AWGAN: A Powerful Batch Correction Model for scRNA-seq Data

Tianyu Liu[1,#], Yuge Wang[2,#], Hongyu Zhao[2,*]

[1] University of Illinois at Urbana-Champaign

[2] Yale University

#: These authors contribute equally to this work.

*: To whom the correspondence should be addressed.

With the advancement of technology, we can generate and access large-scale, high dimensional and diverse genomics data, especially through single-cell RNA sequencing (scRNA-seq). However, integrative downstream analysis from multiple scRNA-seq datasets remains challenging due to batch effects. In this paper, we focus on scRNA-seq data integration and propose a new deep learning framework based on Wasserstein Generative Adversarial Network (WGAN) combined with an attention mechanism to reduce the differences among batches. We also discuss the limitations of the existing methods and demonstrate the advantages of our new model from both theoretical and practical aspects, advocating the use of deep learning in genomics research.

1 Introduction

The advancement of single-cell sequencing technologies and the vast amounts of data generated have contributed significantly to biological research. One major challenge in single cell data analysis is batch effect that is present in essentially all single cell datasets [1–3]. The definition of batch effect refers to the differences in the distributions of single-cell RNA sequences for cells of the same tissue due to many experimental factors, including different

protocols, instruments, operators, and pre-processing methods [4]. The removal of batch effect is not only important to identify true biological signals but also to facilitate multi-omics analyses across studies.

Two main approaches have been proposed to reduce the batch effects [5]. The first one is to assume that data from different batches follow the same distribution so that we can determine the parameters in the distribution and remove the batch effects in the datasets through statistical methods. The second one is to select one batch as the reference batch and others as query batches and get the batch-effect-corrected data by constructing a mapping from the query batches to the reference batch. Representative methods for the first strategy include Limma [6], Combat [7], Liger [8], and scVI [9]. Representative methods for the second strategy include Mutual Nearest Neighbor (MNN) [10], Seurat v4 [11], Batch balanced kNN (BBKNN) [12], iMAP [13], and Harmony [14]. Even with the many methods available, the task of batch correction still presents many difficulties. The most crucial point is to find a way to perfectly mix cells from the same cell type across different batches and to be able to discern batch-specific cell types [13]. The existing methods have different focuses on the above two points, but the different focuses of these methods can lead to poor results sometimes after batch effect removal (see our discussion). Therefore, how to achieve a reasonable trade-off between the two remains a challenge in practice.

Moreover, the variety of single-cell datasets and the volume of data are increasing, challenging the assumptions made in some of the batch effect removal methods. For example, some statistical methods assume that single-cell data follow a specific distribution and need to estimate parameters associated with the distribution (e.g. mean and variance). However, this assumption may not hold for many single cell datasets. For example, we cannot assume that the scRNA-seq data of liver cells and blood cells follow the same distribution because

these two tissues are very different biologically. In addition, the accumulating data make it more difficult to estimate model parameters, and develop algorithms with manageable time complexity. As a result, traditional statistical models may not be the best candidates for batch effect removal for large and complex single cell data. Therefore, researchers have been adopting machine learning models based on approximation theory, where there is more flexibility on the distribution of input data [15]. However, there are also issues with currently deployed machine learning strategies. Firstly, the existing machine learning-based correction models are poorly generalized, with varying results on different datasets. Secondly, the machine learning models sometimes generate incorrect results due to the acquisition of datasets, processing, and the choice of model structure, which will be discussed later in this paper. Therefore, there is a need to overcome this limitation. Finally, machine learning-based models tend to be computationally more expensive and time consuming to train [16].

To address the limitations of the existing methods, we propose Attention Wasserstein Generative Adversarial Network (AWGAN) for batch effect removals. This model has a lighter structure, a relatively shorter training time (using a combination of hardware and software acceleration), and better stability and generalization. These benefits are demonstrated through our results.

2 Results

2.1 Overview of AWGAN

The workflow of AWGAN has three key steps: attention-driven data preprocessing, AWGAN training, and model evaluation (Figure 1):

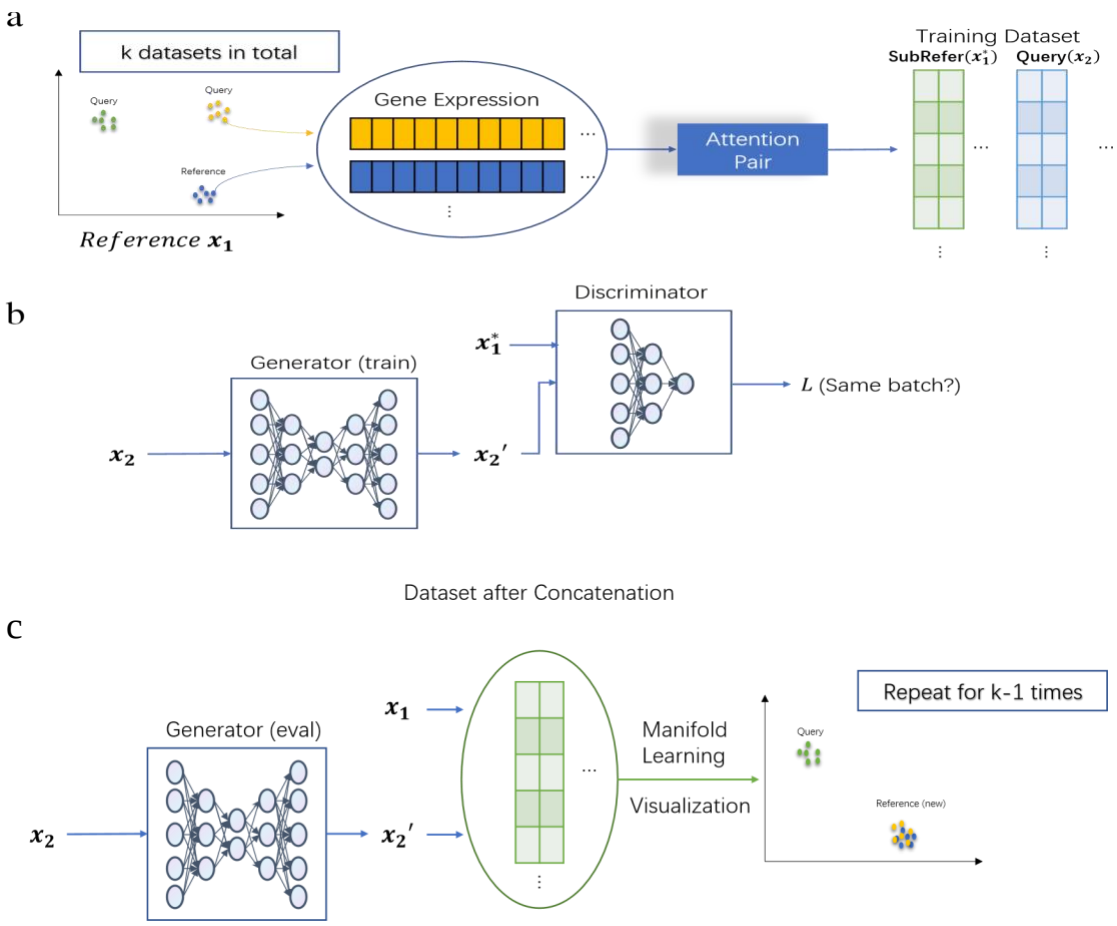


Figure 1: Workflow Chart. (a) This represents how we generate the training set. The attention pair will be transferred into gene expression data pair and fed into the model. (b) Training process for AWGAN. (c) Downstream process for the AWGAN model. The results received from the generator will be utilized for further work shown in Results and Discussion.

In the first step, highly variable genes are selected across batches as common features for data integration. We note that the dataset cannot be directly fed into the training model because we need to ensure that our method can preserve the biological information. Therefore, the number of cells in the training datasets needs to be compatible as $\#(SubReferData) = \#(QueryData)$. In this paper, inspired by the attention mechanism in the nature language processing scheme [17], we utilize a similar attention mechanism that can be

used to generate a subset of the reference data matched to the query data by selecting the nearest neighbor pairs (see Methods). Since training GAN is very tricky and requires the two sample distributions as similar as possible [18], utilizing matching pairs to generate the training set is important. When there are two batches, we only need to train the model once. When there are k ($k > 2$) batches in the target dataset, we utilize the incremental matching learning strategy by running the attention algorithm and the model training process $k - 1$ times. After each round, one query dataset will be integrated into the total reference batch, making the reference batch larger and larger. This strategy is adopted because different query batches may have different distributions.

In the second step, after generating the training dataset, we utilize the WGAN [19] with the gradient penalty [20] to generate a map that can transfer the query data so that their distribution is similar to the reference data. In addition, to reduce the possibility of model collapse, we also allow the self-adaptive mechanism in finding suitable regularization coefficients for the gradient penalty and making adjustment for the number of training epochs. WGAN is composed of one generator model and one discriminator model. The former model is used to generate data that follow a new distribution from the query data, while the latter model is used to discriminate whether the corrected query data have a similar distribution with the reference data. Moreover, the discriminator can find batch specific cell types and maintain their property. We adopt a strategy of confrontation training to improve the ability of the two models and finally achieve the Nash equilibrium between them [21].

