analysis where barcodes are used to sort the sequenced reads into their respective cells or tissue locations [21]. This type of sequencing suffers from a wide array of noise and drop-out events in the data which has been well studied across scRNA-seq methodologies [22]. In barcoding-based single cell data, a single cell's expression profile on its own is not perfectly representative of the true underlying RNA abundances from that cell due to the noise introduced during the sequencing process. There are techniques such as Smart-seq2 [23] and CEL-seq2 [24] that give more representative expression profiles per cell, but these are much lower throughput, resulting in many fewer cells. As a result, a multitude of techniques have been developed to impute the missing expression values and smooth the noise that come directly from the barcode-based scRNA-seq. These techniques help to smooth the data by using similar cells to correct for technical errors in the sequencing data. Specifically, sets of correlated genes and relative cell similarity in transcriptome space is used to impute the drop-out events and eliminate other types of expression errors via machine learning techniques. Some of these tools include SAVER [25] and MAGIC [26]. When applied, the processed expression values give more accurate representations of the true underlying RNA abundances than the raw read counts.

Because most current ST data are also generated from barcode-based technology, it is not surprising that ST data also suffer from similar noise and drop-out events. This means that the distributions of gene expression in the tissue does not always follow the corresponding phenotypic differences in histopathological images at a high resolution. For drop-out events, it can be assumed that zero expression values have some probability of being spuriously missed in the reads simply due to chance. However, ST data, unlike

scRNA-seq, not only has relative similarity in transcriptome space but also in Euclidean space on the tissue slide. That means locations that are adjacent to one another in the ST array will likely be similar tissue and as a result have similar transcriptome profiles. In addition, if two locations in an ST array share a majority of their transcriptome profiles, then their transcriptional differences can be partially explained by drop-out.

Here we present a novel two-factor ST smoothing method that utilizes both the associations of spatial locations in transcriptome space (expression pattern knowledge) and in Euclidean space (spatial knowledge). This is in contrast to the pre-existing one-factor scRNA-seq techniques that only utilize associations in transcriptome space. We believe that by using both the spatial location of a given ST spot and its transcriptome profile we can better smooth the data such that it better represents the true underlying RNA abundances in the tissue.

Methods

Datasets

The datasets that we use in this study including two real world ST dataset, PDAC [10] and DLPFC [27], and a simulating dataset generated from HGSOC single-cell dataset [28]. For the two real world datasets, PDAC include 10 ST slides sourced from traditional ST platform, while DLPFC is a Visium platform dataset with 12 slides. All the data in these datasets consist of two different matrices containing gene is expressions and spatial coordinates. One matrix consists of the gene expression values for each spatial barcode hybridized from its corresponding spot on the ST slide. The other matrix contains the spatial locations in 2D space for each spot's spatial barcode. Using these two matrices, we can generate a 2D representation of each gene's expression value throughout the

biopsied tissue section. Because these datasets sourced from different ST platform, i.e. traditional ST platform and new developed Visium platform, we can explore the influence of smoothing methods more comprehensively. A detailed summary statistics of the data we used is provided in Table S1.

To better explore the ability of smoothing methods on deal with outlier spots, we designed a simulation experiment referred to BayesSpace [29]. Simulating ST data is based on HGSOc single-cell dataset and an immunofluorescence stain image associated with ovarian cancer (OC). In the original single-cell analysis of HGSOc dataset, all the cells were divided into 15 clusters by DBSCAN clustering method and well annotated [28]. Considering the limitation of number of cells, we used only partial of the slide. Ground-truth cluster labels were derived from single-cell level annotation of tumor and stroma compartments within the image. To make simulated data reflect the real nature, we separated the slide into four clusters, including intratumor, stroma and two tumor clusters. Among the spatial clusters, intratumor cluster including dendritic cell and fibroblast cell clusters, stroma cluster corresponded to macrophage cell cluster and tumor clusters associated to two different malignant cell clusters. Detailed information about ground truth clusters is provided in Table S2. For the propose of testing the anti-outlier spot ability of different smoothing methods, we randomly mixed 5% of other types of cells as perturbation for each spatial cluster. We generated 10 groups of simulating data in the simulation analysis.

Spatial and Pattern Combined Smoothing (SPCS) of Spatial Transcriptomic Expression

For each spot on an ST slide, there exists not only the gene expression but also their spatial positions. This means we can improve the quality of the expression values within

each specific spot using the relative similarity to the other spots based on expression pattern and spatial location on the ST slide. In addition, spatial positions can also help to fill the missing spots. To achieve this goal, we put forward a KNN-based method, Spatial and Pattern Combined Smoothing (SPCS), to perform the smoothing and padding. We display the procedure of SPCS method in Figure 1.

Figure 1 placement

In our method, we obtained the smoothed expression of each spot by integrating the contribution-weighted expression of its pattern and spatial neighbors. Let X_i be a vector of gene expression values for spot i , smoothed expression X'_i can be calculated by:

$$X'_i = (1 - \alpha)X_i + \alpha \left(\beta \frac{\sum_{j \in N_s(i)} c_{ji}^s X_j}{\sum_{k \in N_s(i)} c_{ki}^s} + (1 - \beta) \frac{\sum_{j \in N_p(i)} c_{ji}^p X_j}{\sum_{k \in N_p(i)} c_{ki}^p} \right) \quad (1)$$

In eq. (1), there are two ratio parameters α and β . α is used to regularize the ratio of original expression to the corrected expression, which avoids the expression of object spot becoming over-smoothed. β is used to adjust the ratio of spatial-based to pattern-based smoothing for different applications. $N_s(i)$ and $N_p(i)$ are deleted spatial and pattern neighborhood of object spot i respectively. Unless otherwise stated, the neighborhood in this paper is deleted neighborhood. The size of neighborhoods can be determined by parameter τ_s and τ_p which also should be specified in advance. c_{ji}^s and c_{ji}^p are spatial and pattern smoothing contribution of corresponding neighbor spot j of object spot i . For traditional ST platform based PDAC dataset, we set $\alpha = 0.6, \beta = 0.4, \tau_s = 2$ and $\tau_p = 16$; and for the Visium platform based DLPFC and simulating datasets, we make $\tau_s = 4$ due to the more number of spots. We will display the influence of these parameters on data separability and discuss the selection of them in the Discussion section. In addition, SPCS

will also fill the missing spots with multiple non-missing spatial neighbors. Source code of our proposed SPCS method is provided in Github (<https://github.com/Usos/SPCS>). In the next section, we introduce detailed mathematical definitions of neighborhood, contribution and our missing spots padding strategy.