In the last step, we generate an integrated scRNA-seq dataset after removing the batch effect. To visualize the results, we prefer UMAP [22] as a manifold learning choice. Moreover, to quantify the results after removing batch effects, we consider four metrics: ASW [23], kBET

[24], LISI [14], and Graph Connectivity [5] for quantitative evaluation of the performance of batch effect removal in this paper.

2.2 Small-scale scRNA-seq Datasets

We first evaluated our model against other methods on datasets with known cell type labels and moderate data size (less than 30,000 cells), which can be used as a standard to compare the effectiveness of different batch effect removal techniques. The datasets we chose in this part include: Pure Cell Lines (CL) [43], Human Dendritic Cells (DC) [44], Pancreas with certain cell types removed (Pancreas rm) [13] and PBMC3&68K [43], mouse hematopoietic stem and progenitor cells (MHSP) [45] [46], Mouse Cell Atlas [47] [48], human peripheral blood mononuclear cell (Human PBMC) [43], and Mouse Retina [50] [51]. In order to better demonstrate the better performance of our model, we provided two coordinate representations of two different metrics (ASW [23], and LISI [14]) for each dataset. We define the batch specific cell type discerning task as the cell type optimization task and the mixing of common cells in different batches as the batch optimization task. For the cell type optimization task, higher casw and 1-clisi values indicate better cell type separation. For the batch optimization task, higher 1-basw and blisi values indicate a better mixing of cells from the same cell type across different batches. The results based on UMAP and ASW/LISI map for the CL dataset, the DC dataset, the Pancreas rm dataset, and the PBMC3&68k dataset are shown in Figures 2-5, and the results of the MHSP dataset, the Mouse Cell Atlas dataset, the Human PBMC dataset, and the Mouse Retina data can be found in supplementary file 1 part 1.

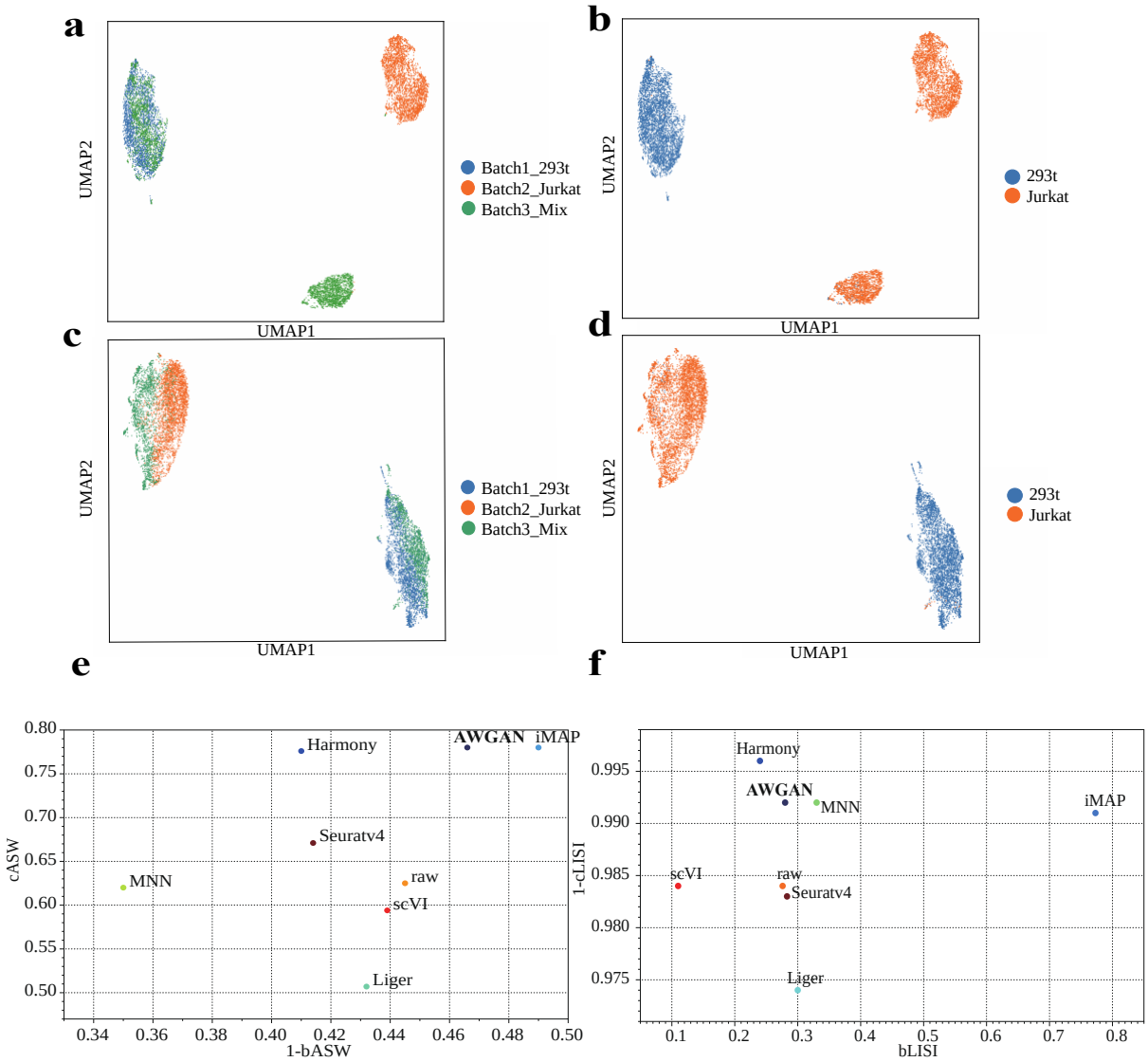


Figure 2: The performance of different batch effect removal methods for the CL dataset. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with cell type label before batch effect removal. (c) Visualization for the data distribution with batch label after batch effect removal. (d) Visualization for the data distribution with cell type label after batch effect removal. (e) ASW assessments of different batch effect removal methods on the CL dataset. (f) LISI assessments of different batch effect removal methods on the CL dataset.

The CL dataset (Figure 2) is composed of three different batches, including two pure batches “Jurkat” and “293T” containing Jurkat cells and 293T cells only and one batch “Mix”

consisting of a 50-50 mixture of Jurkat and 293T cells. We can see from the results that the batch effect is mostly eliminated for the three datasets, and the cells in Mix are mixed into the first two batches. The biological properties are also preserved, as cells in Mix are successfully mapped to their specified cell clusters. We can see that from Figure 2e that no method performs better than AWGAN in cell type optimization and batch optimization simultaneously except iMAP. AWGAN performs better than Liger, Seurat v4 and iMAP in cell type optimization according to Figure 2f, suggesting that AWGAN can better discern batch-specific cell types in this dataset. According to Figure 2 e and f, AWGAN only performs worse than iMAP in mixing cells in different batches for both ASW metric and LISI metric, suggesting good performance of AWGAN in this aspect.

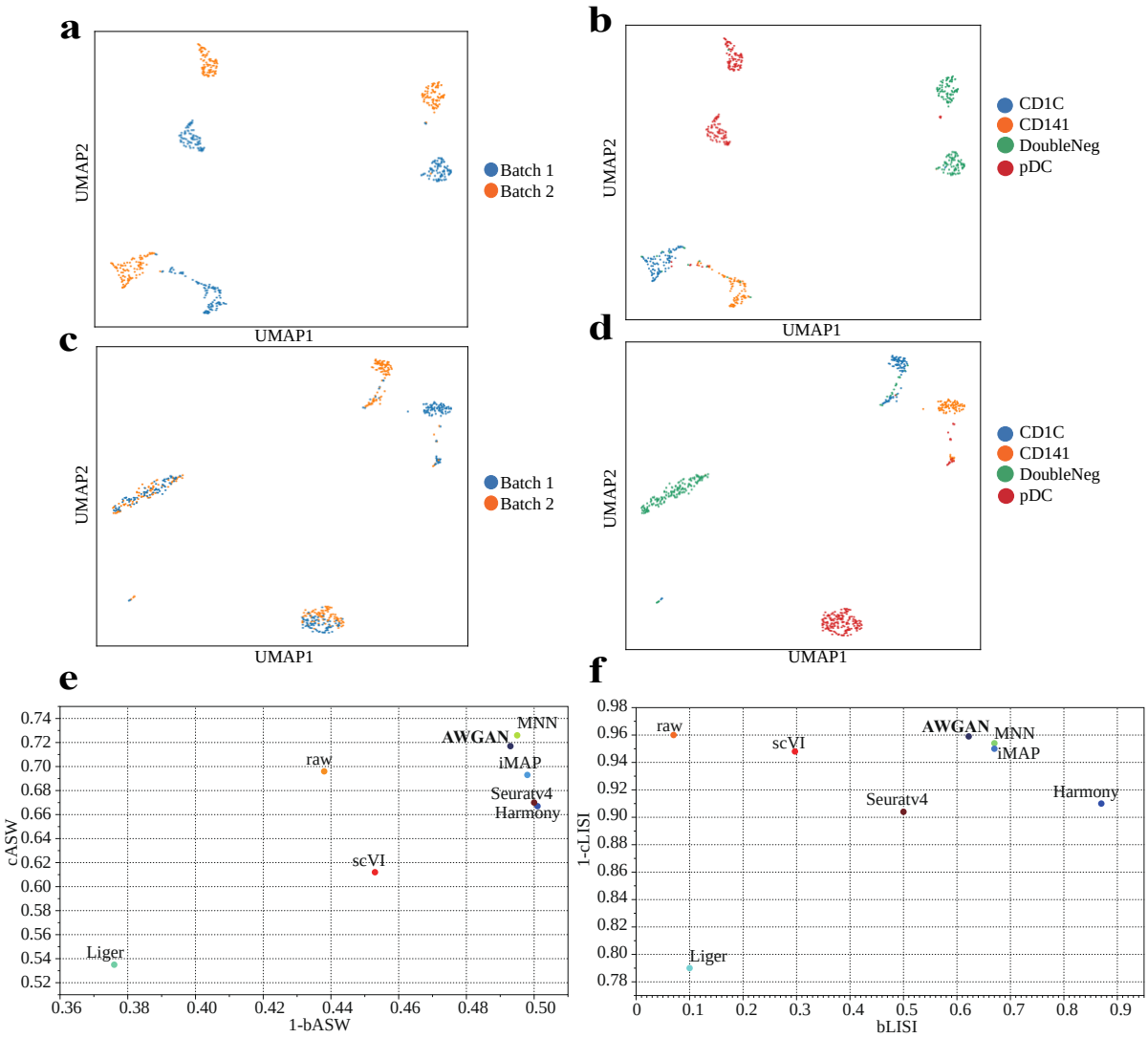


Figure 3: The performance of different batch effect removal methods for the DC dataset. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with cell type label before batch effect removal. (c) Visualization for the data distribution with batch label after batch effect removal. (d) Visualization for the data distribution with cell type label after batch effect removal. (e) ASW assessments of different batch effect removal methods on the DC dataset. (f) LISI assessments of different batch effect removal methods on the DC dataset.

The DC dataset (Figure 3) contains two batches, including four types of human dendritic cells: CD1C, DC, CD141, and plasmacytoid DC (pDC) cells. In the DC dataset, most of the batch variations are removed by AWGAN in both cell type optimization and batch optimization. Moreover, most of the cells are classified in the correct clusters. However, there still exist a few cells which are mixed incorrectly. This may be caused by the misclassification problem in the original data, because according to Figure 3 b, some cells in the prior batch-correction cluster have cell type labels different from neighbors. According to Figure 3 e and f, AWGAN performs better than most of the benchmark methods. Only MNN performs better than AWGAN on both the batch optimization task and the cell type optimization task with the ASW metric. Under the LISI metric, no method is better than AWGAN for both the batch optimization task and the cell type optimization task. Since this dataset contains batch-specific cell type labels, this experiment demonstrates AWGAN's effectiveness in the cell type optimization task.

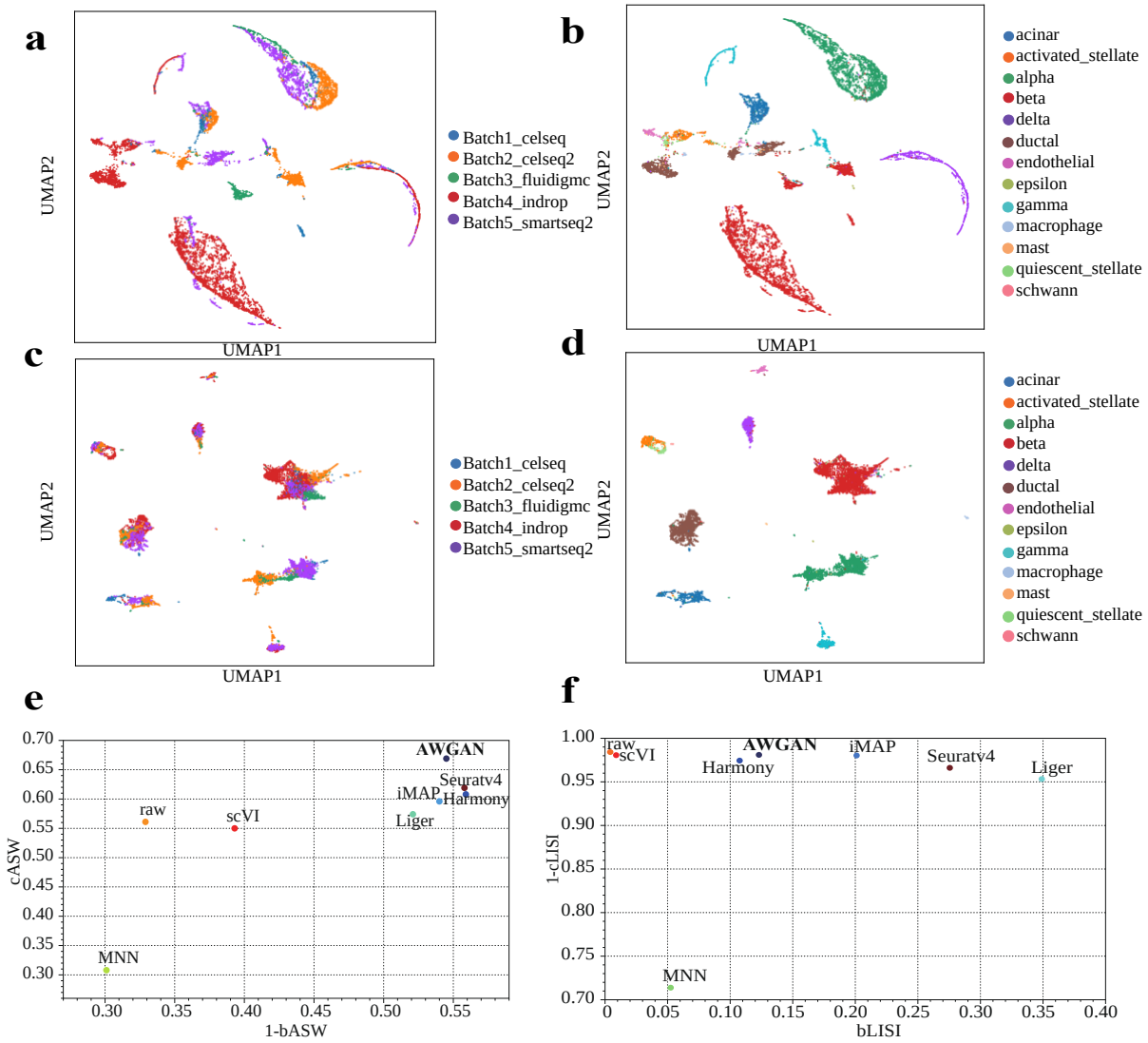


Figure 4: The performance of different batch effect removal methods for the Pancreas rm. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with cell type label before batch effect removal. (c) Visualization for the data distribution with batch label after batch effect removal. (d) Visualization for the data distribution with cell type label after batch effect removal. (e) ASW assessments of different batch effect removal methods on the Pancreas rm dataset. (f) LISI assessments of different batch effect removal methods on the Pancreas rm dataset.

The Human Pancreas dataset is initially composed of human pancreatic cells from five different sequencing methods. To evaluate the ability of AWGAN on both the mixing of the distribution of common cell types across different batches and discerning the batch-specific cell types, we intentionally removed some cell types from this original dataset. Specifically, we removed acinar and alpha cells from the inDrop batch data, and ductal cells from the CEL-seq batch data. In the end, we renamed the new dataset as Pancreas rm. We can conclude from the visualizations that AWGAN maintains a balance between common cell type mixing and specific cell type discernment across different batches according to Figure 4 c and d. In addition, according to the results in Figure 4 e and f, AWGAN performs well in this dataset based on quantitative results. For example, it achieves the best result in the cell type optimization task evaluated by ASW rate. In the function of preserving cell type information, AWGAN performs better than powerful methods like Harmony, Liger and iMAP.