Pattern neighborhood

ST data is a type of transcriptomic data, which means it includes expression patterns similar to scRNA-seq. Like some scRNA-seq data, where cells can be localized to tissue location of origin, spots which have a similar expression pattern are more likely to belong to the same region in a tissue. Therefore, smoothing the expression of a spot with those of similarly expressing spots can improve data quality and is common in traditional one-factor smoothing methods [25, 26]. A group of the most similar spots based on the expression “pattern” of the spot can be defined as that spot’s “pattern neighborhood”. Here, we provide the explicit definition of the pattern neighborhood used by the SPCS method.

Definition 1 (Pattern neighborhood): S_p is gene expression pattern space of a ST slide, $i, j, k \in S_p$ are different spots, d_{ij}^p and d_{ik}^p are pattern distance between spots i, j and i, k respectively. $N_p(i)$ is τ_p -deleted pattern neighborhood of spot i if $|N_p(i)| = \tau_p, \forall j \in N_p(i), \forall k \in S_p - (N_p(i) \cup \{i\})$ s.t. $d_{ij}^p < d_{ik}^p$.

For gene expression data, the overall shapes of gene expression patterns are of greater interest than the individual magnitudes of each feature [30]. Hence, we used Pearson correlation distance to measure pattern distance between different spots. Let ρ_{ij} represent Pearson correlation coefficient (PCC) of coordinate of spots i and j in pattern space, Pearson correlation distance d_{ij} of spots i and j is [31]:

$$d_{ij} = 1 - \rho_{ij} \quad (2)$$

In ST data, some genes are expressed at identical or near-identical levels, which lack of variance to establish an accurate pattern neighborhood. Therefore, we used principal component analysis (PCA) [32], to transform the expression of spots into a 10-dimensional principal component (PC) space before smoothing, to extract uncorrelated components with the largest variance. We named this PC space as gene expression pattern space.

Spatial neighborhood

In contrast to scRNA-seq data, ST data can provide spatial position for each spot in the slide. Regions in proximity on histopathology slides are usually the same tissue type. Aside from the pattern associations between spots, we can also use spatial associations to smooth the expression as a second factor. We named the group of spots that are spatially near a given spot the “spatial neighborhood” of that spot, which is defined explicitly below.

Definition 2 (Spatial neighborhood): S represents the set of spatial location indices of an ST slide, $i, j \in S$, d_{ij}^S is the spatial distance between spot i and j . $N_s(i)$ is τ_s -deleted spatial neighborhood of spot i , if $\forall j \in N_s(i)$ s.t. $d_{ij}^S \leq \tau_s$.

ST spots are spatially distributed in a checkerboard or honeycomb pattern. Manhattan distance is a suitable metric to measure the spatial distance between spots inside an ST slide. Thus, we chose Manhattan distance as the spatial distance to define our spatial neighborhood. Due to the difference of spatial pattern of spots, the shape of spatial neighborhood could be different in different platform. Figure 1(b) illustrated the shape of second-order neighborhood of both traditional ST platform (checkerboard spatial pattern) and Visium platform (honeycomb spatial pattern).

Contribution of neighbors on smoothing

Different neighbors in the spatial or pattern neighborhood, will have different impacts on the smoothing for a given spot, which we refer as “contribution”. Since the definitions of spatial and pattern distance are different, we model the corresponding contributions in different ways. The contributions of both spatial and pattern neighbors are still comparable since the range of both is $[0,1]$. For spots outside the neighborhood (both spatial and pattern) of object spot, we assigned their corresponding contribution to 0, which means they have no contribution to smoothing of the object spot.

In pattern space, to better capture global gene expression patterns, we used Pearson correlation coefficient distance described in eq. (2) as distance metric, whose range is $[0,2]$. If the expression of two spots has a negative correlation, the distance based on eq. (2) will become greater than 1. However, smoothing with negative correlation spots is not performed since they are dissimilar. Therefore, we set the contribution to 0 if pattern distance between the object spot and one of its neighbors is greater than 1. We used an exponential transformation to achieve this goal. For object spot i and its pattern neighbor j , pattern contribution c_{ji}^p can be defined as:

$$c_{ji}^p = \begin{cases} \exp\left(-\left(\frac{d_{ij}^p}{\sigma}\right)^2\right) & d_{ij}^p < 1 \\ 0 & \text{otherwise} \end{cases} \quad (3)$$

The exponential function in eq. (3) limits the range of c_{ji}^p to $[0,1]$ and ensures c_{ji}^p decreases as d_{ij}^p increases. σ in this equation is a parameter to tune the initial position of exponential function descending. When $d_{ij}^p > \frac{3\sigma}{\sqrt{2}}$, c_{ji}^p will quickly decay to 0 [33, 34]. Hence, setting σ to $\frac{\sqrt{2}}{3}$ will cease the effect of negative correlation neighbors and we can simplify the eq. (3)

242 as:

$$243 \quad c_{ji}^p = e^{-4.5(d_{ij}^p)^2} \quad (4)$$

244 For spatial neighbors, we used Manhattan distance, an integer greater than 0, as spatial
245 distance to measure their distance to object spot. Since the inverse proportional function
246 has a similar decay nature as exponential function, we used the inverse of spatial distance
247 as the spatial contribution, i.e.:

$$248 \quad c_{ji}^s = \frac{1}{d_{ij}^s} \quad (5)$$

249 In this case, i is the spot being smoothed and j is the spatial neighbor of spot i .

250 **Padding of missing spots**

251 For ST slides, there are two types of missing values. The first type of missing values is
252 gene level missing, which means the spot itself is located in tissue region, but some
253 specific gene expression is missing. This is the main form of drop-out events and also
254 frequently happen in single-cell data, which most of the smoothing methods can handle.
255 The second type is spot level missing, i.e. expression of all genes in the spot is 0. This
256 kind of missing is unique in ST data and cannot be padded without spatial position
257 information, which often indicates there's no tissue on the spot or the spot has been
258 removed due to the quality problem. Hence, it is necessary to judge whether the missing
259 spot need to be padded. As shown in Figure 1(c), for each blank spot in a slide, SPCS will
260 only pad the ones whose pre-determined spatial neighborhood has more than 50% non-
261 blank spots. This criteria ensures that boundary of the tissue will not be errorly expanded.
262 Since there's no expression on missing spots at all, we estimate the expression of missing
263 spots by their spatial neighbors only. Let i become a missing spot, its expression X_i can

264 be estimated by:

$$265 \quad X_i = \frac{\sum_{j \in N_s(i)} c_{ji}^s X_j}{\sum_{k \in N_s(i)} c_{ki}^s} \quad (6)$$

266 In Eq.(6), $N_s(i)$ is τ_s -order spatial neighborhood of spot i , c_{ji}^s and c_{ki}^s are spatial
 267 contribution of spot j and k to spot i . Spot padding is performed after non-missing spot
 268 smoothing. To evaluate ARI score for padding spots, we assign ground truth labels for
 269 them. The ground truth labels of the padded spots are determined by spatial contribution
 270 weighted major voting of their spatial neighbors.

271 Performance evaluation

272 To evaluate the effectiveness of our proposed SPCS method, we first performed SPCS
 273 and other one-factor smoothing methods (SAVER and MAGIC) on PDAC, DLPFC and
 274 simulated ST datasets. SAVER and MAGIC are two representative one-factor smoothing
 275 methods on different technical routes (i.e., statistical model-based and kNN-based). Then
 276 we partitioned both smoothed and the original unsmoothed real world slides using K-
 277 medoids clustering method [35] and judged how well the clusters can be separated after
 278 performing multiple smoothing methods by internal evaluation of the unsupervised
 279 clustering. In addition, we also explored how the parameters in SPCS will influence the
 280 smoothing. After all, we performed multiple unsupervised clustering methods, including K-
 281 medoids, Louvain [36], mclust [37], BayesSpace [29] and spaGCN [38], on all the
 282 smoothed and unsmoothed slides to find out how well the clusters match the
 283 histopathological labels from the corresponding image as an external evaluation.
 284 Simulating data also was used here to detect the ability of anti-outliers for each smoothing
 285 method. As a gene filter, we only kept genes with less than 70% zero expressed spots in
 286 our analysis. For normalization, we performed logarithmic count per million (CPM)

normalization before smoothing using SPCS, SAVER, and MAGIC. The distance metric used during clustering was Pearson correlation distance as described in eq. (2). For each dataset, number of clusters is pre-determined by ground truth which is provided in Table S1-S2. PCA was performed prior to clustering, then we selected the eigenvectors with top 20 eigenvalues to reduce the dimensions of expression matrices and enhance the clustering results.

Internal evaluation

For the internal evaluation, silhouette score was used as the evaluation indicator [39]. Silhouette score, a metric whose range is $[-1, 1]$, estimates the average distance between clusters. A greater silhouette score indicates better cluster separations. For a ST spot i , let $a(i)$ be the average distance between i and all the other ST spots within the same cluster and $b(i)$ be the smallest average distance between i and all the ST spots with any other clusters. The silhouette coefficient $S(i)$ of ST spot i can be expressed by:

$$S(i) = \frac{b(i) - a(i)}{\max\{a(i), b(i)\}} \quad (7)$$

Silhouette score of an ST slide is the average silhouette coefficient of all spots in it.

In this piece of work, Pearson correlation based measurement in Eq. (2) was used to calculate the dissimilarity between ST spots from the imputed ST data. Traditionally, Euclidean distance is used but here we adopted the dissimilarity metric defined in Eq. (2) since our clustering method is based on Pearson correlation distance. Padded spots are excluded in this analysis. To make the average silhouette coefficient of clustering under different smoothing methods comparable, we used the same distance matrix which was based on dimensionality reduced original unsmoothed ST expression. Hence, in our experiments, comparing with unsmoothed slides, an increase or lack of change in

silhouette score on smoothed slides represents the smoothing method has enhanced the original data distribution while a decrease indicates the smoothing method has changed the distribution without keeping the originl one.

External evaluation

For the external evaluation, we obtained the histopathological labels of the ST spots. The correspondence between smoothed slides and histopathological labels was evaluated at both clustering and gene expression level. At the clustering level, the imputed ST data is clustered using multiple clustering algorithms. Technical details of the clustering methods is shown in Table S4. Next, the concordance was evaluated between the clusters from the imputed ST data and the labels from the corresponding histopathological images. Furthermore, the distribution of marker gene expression was compared to the locations of histopathological labels.