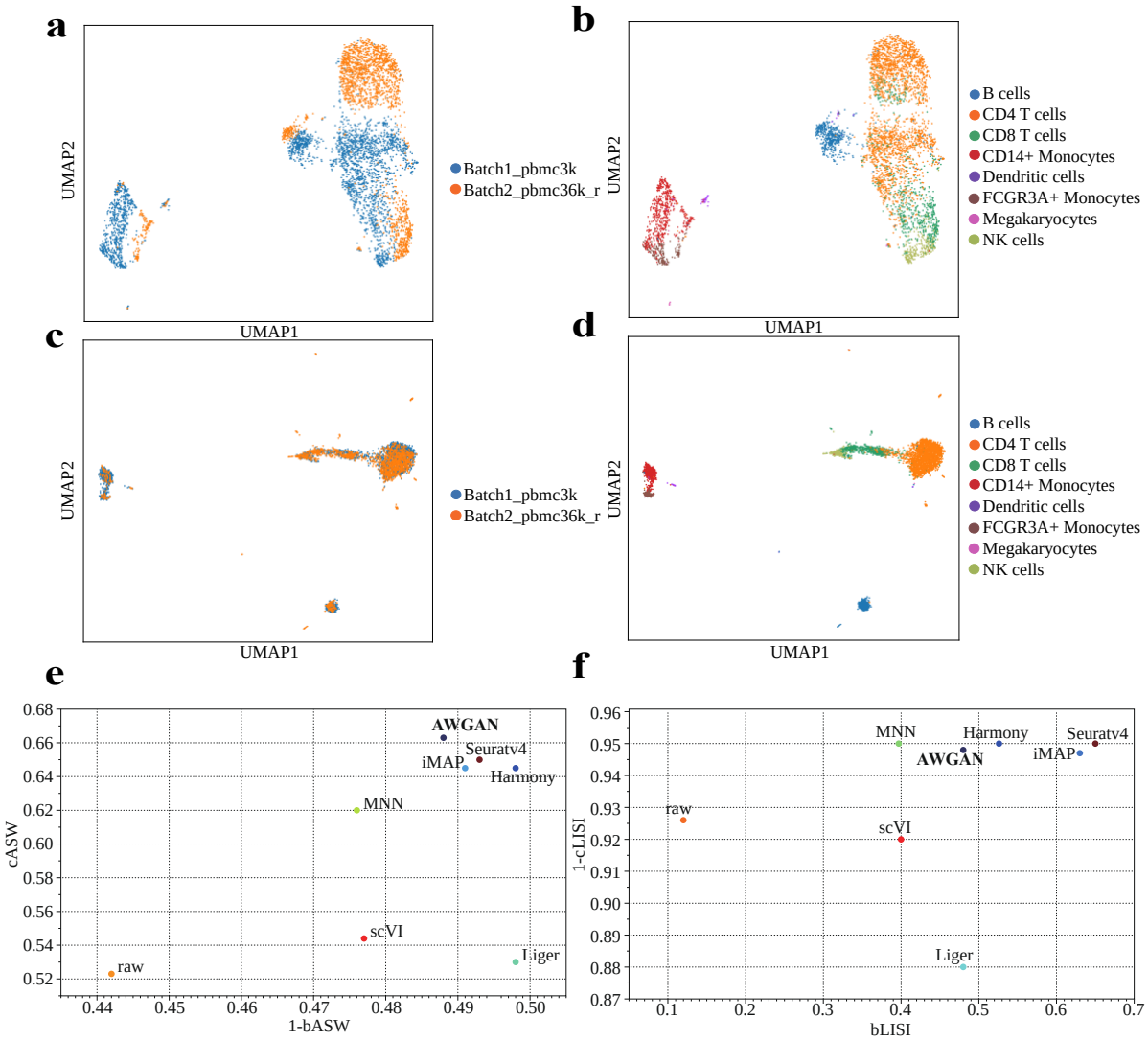


Figure 5: The performance of different batch effect removal methods for the PBMC 3&68K. (a) Visualization for the data distribution with batch label before the batch effect removal. (b) Visualization for the data distribution with cell type label before the batch effect removal. (c) Visualization for the data distribution with batch label after the batch effect removal. (d) Visualization for the data distribution with cell type label after the batch effect removal. (e) ASW assessments of different batch effect removal methods on the PBMC3&68K dataset. (f) LISI assessments of different batch effect removal methods on the PBMC3&68K dataset.

The PBMC 3&68K dataset (Figure 5) contains two PBMC datasets with different numbers

of cells. For PBMC 3K, we utilized the entire dataset whose number of cells is near 3,000, but for PBMC 68K, we sampled the raw version with 2,000 cells to decrease the workload. The main cell types contained in PBMC are B cells, CD 4/8 T cells, Megakaryocytes, and NK cells. According to the steps in the Scanpy document, we first mapped cell type labels from the PBMC 3k dataset to the PBMC 68k reduced dataset by using INGEST [25] and then performed batch correction. On this dataset, AWGAN still maintains good performance. Because the two batches do not contain batch specific cell types, we reached the batch removal target easier than the previous work. From Figure 5 c and d, we can conclude that AWGAN is able to remove the batch effect in this dataset effectively, and most of the benchmark methods have worse performance based on the results shown in Figure 5 e and f.

We plot running time for different batch correction methods in Figure 6.

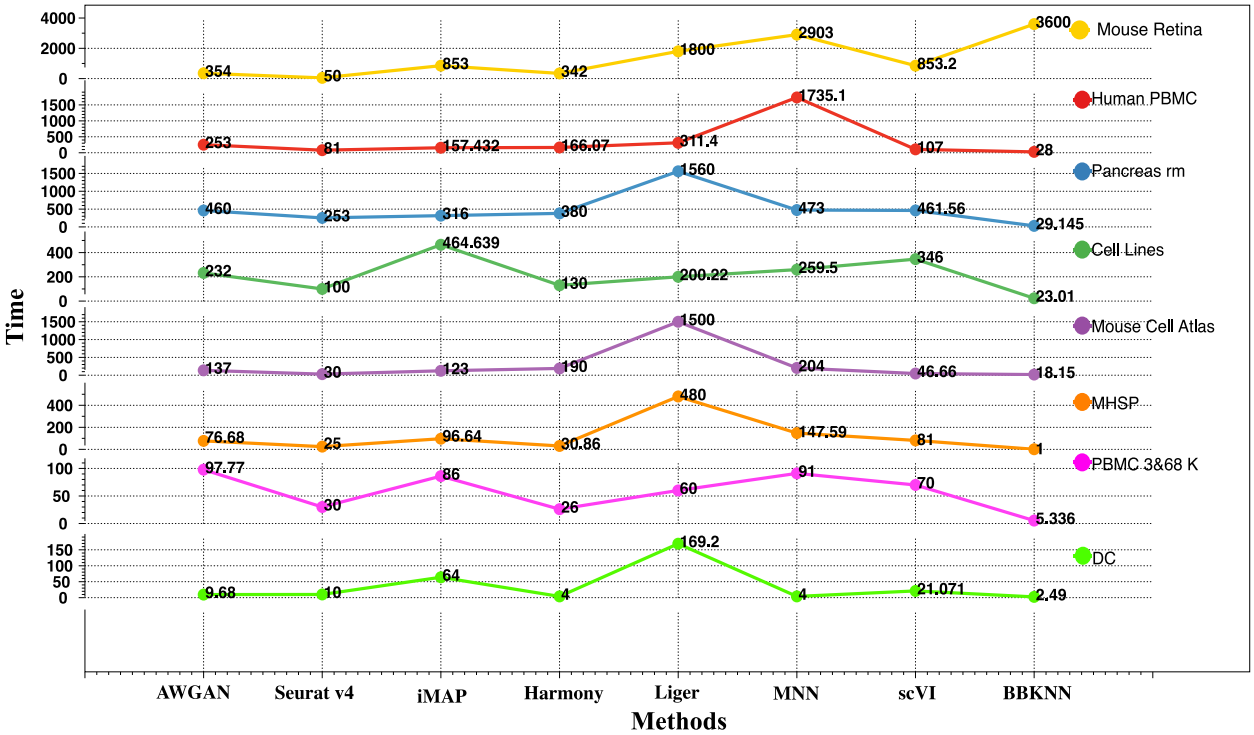


Figure 6: Running time comparisons across different datasets. We ordered the methods by the rank we calculated in Table 2 and ordered the datasets by their size. Methods

closer to the left have smaller overall rank in Table 2, and datasets closer to the bottom have smaller number of cells.

It can be seen that MNN and Liger are the two most time-consuming methods in some datasets, while BBKNN and Seurat v4 are the two fastest methods for running on most datasets. This might be because BBKNN and Seurat v4 do not involve large matrix decomposition as Liger does or complex matching learning process as other methods. As for AWGAN, its running time is relatively short in most datasets among all the machine-learning based methods (e.g. iMAP, Harmorny, and scVI).

2.3 Large-scale scRNA-seq Datasets

In this part, we consider two scRNA-seq datasets with more than 100,000 cells and investigate the ability of AWAGAN to integrate large-scale datasets. The first dataset is from the Human Cell Atlas [49], which is a large dataset containing more than 700,000 cells. The second dataset is a Mouse Brain dataset [52] [53], which contains over 800,000 cells. The analysis of large-scale dataset can take much space, memory, and time. Our model utilizes a more economical approach to handle large datasets. We show the results for the Human Cell Atlas dataset and the Mouse Brain dataset in this section.