The Adjusted Rand Index (ARI) was used to evaluate the similarity between the clustering results from the imputed ST data and the histopathological labels [40]. For clusters from the imputed ST data $\{I_1, I_2, I_3, I_4\}$ and the histopathological categories of ST spots $\{H_1, H_2, H_3, H_4\}$, we denoted n_{ij} as the number of ST spots that are both in I_i and H_j . The ARI is defined as:

$$ARI = \frac{\sum_{ij} \binom{n_{ij}}{2} - \frac{\sum_i \binom{n_i}{2} \sum_j \binom{n_j}{2}}{\binom{n}{2}}}{\frac{1}{2} \left[\sum_i \binom{n_i}{2} + \sum_j \binom{n_j}{2} \right] - \frac{\sum_i \binom{n_i}{2} \sum_j \binom{n_j}{2}}{\binom{n}{2}}} \quad (8)$$

where n_i is the number of ST spots in I_i and n_j is the number of ST spots in H_j . A higher ARI value indicates that the imputed ST clusters and the histopathological labels are more

similar. In this analysis, unable to cluster padding spots with ground truth will be treated as error clustered.

In the marker gene evaluation, For PDAC dataset, two marker genes, *PRSS1* and *TM4SF1*, were used to compare their expression distribution spatially. Both genes are protein-coding genes. *PRSS1* encodes a trypsinogen, which is often highly expressed in normal pancreatic tissues while *TM4SF1* is a common proto-oncogene and highly expressed in pancreatic cancer among other malignancies [41, 42]. High expression of *TM4SF1* in the cancerous regions of the PDAC dataset has been detected in previous research [10]. For DLPFC dataset, three other marker genes, *MOBP*, *PCP4* and *SNAP25*, is used in the analysis. Previous research has reported that these genes can delineate different cortical layers [27]. The expression distribution of these representative genes can stratify the histopathological regions and can be used to measure the accuracy of that partition.

Biological analysis

To examine how our algorithm aids in informing the biology of a ST sample, we identified differentially expressed genes (DEGs) and performed gene ontology enrichment analysis (GOEA). DEGs were identified in the PDAC slides by comparing two groups of spots with *TM4SF1*-high (neoplastic tissue) and low expression (non-neoplastic tissue). To define the two groups, we first linearly transformed the expression values of *TM4SF1* between [0,1] by dividing each value by the maximum expression value. The spots on each slide were split into two groups with one having a transformed expression value less than 0.7 (*TM4SF1* under-expressed) and the other greater (*TM4SF1* over-expressed). The cutoff 0.7 was chosen as it can reflect the boundary of the cancer region accurately.

A Kruskal-Wallis test was performed between the over- and under-expressed groups and the p-values were adjusted using the Benjamini-Hochberg method to account for the multiple comparisons [43]. Only genes with an adjusted p-value less than 0.05 were included. In addition, we used logarithmic foldchange (logFC) to determine the up-regulated and down-regulated events for genes. Genes with a logFC greater than 1 were considered up-regulated, and down-regulated if the logFC were less than -1.

GOEA was performed on the DEGs using *g:Profiler* [44]. The significance of enriched terms were tested by cumulative hypergeometric test and p-values were corrected by *g:SCS* method [45]. Only terms with an adjusted p-value below 0.05 were reported. All data sources offered by *g:Profiler* were used including Gene Ontology (GO) [46], Reactome [47], KEGG [48], WikiPathways [49], Transfac [50], CORUM [51], Human Protein Atlas [52], and the Human Phenotype Ontology [53]. Heatmaps for each sample were then generated to compare the terms found by each smoothing algorithm.

The DLPFC slides were processed slightly differently than the PDAC slides. To find the differentially expressed genes, one layer of the cortex was compared with the other six layers. For example, the expression values for all the spots in Layer 1 (L1) were compared to the expression values for the spots in layers L2, L3, L4, L5, L6, and WM (white matter). A Kruskal-Wallis test was performed between one layer of the cortex and all other layers, p-values corrected using the Benjamini-Hochberg method, and logFC calculated. This resulted in 7 sets of comparisons. The same cutoff of p-value less than 0.05 and logFC greater than one were used. The significant DEGs were then compared to a set of DLPFC layer marker genes published by Zeng et al. [54]. Enrichment analysis was performed using *g:Profiler* in the same way as the PDAC slides were processed.

Results

In this study, we applied our two-factor smoothing method SPCS and two state-of-the-art one-factor smoothing methods (MAGIC and SAVER) to smooth ten PDAC ST slides. To compare the performance of different smoothing methods, we evaluated the quality of generated clusters after performed K-Medoids unsupervised clustering on both smoothed and unsmoothed expression. The generated clusters and the distribution of marker gene expressions were compared with pre-labeled histopathological partitions to check if the smoothed expressions are more accurately reflecting the pathology features of the corresponding images. In addition, the DEGs were identified in different regions with respect to the marker gene *TM4SF1* expression level and GOEA was performed to reveal the biological meanings revealed by the smoothing.

Figure 2 placement

Internal Evaluation

The silhouette score indicates average distance between clusters in a slide. Since we used a same unsmoothed distance matrix for each slide smoothed by different smoothing methods, silhouette score in this analysis also can reflect how well the smoothing methods keep the original data distribution. For different smoothing methods, a greater silhouette score than unsmoothed slide usually represents smoothed expressions have better separability based on original data distribution. In the contrast, the decrease of silhouette score indicates the smoothing method has changed the original data distribution. After clustering the spots into corresponding clusters, we calculated the silhouette score of ten PDAC slides and eight DLPFC slides with seven clusters for each smoothing method in Figure 2(a) and Figure S1. In most of the slides in these two datasets, SPCS and MAGIC

got similar or even greater silhouette score to the unsmoothed slide while SAVER got a significantly lower score. In addition, SPCS got the greatest average silhouette score than other smoothing methods, even slightly higher than the unsmoothed slide. This result indicates that kNN-based smoothing generally can keep the characteristics of original data distribution better and SPCS can contribute to better data separability.

Figure 2 (b)-(c) illustrated how the parameters will influence the data separability of SPCS smoothed PDACA1 slide. For the two ratio parameters α and β , while $\beta \leq 0.4$, silhouette score will not be affected a lot by the changing of α , but the result will become more sensitive to α if β is large enough. This result indicates that pattern based smoothing can help to keep the original data distribution. In addition, the results also reveal a truth that data distribution nature of spatial neighbors is different from the pattern neighbors. For the neighborhood size parameters τ_s and τ_p , the results show that they will have no significant influence on data separability.

External Evaluation

In the external evaluation, different smoothing methods were evaluated based on the consistency of unsupervised clusters with histopathological labels. The result of PDACA1 was shown in Figure 3 since the histopathological labels were available for PDACA1. This slide can be well clustered by k-Medoids clustering method even without smoothing. When examining the results in detail, Figure 3(b) shows that most ST spots in the cancer cells and desmoplasia region can be separated from other clusters. Most ST spots in the duct epithelium region can also be well separated, with slight mixing of the interstitium and the normal pancreatic tissue regions. It is worth mentioning that by including spatial position information, SPCS can pad missing spots, which other one-factor smoothing method cannot do. Figure 3(c, d) show the influence of different smoothing methods on the two

marker genes expression. Compared with other smoothing methods, SPCS generated better marker gene contrast between distinctive histopathological areas due to two reasons. 1) By performing smoothing, marker gene expressions showed fewer drop-outs and better reflected expression patterns across different tissue regions with SAVER and SPCS (MAGIC does not impute the missing expressions). 2) In contrast to SAVER, SPCS imputed the missing values and kept the spatial distribution of non-missing values relatively stable.

Figure 3 placement

Figure 4 illustrates external evaluation results from 12 slides in DLPFC dataset. Figure 4(a) shows the influence of smoothing methods on different clustering methods. To make the evaluation comprehensive, we chose three commonly used general clustering methods (i.e. k-Medoids, Louvain and mclust) and two state-of-the-art clustering methods developed specifically for ST slides (i.e BayesSpace and SpaGCN). The results revealed that smoothing helps to get a higher ARI score for various clustering methods and SPCS outperform other one-factor smoothing methods for every clustering method. When combining SPCS and Louvain or mclust clustering methods, we can get a ARI score near BayesSpace for DLPFC dataset, which means the combination of SPCS with general clustering methods is good to performing spatial clustering on ST slides. In addition, combining of SPCS and BayesSpace got the highest ARI score among all the combinations. Due to the difference of preprocessing steps, the results of SpaGCN clustering were moderately lower than the original publication [38], which indicates that hyperparameter settings of SpaGCN is sensitive to preprocessing steps and we hope to optimize this in the future. Figure 4(b) and Figure S2(a) illustrated clustering ground truth and BayesSpace clustering results while combining with different smoothing methods on slide 151675 and 151673. Compared with other one-factor smoothing or without

smoothing, SPCS can help to get a more clear and accurate boundaries between different regions, which contribute to a higher ARI score. In addition, smaller spatial neighborhood (τ_s) for SPCS can help to capture long narrow regions in the slide but may cause over clustering in thicker regions with similar length and width. Similar to PDAC dataset, SPCS also helps to obtain a better marker gene contrast between distinctive cortical layers in DLPFC dataset as shown in Figure S2(b-d).

Figure 4 placement

We also performed simulation analysis to test the clustering accuracy for different smoothing methods. Louvain and BayesSpace, the two best general and ST dedicated clustering methods in DLPFC experiment, were used in this analysis. Clustering ground truth is shown in figure 5(a). For the 10 simulated slides, Figure 5(b) illustrated distribution of ARI score of each combination of smoothing and clustering method. In general, since the original single cell dataset was well clustered, every combination of smoothing and clustering methods got high ARI score in this experiment. On the track of Louvain clustering, there's only a little difference of ARI score on different slides. For different smoothing methods, SPCS gets a slightly higher average ARI score which indicates that spatial information helps general clustering methods to better clustering ST slides. However, Louvain separates the spots into five clusters instead four, which means Louvain tends to cluster the spots according to cell types. For the BayesSpace experiments, the ARI score was lower than Louvain in average and greater in variance. To better review the higher variance of BayesSpace, we illustrated BayesSpace clustering results of two simulating slides in Figure 5(c) and (d). Obviously, BayesSpace tends to merge small clusters into nearby larger ones which leads to oversmoothing in the simulating dataset, and SAVER and SPCS smoothing will aggravate this problem. Overall all methods performed well on the simulated data (ARI > 0.85) but it is worth noting that

SPCS has higher potential ARI as evaluated by the 75th quantile. This is surprising considering the other smoothing methods are specifically designed for scRNA-seq data from which these simulated ST slides are generated.

Figure 5 placement

Biological Analysis

The biological interpretability of the smoothed results was compared between different smoothing methods. When comparing between *TM4SF1* over- and under-expressed regions for the PDAC slides, the significant DEG numbers were displayed in Figure 6(a). There were no DEGs found in any smoothed or unsmoothed slides of PDACB2 and PDACG because they lacked *TM4SF1* while a higher number of DEGs was detected by SPCS in six out of the rest eight ST slides. Correspondingly, the number of enriched GO terms was also more from SPCS than the other methods, as shown by Figure 6(b), which are further examined below.

The top ten most significant GO terms found in slide PDACA1 for each algorithm are shown in Figure 6(c) for the up-regulated DEGs and Figure 6(d) for the down-regulated DEGs. More than ten terms are shown in the heatmap because most GO terms were not shared between smoothing methods. Without smoothing the slides, the up-regulated DEGs identified are related to interleukin-1. Instead, the terms found by SPCS smoothing are related primarily to cell adhesion, extracellular matrix (ECM) organization and MET activates PTK2 signaling. The GO terms from all smoothed and unsmoothed up-regulated slides can be seen in Figure S3. *g:Profiler* could not find any enriched terms for slides PDACB2, PDACD, and PDACG. Up-regulated enriched terms for all SPCS slides except PDACB1 appear similar to PDACA1. The results for the other methods had a small

number of terms which do not seem related to PDAC, except for slide PDACF which contained cell adhesion and ECM terms for the no smoothing and MAGIC.