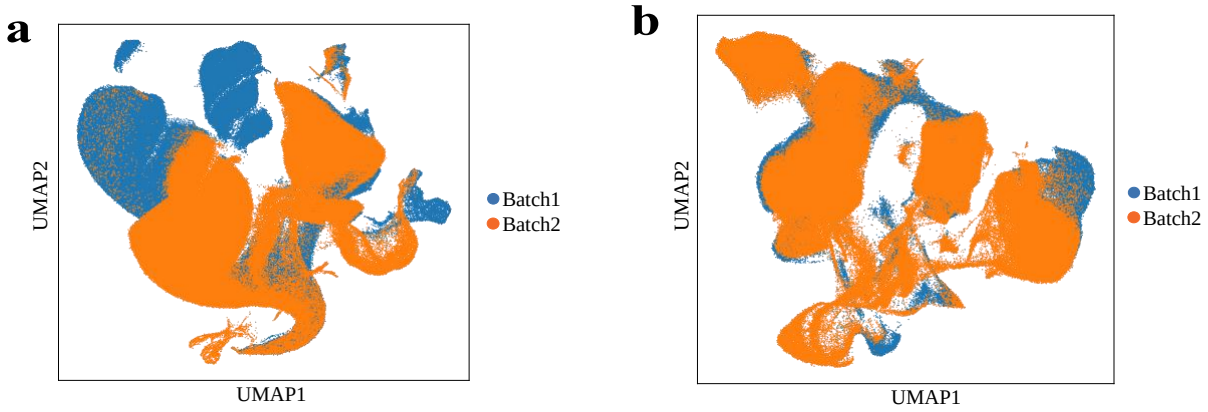


Figure 7: Human Cell Atlas dataset. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with batch label after batch effect removal.

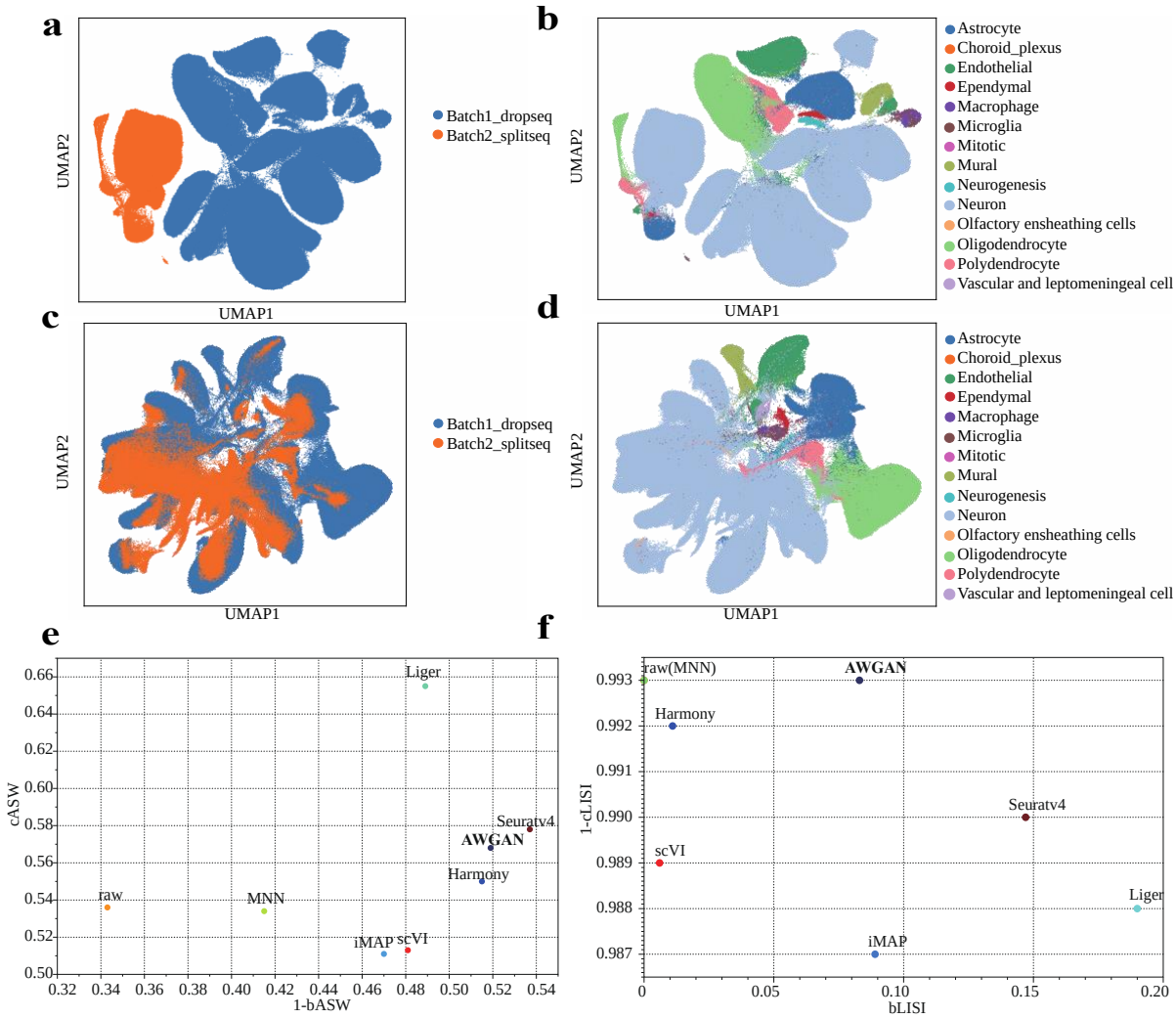


Figure 8: The performance of different batch effect removal methods for the Mouse Brain dataset. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with cell type label before batch effect removal. (c) Visualization for the data distribution with batch label after batch effect removal. (d) Visualization for the data distribution with cell type label after batch effect removal. (e) ASW assessments of different batch effect removal methods on the Mouse Brain dataset. (f) LISI assessments of different batch effect removal methods on the Mouse Brain dataset.

As can be seen from Figures 7 and 8, AWGAN also performs well for large-scale datasets. These two selected single-cell datasets have significant batch effects. Moreover, both of them have over 700,000 cells, making batch effect removal very challenging. To save time, we simplify the attention process for the two datasets. AWGAN is able to remove the batch effect for these two large-scale datasets, demonstrating its scalability to large datasets.

The running time and memory usage of an algorithm are important indicators of its efficiency, especially for models based on deep learning. We compare AWGAN with another GAN-based model, iMAP, from the perspective of resource utilization. The results are listed in Table 1.

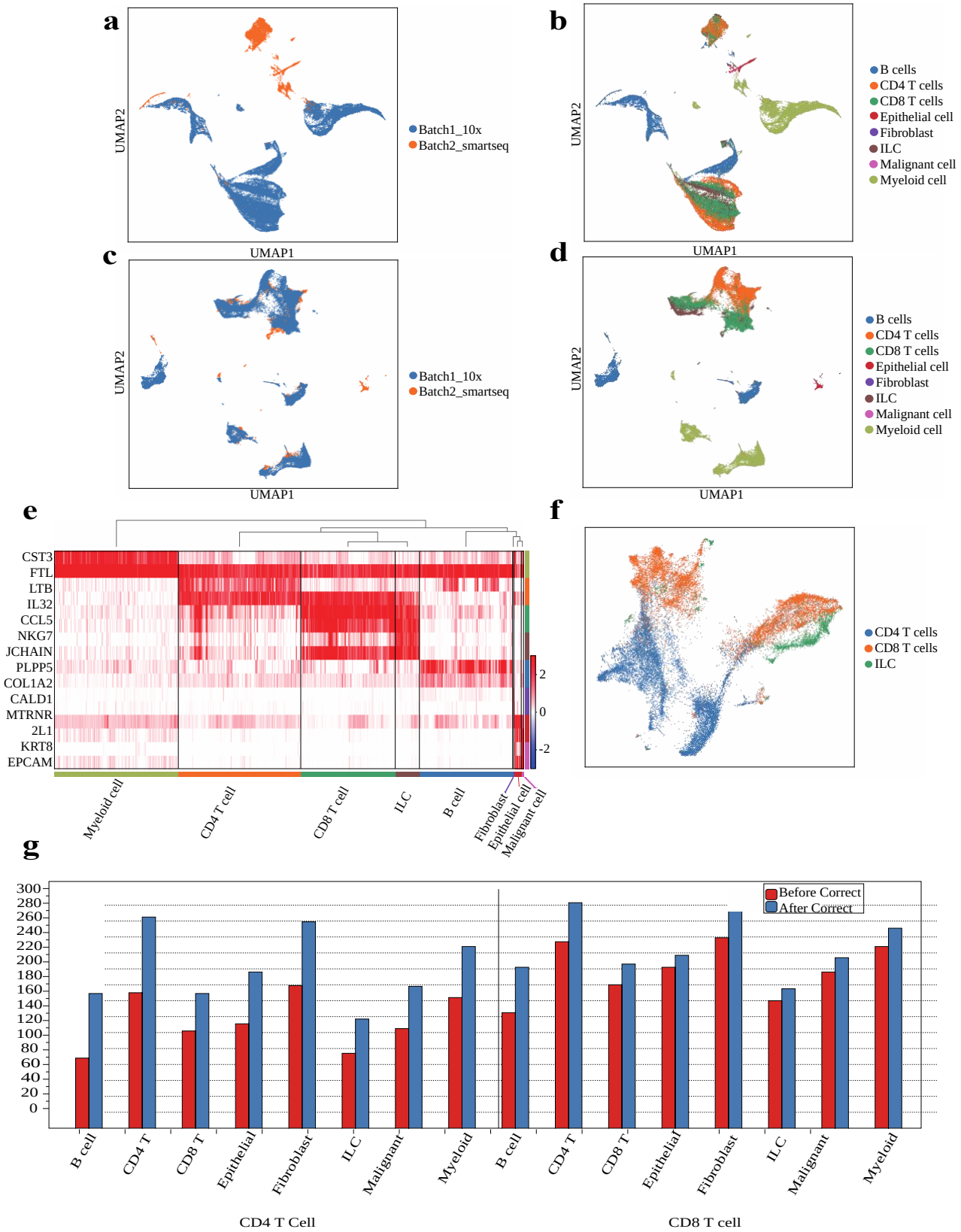
Datasets	Method	Running Time	Memory Usage
Mouse Brain	iMAP	7920 s (HPC)	39 GB
	AWGAN	3500 s	25 GB
Human Cell Atlas	iMAP	2040 s (HPC)	30 GB
	AWGAN	1800 s	20 GB

Table 1: Comparison on Running Time and Memory Usage between AWGAN and iMAP in the two large-scale datasets.