Figure 6 placement

The enriched GO terms for the down-regulated DEGs (Figure 6(d)) tell a different story. Without smoothing, GO terms related to digestion as well as infection and autoimmune related pathways and complement cascade components (C3, C5) were identified. GO terms related to digestion were seen with SAVER as well. After performing smoothing, we found many more GO terms related to the ECM, more complement binding, as well as apoptosis regulation, but the infection and autoimmune-related pathways were absent. Applying MAGIC and SPCS also helped to find other complement cascade components such as C1q complex binding. The other slides showed are in Figure S3 and appear similar results to the terms described above.

Figure 6(e) contains terms enriched in the SPCS smoothed data from up-regulated DEGs that have previously been reported in PDAC [55-58]. These terms, such as cell adhesion, Cadherin binding, PI3K-Akt signaling pathway and focal adhesion, are all significantly enriched in DEGs from SPCS-smoothed data (albeit not among the top-ten enriched terms), but were absent from unsmoothed data, reflecting the enhancement of biological interpretability by performing our novel SPCS smoothing method.

The results of the biological analysis of the DLPFC slides are shown in Figure S4. The significant DEGs are shown in Figure S4(a) and show a similar number of DEGs for each algorithm. When comparing the DEGs to those reported by Zeng et al. [54], a similar number of genes were shared for each smoothing method as shown in Figure S4(b). These genes include *MOBP* (an oligodendrocyte and white matter marker), *PCP4* (associated with L5 and L6) [27, 54], and *GFAP* (associated with astrocytes and many

neurological disorders) [59]. In addition, comparing with other smoothing methods, SPCS helped to discover five DEGs (i.e., *CADM4*, *ELOVL5*, *AGR2*, *LGALS3BP* and *SCARB2*) which not previously reported. These genes were all found in the white matter layers of the brain. The expression of *CADM4* and *SCARB2* in sample 151673 are shown in Figure S4(c) and Figure S4(d), respectively.

Discussion

Importance of smoothing on ST data

ST data is based on highly multiplexed sequence analysis where barcodes are used to split the sequenced reads into their respective tissue locations. However, this type of sequencing suffers from high noise and drop-out events. To keep enough genes to perform biological analysis, we set a relaxed filtering threshold (less than 70% zero expressed spots) to filter out non-expressed genes in preprocessing steps. Even with this rather relaxed threshold, less than 10% of genes in slides of both PDAC and DLPFC datasets were left, which indicated that drop-out events are highly frequent in ST datasets. By visualizing the expression distribution of two marker genes, *PRSS1* and *TM4SF1* in Figure 3, the gene-level drop-out events can be easily seen, as indicated by the black-colored spots on the slides. For both genes, drop-out events frequently occurred in interstitium regions of the slide, leading to a failure cluster this area on unsmoothed ST data. In addition, there's also entire spots missing in multiple ST slides, which can negatively influence spatial clustering. By performing smoothing methods, the missing and noisy expression values were controlled to a certain extent so that the partition of regions and downstream analysis are greatly improved. Therefore, smoothing is an important and necessary step for analyzing ST data.

543 SPCS Improves Data Quality

544 Compared with unsmoothed data, smoothing step improves data quality. We have
 545 demonstrated that various smoothing algorithms increase the separability and partition
 546 accuracy of ST spots. Moreover, by performing internal and external evaluations, we
 547 confirm that SPCS-smoothed data show better quality on both aspects as compared with
 548 the two existing one-factor smoothing methods MAGIC and SAVER. For internal
 549 evaluation, from the results shown in Figure 2, SPCS smoothing method produces greater
 550 silhouette scores than MAGIC and SAVER, which means the ST data smoothed by SPCS
 551 has better separability. In addition, a more similar silhouette score of SPCS smoothed and
 552 original unsmoothed data indicates that SPCS can better preserve original data
 553 distribution of non-missing values, which helps to keep an accurate biological analysis
 554 results while improving spatial clustering accuracy.

555 Our external evaluation verifies the partition accuracy of smoothed data. One main
 556 objective of slide partition is to identify different histopathological regions in the slide.
 557 Hence, we measured the degree of overlap between unsupervised clusters on smoothed
 558 data and histopathological partitions using ARI. Since the spots in the same
 559 histopathological region are usually connected, incorporating spatial knowledge is
 560 expected to boost partition accuracy. Moreover, spatial knowledge can also help to detect
 561 and pad missing spots. Indeed, as expected, results in Figure 3-5 reveal SPCS method
 562 generates a higher ARI score than existing one-factor methods, which means a more
 563 accurate histopathological partition can be acquired by performing two-factor SPCS
 564 method. In addition, it is also cleared that SPCS can be used before various clustering
 565 methods to improve clustering accuracy. From the marker gene analysis, SPCS method
 566 recovers the drop-out events and enhance the expressions of marker genes in the

corresponding regions. This evidence proves that SPCS can improve the accuracy of the ST spot partitions.

SPCS Enhances Biological Interpretability

The GO terms reported in PDACA1 from data using the SPCS smoothing method are more interpretable in the context of pancreatic cancer and the role of *TM4SF1* in cancer than non-smoothed data and the two one-factor smoothed data. *TM4SF1* has been found to be overexpressed in pancreatic cancer and has roles in apoptosis, proliferation, and cell migration [41]. Previous work found that collagen 1 (GO:0030199) binds with DDR1 which then interacts with *TM4SF1* to activate the focal adhesion kinase (FAK) [41, 60, 61]. This results in disruption of E-cadherin (GO:0045296) leading to the Wnt signaling pathway and loss of cell-cell adhesion (GO:0050839) [41, 60]. The FAK pathway also increases expression of N-cadherin (GO:0045296) which results in migration of cancer cells [41, 60]. *TM4SF1* can also activate the *AKT* pathway (KEGG:04151) leading to anti-apoptotic effects and angiogenesis [41, 62]. All of these previously identified GO terms are important factors for PDAC survival and metastasis. They were also identified using SPCS-smoothed data only, which would have been missed using other smoothing methods or with the unsmoothed ST data.