Note that for the Human Cell Atlas, iMAP failed on our cloud platform since in the step of finding the rwMNN pairs, memory usage was larger than available memory (25GB). As a result, running iMAP in a large-scale dataset requires larger memory than AWGAN. However, using AWGAN, our cloud platform can still generate the results because AWGAN only runs the attention step once to use memory more wisely in the computer, compared with iMAP.

2.4 Biological Characteristic Discovery based on AWGAN

The Colorectal Cancer (CRC) dataset is a scRNA-seq dataset containing tumor-infiltrating immune cells with 54,285 cells in total [26]. In this part, we took a deeper dive into the functional diversity of AWGAN by looking into the CRC dataset. Results shown in Figure 9 a-d suggest that AWGAN can mostly eliminate the batch effect in the CRC dataset and preserve the biological variation of the dataset. From Figure 9 d, we note that innate lymphoid cells (ILCs) and CD8 T cells are close to each other in the UMAP after removing batch effects. We assumed that the ILCs are transformed from CD8 T cells. To verify this part, in Figure 9 e, based on the heat map clustered by cell type labels, we found the gene expression amount between CD8 T cells and ILCs are very similar. Moreover, from Figure 9 f, a specific UMAP graph for CD4 T cells, CD8 T cells and ILCs, we also found that some ILC cells are mixed within the cluster of CD8 T cells. Therefore, it is possible that some ILCs may be incorrectly marked. This may be caused by the expression levels of some maker genes being too close or noises in the original data.


a
UMAP2
UMAP1
Batch1_10x
Batch2_smartseq
b
UMAP2
UMAP1
B cells
CD4 T cells
CD8 T cells
Epithelial cell
Fibroblast
ILC
Malignant cell
Myeloid cell
c
UMAP2
UMAP1
Batch1_10x
Batch2_smartseq
d
UMAP2
UMAP1
B cells
CD4 T cells
CD8 T cells
Epithelial cell
Fibroblast
ILC
Malignant cell
Myeloid cell
e
CST3
FTL
LTB
IL32
CCL5
NKG7
JCHAIN
PLPP5
COL1A2
CALD1
MTRNR
2L1
KRT8
EPCAM
2
0
-2
Myeloid cell
CD4 T cell
CD8 T cell
ILC
B cell
Fibroblast
Epithelial cell
Malignant cell
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CD8 T cells
ILC
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300
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240
220
200
180
160
140
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80
60
40
20
0
B cell
CD4 T
CD8 T
Epithelial
Fibroblast
ILC
Malignant
Myeloid
B cell
CD4 T
CD8 T
Epithelial
Fibroblast
ILC
Malignant
Myeloid
CD4 T Cell
CD8 T cell

Figure 9: Results on the CRC dataset. (a) Visualization for the data distribution with batch label before batch effect removal. (b) Visualization for the data distribution with cell type label before batch effect removal. (c) Visualization for the data distribution with batch label after batch17 effect removal. (d) Visualization for the data distribution with cell type label after batch effect removal. (e) Gene heatmap of the CRC dataset after batch effect removal. (f) Visualization for the misclassification data distribution with cell type label after batch effect removal. (g) Significant ligand-receptor pairs ($p < 0.05$) defined between all other cell types and T cells.

We also studied the interactions among different cell types. The two batches of the CRC dataset were sequenced based on different methods, and the two methods had their own characteristics. The 10X data (10x original data) are sparser but have higher throughput in terms of cell numbers, while the Smart-seq2 data measure more accurate gene expression levels [13,26,27]. Therefore, relying on the noisy 10X data only may makes it difficult to estimate correlation between cells and calculate cell-cell communication scores. AWGAN provides a potential solution to this problem by mapping the 10X data to the Smart-seq2 data. We used CellPhoneDB [28] to calculate cell-cell communication scores and generate visualization results. Based on Figure 5 in supplementary file 1 part 2, 10X data after batch correction (10x corrected data) could find some new ligand-receptor pairs, which could be defined as novel cell-cell interactions. For example, the pair XCL1-CCR5 between CD4 T cells and B cells could only be uncovered by 10x corrected data but not by 10x original data. Based on Figure 9 g, we found that the number of ligand-receptor pairs of the whole dataset increased after running the AWGAN correction. For more interesting pairs, please refer to supplementary file 1 part 2.

2.5 Benchmark Results Comparisons

It is difficult to evaluate and compare the performance of different models from visualization results, so we utilized four metrics to quantitatively evaluate AWGAN and other methods in removing batch effect: ASW, kBET, LISI, and Graph Connectivity. Moreover, since different metrics also have their own limitations, we need to consider results from the four metrics

together to judge the model's performance. For example, kBET value cannot accurately measure batch effect removal results in the presence of varying proportions of cell types in different batches [14]. In addition, the value of kBET is also very sensitive to its parameters as shown in supplementary file 1 part 3. Therefore, kBET may not be the only metric to evaluate different models. Instead, LISI, ASW, and Graph Connectivity can be considered together with kBET to provide comprehensive evaluations of the performance of different methods. The details of these four metrics can be found in the Methods section. In order to summarize the overall performance of the mth method for the dth dataset, we consider:

$$TotalRank(d, m) = \sum_{i \in M} Rank(d, i, m) \tag{1}$$

where we sum the rank of this method over different metrics in set M that consists of the four metrics discussed above. Therefore, a smaller $TotalRank$ for a batch effect removal method implies its better performance. Table 2 shows the results for different methods based on $TotalRank$.

	Total Rank Summary by Methods						
Datasets	AWGAN	Harmony	iMAP	Liger	MNN	scVI	Seurat v4
Human PBMC	**11**	19	16	**12**	20	24.5	**9.5**
Pancreas rm	**11**	**10**	**13**	18	29	22	14
Cell Lines	**15**	**19**	**9**	20	20	26	20
Mouse Cell Atlas	**13**	15.5	18	**14**	15.5	25	**14**
MHSP	**11.5**	17.5	**11**	19	18	22	**17**
PBMC 3&68K	**11.5**	**9.5**	17	23	24	24	**8**
DC	**11**	**7**	16	31	**13**	20	22
Mouse Retina	**14**	22.5	**14**	**13**	21	**12**	20
Mouse Brain	**14**	16	22	**11**	21.5	17.5	**10**
Final Total Rank	**112**	**136**	**136**	161	182	193	**134.5**
Overall Rank	**1**	**3.5**	**3.5**	5	6	7	**2**

Table 2: Total Rank Results. The total rank of the top three performers for each dataset (row) are highlighted in bold font.

When calculating the final total ranks, we did not consider the Human Cell Atlas dataset and the CRC dataset because the former dataset did not have available cell type labels and the latter dataset was mainly used to show the potential of AWGAN on downstream analysis. From table 2 we can see AWGAN performs better than other methods in the overall ranking and it is the only method that is always among the top three performers across all datasets. This result proves that AWGAN has a strong generalization ability.

Moreover, since Seurat v4 and iMAP are the two strong competitors in most datasets, we also conducted pairwise comparison between AWGAN and Seurat v4, and between AWGAN and iMAP on each dataset and with each metric (Figure 10).

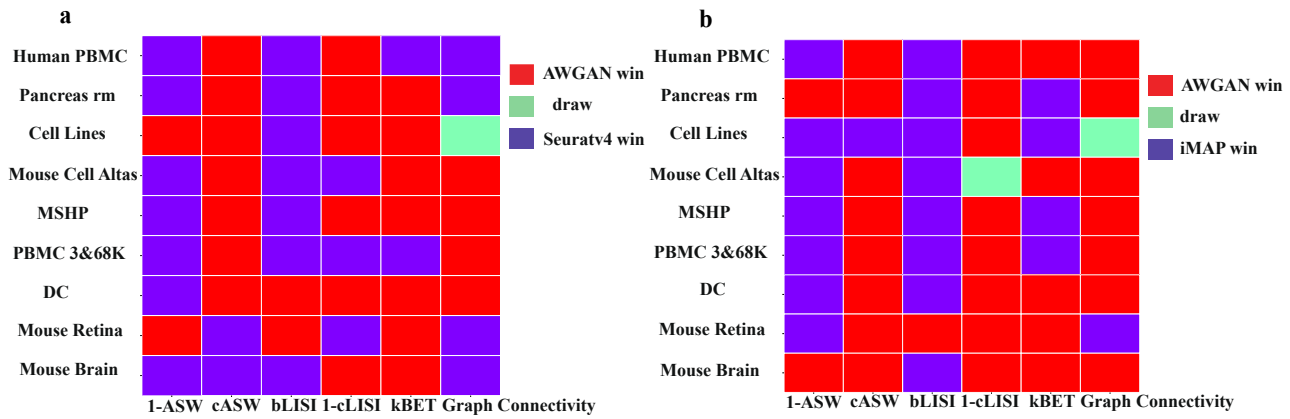

Figure 10: Comparison Heatmap. (a) Results for AWGAN vs. Seurat v4 in each dataset (row) and by each metric (column). (b) Results for AWGAN vs. iMAP in each dataset (row) and by each metric (column).

As we can observe from Figure 10 a, AWGAN performs better than Seurat v4 in terms of three metrics for cell type optimization (casw, 1-clisi, and Graph Connectivty) in the MHSP dataset, and the DC dataset. From Figure 10 b, AWGAN performs better than iMAP in terms of all these three metrics for cell type optimization in the Human PBMC dataset, the Pancreas dataset, the MHSP dataset, the PBMC 3&68K dataset, the DC dataset, and the Mouse Brain dataset. This explains why AWGAN has better overall performance in Table 2 than Seurat v4 and iMAP in datasets that contain a larger proportion of batch-specific cell types, such as the DC dataset, the MHSP dataset, and the Pancreas dataset. Moreover, AWGAN performs better

than iMAP in terms of nearly all the metrics in the Mouse Brain dataset, which suggests the advantage of AWGAN when dealing with such a large-scale dataset. Apart from better performance on preserving batch-specific cell types, the benefits of AWGAN in terms of lighter structure and less memory usage, compared with iMAP, are important factors for researchers to choose more suitable batch correction tools.

The detailed results for each of the four metrics are included in supplementary file 2. Overall, AWGAN performs well for both small-scale and large-scale datasets, especially for datasets with batch-specific cell types, for example, the DC dataset, the MHSP dataset, and the Pancreas dataset.

3 Discussion

In this paper, we have proposed and evaluated AWGAN, a novel model for removing batch effects. With a framework based on deep learning theory and generative models, AWGAN utilizes attention strategy to generate nearest neighbor pairs as a training set and has good performance both in terms of effectiveness and faster time compared with other methods.

More specifically, except for iMAP, Seurat v4 and Harmony, AWGAN achieves better results than the traditional batch effect removal algorithms in both ASW score and LISI score in most datasets. We note although both AWGAN and iMAP have advantages over other methods on different datasets, there are disadvantages of iMAP compared to AGWAN. Firstly, iMAP sometimes crashes (in 20% conditions, see example in Figure 6 in supplementary file 1 part 4), which is triggered by the improper neural network initialization. Secondly, according to Wasserstein Generative Adversarial Network with Gradient Penalty (WGAN-GP) [20], iMAP chooses the same value for the weight of gradient penalty (λ), to satisfy 1- Lipschitz condition, which can be interpreted as one condition to ensure the robustness of the generative model. However, WGAN-GP chooses this value based on experiments on image datasets, so it is not

suitable for iMAP to choose the same λ. AWGAN also has a similar design to WGAN-GP, but we find that we need to apply different λ for some datasets. Therefore, this affects iMAP performance. Thirdly, since iMAP needs to train five different neural networks in total, its efficiency cannot be guaranteed, and this design extends its running time for training the whole model especially when dealing with large-scale datasets. Finally, iMAP has poor performance on preserving batch specific cell types as we discussed in the last section of the benchmark results.

Some quantitative assessments of the performance of Seurat v4 are very favorable. However, Seurat v4 suffers from the drawback of not considering the conservation of cell type labels. For example, it ignores biological variation and overemphasizes data integration function. This drawback may negatively impacts biological research. We can see from Figure 7 in supplementary file 1 part 4 that in the DC dataset, CD1C cells are mapped to another type of cells, which is not desirable. In addition, we can also observe that in the Cell Lines dataset, after processing based on Seurat v4, the cells with the label ”Jurkat” still have batch effect from Figure 8 in supplementary file 1 part 4. Harmony also has issues related to batch-specific cell types, shown in Figure 9 in supplementary file 1 part 4. So overall, AWGAN performs better than Seurat v4 and Harmony with better visualization results.

Moreover, for batch correction methods like scVI, BBKNN, and Liger, there are some issues in the method design principles. For scVI, it extracts the gene expression statistics as an embedding in another space, so we need to regenerate the count matrix through a decoding model. Therefore, this variational inference method is even more complex. For BBKNN, it is similar to the neighbors function in Scanpy [25], which can only make the visualization result better. BBKNN does not process the raw data, so it cannot contribute much to downstream analysis. Liger has similar issues as it only returns integration results in a low dimensional space. Therefore, we cannot directly obtain corrected gene expression profiles in the original feature

space from Liger. In contrast, methods like AWGAN, MNN, Seurat v4, and iMAP can adjust the original gene expression data and provide a better basis for downstream analysis.

For large-scale data, AWGAN can overcome the memory limitation problems and finish the batch correction task without collapse error. In addition, in the research related to the CRC dataset, AWGAN shows its ability to identify potentially misclassified cells and uncover the hidden cell-cell interactions. Overall, in most cases, the performance of AWGAN on the datasets mentioned in this paper is better than the other approaches.

4 Conclusion

We demonstrate that AWGAN is a valuable method for batch effect removal through experiments on ten different real-world datasets in comparison with seven different batch effect removal methods using four metrics. It takes the advantages of deep learning models and uses relatively few resources to achieve better results. Firstly, AWGAN can remove the batch effect in different datasets and preserve the biological variation (for example, gene expression difference and cell type distribution), while other state-of-the-art models, for example, iMAP, Harmony, and scVI, fail to keep it in some conditions. Secondly, our model supports various improvement extension applications, and researchers can adopt many hyper-parameters in our model. The strong adaptability of our model also allows it to be improved to achieve better results. Thirdly, running our model does not require very high technique or advanced device. Researchers can choose to use either cloud platform or CPU core in a personal computer to train our model and get a satisfactory result. We believe that the output of our model can better inform downstream analyses, such as specific cell function analysis, cell heterogeneity analysis, and others.

5 Methods

5.1 Attention-driven Data Pre-processing

We observed that most studies on batch effect removal chose the gene number of one cell within the range of at least 200 and up to 2000 ($N \in (200,2000)$) [13,29] based on the highly variable gene (HVG) filtering method designed in Seurat v4 [11]. In this paper, we follow the above rules and process the raw count matrix. After obtaining the filtered count matrix, we proceed to the step of generating the training set. In this paper, inspired by the basic attention mechanism, we construct nearest neighbor cell pairs using cosine distance. For the set of cells with larger cosine similarity, we consider them as the nearest cell pair, and their attention to each other is 1, while the attention of the rest of the cells to them is 0. This type of relationship of attention is not transmissive but satisfies the principle of superposition. Therefore, any cell in the query dataset could find its own nearest neighbor cell in the reference dataset, generating a neighbor cell pair. The definition of nearest neighbor cell pairs can be explained as the Figure 10 in supplementary file 1 part 5.

To find the NNP, we only need to solve the following optimization problem. Assume there are two scRNA-seq datasets $Data_{true}$ (usually reference dataset) and $Data_{false}$ (usually combined with all the extra batch data, which is defined as query dataset), our target is to find:

$$(C_1^*, C_2^*) \in Pair = argmax_{C_i,C_j} CosineSimilarity(C_i, C_j) \tag{3}$$

$$C_i \in Data_{true}, C_j \in Data_{false} \tag{4}$$

We utilize the iteration algorithm with Numba acceleration [30] to solve this problem and collect the ideal solution. Here we declare that our algorithm always prefers to find cell pars with the largest cosine similarity (based on cosine value of two vectors). Based on this rule, sometimes NNP may not choose cells with the same cell type label, but since this situation is very rare and to save time for running the attention matching algorithm, we choose this

mechanism (other methods like MNN [10], iMAP [13] also ignore the effect of cell types). NNP with different cell type labels has little effect on the normal operation of the model, and we verify this in part 5.2. The algorithm for finding the target pairs is shown in Algorithm 1.

We also considered some variations in this algorithm, including selecting more pairs and considering the construction of training sets after averaging multiple attention pairs. However, the above selection effects were not as good as the algorithm in Algorithm 1. Details of the experiment can be found in the supplementary file 1 part 6.

Algorithm 1 NNP Generating Algorithm. This algorithm is based on iterative approach and optimal solution searching approach. Its target is to generate the NNPs.