Identifying the DEGs in different histological regions and check their associated GO terms can also help evaluate the biological interpretability of a smoothed slide. It is worth noting that the gene ontology (i.e. gene set database) and the enrichment tool can yield different results. The simplest example would be the use of a hypergeometric test to determine enriched gene sets opposed to gene set enrichment analysis which accounts for the significance of DEGs. The hypergeometric test has a longer history of use and is easily interpretable whereas gene set enrichment analysis is a newer approach. Furthermore,

the gene sets themselves differ between databases such as KEGG and GO. There could potentially be more enriched KEGG pathways for one sample and more enriched GO terms for another. For these reasons, we use g:Profiler since it incorporates many gene set databases in the analysis and relies on the well-established hypergeometric testing approach.

In our analyses we chose to use Our SPCS method contributed to identify many more differentially expressed genes than the other smoothing methods for most of the slides. Among the DEGs, the up regulation of *IL1RN*, *KRT7*, *LAMB3*, *LAMC2*, *NOTCH3*, and *S100A16*, which have previously been reported associate with poor survival in PDAC, was also detected in the SPCS smoothed PDACA1 slide [56, 63]. *IL1RN* and *LAMB3* were also found using unsmoothed data and *LAMB3* was detected with MAGIC smoothed data, but the other genes were uniquely detected by SPCS. Generally, a high number of DEGs also associate with more GO terms. However, some unsmoothed, SAVER, and MAGIC-smoothed slides contained DEGs but no GO terms, which may indicate the insufficient imputing or smoothing without considering the location information. Specifically, only slides PDACE and PDACF contained up-regulated DEGs and GO for SAVER and MAGIC; although PDACD was the only slide which had DEGs and no GO terms for the down-regulated genes. The unsmoothed up-regulated slides PDACA2, PDACB3, PDACD, and PDACE, and down-regulated slides PDACA3, PDACB3, and PDACD also had DEGs but no GO terms. In contrast, every SPCS smoothed slides that had DEGs also had GO terms. The SPCS data demonstrated a *TM4SF1* expression landscape that matched the original histopathological assignment more accurately than the other smoothing methods. For this reason, it's possible SPCS found more DEGs because other genes in the slide had more defined expression after smoothing.

The terms found through GOEA were more interpretable in the scenario of PDAC and its

pathogenesis when applying SPCS smoothing. Many of the SPCS top ten terms in Figure 4(c) and previously reported terms in Figure 4(e) are similar to those found in the literature [55-58] and are consistent between slides. Without applying SPCS, some of the top terms found in Figure 4(d) are involved in typical pancreatic activity such as peptidase regulator activity (GO:0061134), digestion (GO:0007586), and triglyceride lipase activity (GO:0004806) which indicates that important PDAC-pathology related GO terms may be missed when data is not smoothed or not properly smoothed.

Using the cortical layers in the DLPFC slides, we found a similar number of DEGs and shared genes between the smoothing methods. Most of these genes were previously reported. However, profiting from contrast enhancement, our proposed SPCS method helped to identify five DEGs in the white matter while the other methods did not. The Cell Adhesion Molecule 4 (*CADM4*) and scavenger receptor class B Member 2 (*SCARB2*), to the author's knowledge, have not been identified as white matter markers in the DLPFC. Cell adhesion molecules (CAMs) like *CADM4* play an important role in myelination by oligodendrocytes. Higher expression of *CADM4* leads to many short myelin internodes that disrupt the normal myelination process [64]. *SCARB2* is a lysosomal membrane receptor for the glucocerebrosidase enzyme. It has been associated with Parkinson's disease and Lewy Body Disease. Glucocerebrosidase degenerates sphingolipid, which is important for brain development. There is some evidence that decreases sphingolipids can lead to demyelination [65, 66]. While the authors do not intend to present *CADM4* and *SCARB2* as marker genes for white matter, we believe that SPCS can be used to help aid with this task. The enrichment analysis results for DLPFC are not shown but are similar between all smoothing methods. This is likely because the number of GO terms found is correlated with the number of DEGs. It's possible that using a gene markers instead of layers to identify DEGs could produce different results. Given the generally clearly defined

boundries of the brain, using layer data to get differential gene expression seems more appropriate.

Determination of SPCS parameters

There are four parameters in SPCS: τ_s , τ_p , α and β . These parameters affect the performance of SPCS algorithm by adjusting neighborhood size and smoothing intensity and should be tuned according to the quality and distribution of original ST data.

τ_s and τ_p are designed to adjust the size of spatial neighborhood and pattern neighborhood, respectively. Obviously, including more information while performing smoothing is beneficial for a more robust result, and increasing the size of neighborhood is a good way to achieve that goal. However, Considering blindly expanding neighborhoods will incorporate some spots that are not similar to the spot being smoothed, SPCS method use contribution weighting to reduce this effect, which gives the size of both spatial and pattern neighborhoods limited influence on data separability as shown in Figure 2(e). In addition, as shown in Figure 4 (b) and (c), spatial neighborhood will also influence clustering sensitivity. A smaller spatial neighborhood (τ_s) for SPCS can help to capture long narrow regions in slides but may cause over clustering in thicker regions with similar length and width. Therefore, we recommend a modest selection of these two parameters to balance the tradeoff. For most cases, $\tau_p \leq 16$ and $\tau_s \leq 4$ is recommended.

The parameters, α and β , are designed to balance the original expression and corrections from spatial and pattern neighbors, which has a significant effect on smoothing quality. Due to the pattern similarity between object spot and its pattern neighbors, corrections from pattern neighbors trend to enhance the original data distribution features, which can help to increase or stabilize on silhouette score. In the contrast, corrections from spatial

neighbors try to make the object spot expression consistent to its spatial neighbors, which is beneficial to spatial clustering but may change the original data distribution leading to a worse silhouette score. To keep the accuracy of spatial clustering and biological analysis simultaneously, it is important to balance the intensity of correction with the underlying expression signatures. Results in Figure 2(d) indicate that the influence of smoothing strength (α) on data separability is heavily reliant on the proportion of spatial and pattern correction (β). Hence, we recommend setting α between 0.2 to 0.8 and β less than 0.6 in most cases and should be selected carefully according to data distribution. In addition, as indicating in Figure 5, when combining with spatial clustering methods like BayesSpace, smaller α and β are recommended to avoid erroneous merging of small clusters.

Conclusion

Due to the limitation of barcoding based sequencing technologies which result in noisy and high drop-out in transcriptomics data, smoothing has become an essential data processing step before performing the downstream analysis on ST data. In this paper, we proposed a novel two-factor ST data smoothing method SPCS, which can take full advantage of both the expression patterns and the spatial patterns contained in ST data. Compared with traditional one-factor smoothing methods, we have demonstrated that SPCS can recover the missing or erroneous expression in ST data better and improve the separability, partition accuracy, and biological interpretability. This reflects that our proposed SPCS method can effectively improve data quality of ST and contribute to more accurate and meaningful downstream analysis results. SPCS smoothing is a state-of-the-art ST smoothing algorithm with implications in numerous diseases where ST data is being generated.

Key Points

- Due to the common issue of noise and drop-out events in ST data, smoothing has become a necessary step before downstream analysis on ST data.
- SPCS is a novel kNN-based two-factor smoothing method that can fully utilize both expression pattern and spatial knowledge in ST data.
- Compared with traditional expression pattern knowledge-based one-factor smoothing methods, SPCS can provide better separability, partition accuracy and biological interpretability.

Data Availability

All of the data used for the analyses in the manuscript are freely available from their original publications.

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735

736 **Figures**

737

738 **Figure 1** Workflow of our proposed SPCS smoothing method. (a) An example of
739 smoothing. In this sample, SPOT 1 is the object spot that going to be smoothed. For ST
740 slides, preprocessing steps normalize the expressions and filter out genes with zero

741 expression in most spots. Spatial neighborhood is second order neighborhood of SPOT 1
742 with nine spots. Spots where all gene expression is 0 are treated as non-tissue regions
743 and excluded from spatial neighborhood. Pattern neighborhood consists of the top eight
744 spots with most similar expression pattern to SPOT1. Both spatial and pattern
745 contributions are normalized which means total contribution of all neighbors is equal to 1.
746 Smoothing is performed by integrating parameter- and contribution-weighted expression
747 of the object spot itself and both its spatial and pattern neighbors. (b) Shape of second
748 order spatial neighborhood for ST and Visium platforms. Spatial neighborhood is
749 determined by Manhattan distance. Due to the difference of spatial pattern of spots, the
750 shape of spatial neighborhood could be different in different platforms. (c) Padding
751 strategy of SPCS. A missing spot will not be padded unless there are more than 50% of
752 unmissing spots inside its spatial neighborhood.

753

754 **Figure 2** Influence of smoothing on data separability and data distribution. (a) Box plot of
755 average silhouette score of ten PDAC samples and eight 7-layer DLPFC samples without
756 smoothing and with smoothing by different methods (MAGIC, SAVER and SPCS). (b)
757 Influence of parameters α and β on average silhouette score for SPCS smoothed
758 PDACA1 slide when $\tau_p = 16, \tau_s = 2$. (c) Influence of parameters τ_p and τ_s on average
759 silhouette score for SPCS smoothed PDACA1 slide when $\alpha = 0.6, \beta = 0.4$.

Figure 3 Influence of smoothing on clustering accuracy in PDACA1. (a) Original ST slide of PDACA1 and its histopathological partitions, (b) Results of K-Medoids clustering on expression smoothed by different methods. ARI score, which reflects the correlation between clusters and histopathological labels, is marked at bottom-right corner of each figure. Clusters are ordered by their size. Heatmap of smoothed expressions of two marker genes, (c) *PRSS1* and (d) *TM4SF1*, are also shown. For demonstration purposes, expressions of genes are linear transformed into range of 0 to 1 as normalization.

Figure 4 Influence of smoothing on clustering accuracy in DLPFC dataset. (a) Box plot of ARI score on various combination of smoothing (No smoothing, MAGIC, SAVER and SPCS) and clustering methods (k-medoids, Louvain, mclust, BayesSpace and SpaGCN) in 12 DLPFC samples. (b) Ground truth label and BayesSpace clustering results of smoothed and unsmoothed sample 151675. Clusters are ordered by their size.

774

775 **Figure 5** Influence of smoothing on clustering accuracy in HGSOc simulating dataset. (a)
776 Ground truth labels of a simulated ST slide. (b) Box plot of ARI score on Louvain and
777 BayesSpace clustered 10 smoothed simulated ST slides. (c) and (d) illustrate BayesSpace
778 clustering results of two of the simulated ST slides. Clusters are ordered by their size.

779

780 **Figure 6** Biological analysis of unsmoothed and different methods (MAGIC, SAVER and
781 SPCS) smoothed PDAC slides. (a) Number of DEGs identified in each slide and (b) the

782 number of GO terms found from GOEA in each slide shown on a log10 scale. DEGs which
 783 are down-regulated in *TM4SF1* over-expressed region and their corresponding GOEA
 784 terms are marked as slash filling texture. The ten most significantly enriched terms from
 785 each smoothing method found in the (c) up-regulated DEGs and (d) down-regulated DEGs
 786 groups in *TM4SF1* over-expressed region for PDACA1 slide are shown, shaded by -
 787 log10(p-value). (e) GO terms enriched in SPCS smoothed slides from up-regulated DEGs
 788 that have previously been found in PDAC.

789

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