Input: Input parameters: Reference dataset $Data_{ref}$; Query dataset $Data_{que}$
Ouput: Output parameters: Training dataset $Data_{true}$; Training dataset $Data_{false}$.

```
INIT: initialize all parameters.
for item1 in Data_que do
    MaxSim = -inf; Q = None; R=None;
    for item2 in Data_ref do
        sim = CosineSimilarity(item1,item2)
        if sim > MaxSim then
            MaxSim = sim
            Q = item1
            R = item2
    Data_true.append(R)
    Data_false.append(Q)
Return Data_true, Data_false
```

5.2 AWGAN Training

Description of notations used in this section can be found in table 3 and the structure of AWGAN is shown in the table 4. For the generator model, we apply the batch normalization [31] technique to accelerate the training speed, and we also utilize the Dropout method [32] based on momentum to prevent the model from overfitting. Based on the requirement in WGAN-GP [20], we remove the BatchNorm layer for the discriminator model because we

intend to ensure the smooth requirement for Lipschitz condition could not be affected by batch normalization layers.

Notation	Description
G	Generator Model in AWGAN
D	Discriminator Model in AWGAN
L	Total loss for AWGAN
L_G	Loss in Generator Training
L_D	Loss in Discriminator Training
$Data_{false}$	Data need to remove the batch effect
$Data_{true}$	Data without the batch effect (reference dataset)
$Data_{fixed}$	Data after batch effect correction
$Pair$	Pair set after running the attention method
λ	Coefficient for gradient penalty
P_g	Distribution of generator data
P_r	Distribution of real data
P_x	Distribution of interpolation data
N	Selected number of genes

Table 3: Notation List

Generator		Discriminator	
Layer Type	#	Layer Type	#
Input Layer	N	Input Layer	N
Hidden Layer	1024	Hidden Layer	1024
Batch Norm	1024	Hidden Layer	512
Hidden Layer	512	Output Layer	1
Batch Norm	512	-	-
Hidden Layer	1024	-	-
Batch Norm	1024	-	-
Output Layer	N	-	-

Table 4: AWGAN Structure

The minimization optimization target of our model can be concluded as :

$$L = E_{x^* \sim P_g}[D(x^*)] - E_{x^0 \sim P_r}[D(x^0)] + \lambda E_{x \sim P_x}[(\mathrm{k}\nabla_x D(x)\mathrm{k}_2 - K)^2] \quad (5)$$

The first two terms mean the Wasserstein Distance, which is the original expression in WGAN that we intend to minimize. However, if we minimize this value directly, the K-Lipschitz

condition may not be satisfied. To satisfy the K-Lipschitz condition, the original WGAN uses a direct parameter range limitation. However, in the training progresses, the parameters will spontaneously concentrate on the boundary, so the gradient will not be well passed back to the model. Therefore, we introduce the third term and the gradient penalty mechanism [20]. The third term, gradient penalty, will first interpolate the sampling between true data and fake data and obtain a new distribution. In this problem, proved by the WGAN-GP paper, we take $K = 1$ because the L2 norm of the ideal gradient for WGAN is 1 (1-Lipschitz condition) [20]. We then use this new distribution to compute the gradient penalty to match the Lipschitz condition over as large as possible space, thus eliminating the negative effects of parameters on the boundary. Some researchers agree with the idea to set K = 0, which is more reasonable [33]. We provide our experiment results with this choice in the supplementary file 1 part 7 and show that there is no significant difference between these two choices. The interpolation function between the $True$ dataset and $Fake$ dataset (including raw Fake data and temporary $G(Fake)$ data) is defined as:

$$\begin{cases} Interpolation(True, Fake) = \epsilon \cdot True + (1-\epsilon) \cdot Fake \\ \epsilon \sim Uniform(0,1) \\ x \sim P_x(Interpolation(True, Fake)) \end{cases} \tag{6}$$

For the choice of the activation function, we utilize the state-of-the-art activation function: Mish [34] to construct our model. The expression of Mish function can be defined as:

$$Mish(x) = x \cdot tanh(ln(1 + e^x)) \tag{7}$$

where $f(x) = ln(1 + e^x)$ is also known as SoftPlus function. The Mish function has smoother gradient compared with the Rectified Linear Unit (ReLU) [35] function, and smoother activation function allows better information integration and information mining performed by the deep neural network [34].

For the output layer of the generator, we adopt the ReLU function with ResNet structure [36], which is also known as identity shortcut connection, to generate more stable results. To implement the ResNet structure, we change the expression of output layer into this form:

$$Y = T(F(X,\{W_i\}) + W_sX) \qquad T,F \in set(Activation \quad Functions) \tag{8}$$

where $\{W_i\}$ means the set of inner weight matrix. W_s is a linear map which helps to ensure the feature dimensions of X and F(X) are consistent. In this problem, their feature dimensions are the same, so W_s can be treated as identity mapping. To make our model perform better in different dataset situations, we define a new function for output layer, the transformation function T. For datasets which originally utilize log preprocessing method, we set $T = 1$, otherwise, $T = ReLU()$.

From the perspective of the back-propagation algorithm, we always meet non-zero terms when we are trying to find gradients based on ResNet structure, so the problem of gradient vanishing and white noise (very small gradient) can be solved to some extent [37].

For the overall formal design of the generator, inspired by the encoder-decoder model, we also add features such as embedding generation and embedding decoding in our network, which can extract the biological variation in scRNA-seq data.

In most of the time, since cells are highly differentiated, there are significant differences in gene expressions between different types of cells, so theoretically, it is easy for us to distinguish batch specific cell types [38,39]. However, sometimes there may exist cells with similar gene expression levels but different cell identification. For such problem, we utilize the Discriminator and Dropout method design to help keep the balance between mixing the cells in different batches and discerning batch specific cell types. Firstly, the discriminator is designed to identify the cells in different batches. For batch specific cell types, the discriminator is able to quickly treat them as cells in other batches, and the model will not

evaluate this situation. Secondly, mixing these cells into known cells can be treated as an overfitting problem. The dropout method is designed to solve this kind of problems. Moreover, adjusting the training epochs and batch sizes can also generate a better result, depending on the chosen dataset. The Figure 11 supplementary file 1 part 5 illustrates the function of discriminator.

In total, the training algorithm is shown in Algorithm 2.

Algorithm 2 Training process with AdamW [40] optimization method for AWGAN. The training epoch is the number of training iterations, and the batch size set for this model is k. They are all hyper parameters.

Input: Input parameters: Model D,G; Dataset $Data_{false}$, $Data_{true}$; Batch marker set $Batch$.
Ouput: Output parameters: Model G; Dataset $Data_{true}$

```
INIT: initialize all parameters.
for Type in Batch do
    Data_batch = Data_false[Type]
    for number of training iterations do
        for k steps do
            Refer = Data_true (sample with batchsize=k)
            Query = Data_batch (sample with batchsize=k)
            L_D = -E[D(Refer)]
            L_G = E[D(G(Query))]
            x = Interpolation(Refer,Query)
            Div = E[(||∇_x D(x)||_2 - 1)^2]
            L = L_D + L_G + Div/λ
            AdamW(D, L) # Training D
            L_G = -E[D(G(Query))]
            AdamW(G, L_G) # Training G
            if G is not convergent then
                Adjust λ = λ_new
                goto INIT
    Data_fixed = G(Data_batch)
    Data_true = concat(Data_true,Data_fixed)
Return G, Data_true
```

5.3 Model Adjustment and Evaluation

There are five kinds of parameters that can be adjusted in our model: epoch, batch size, gradient penalty, dropout rate, and learning rate. Moreover, for the choice in calculating the gradient penalty, we prefer using raw $Fake$ data for the first attempt. If the results are not satisfactory, we shift to temporary $G(Fake)$ data. The strategy we choose to improve the performance of our model is based on both the visualization result and quantitative assessments (especially ASW rate and LISI rate). We also refer from the stage2 of iMAP's structure for network design [13]. Finally, we choose the combination of different parameters which could optimize the quantitative rates and visualization performance. The choice for all key hyperparameters can be found in supplementary file 1 part 8. Moreover, for all the datasets, we run our model 10 times in each dataset to ensure that our output converge and is stable.

To evaluate AWGAN and other benchmark methods except for Liger and BBKNN, we used the top 20 PCs [29] based on the PCA method [41] with batch information and cell type information of the processed data as the input part for all the four quantitative metrics. For Liger, we chose to evaluate the norm matrix of the data generated by Liger with the original batch and cell type information. For BBKNN, since this method does not generate either gene expression information or latent space information, we did not compute metrics on this method. To evaluate batch optimization task more reliably, we removed the batch specific cell types when we generated the metrics related to this part. The visualization of quantitative comparisons can be found in the 'Results' section and the detailed quantitative results can be found in the supplementary file 2.

5.4 Metrics Details

- ASW: Average silhouette width, also known as ASW, is computed based on cluster and average distance. For one data point, we first calculate the difference between the average distance between this point and the other points in a neighboring cluster to the average distance between this point and other members in the same cluster. After finishing this step, we divide this result by the larger distance to get the silhouette value. We repeat the process for each data point for a certain number of iteration times and take the average across them. More specifically, ASW is defined as:

$$ASW = \frac{(d_{neighbors} - d_{members})}{max(d_{neighbors}, d_{members})} \tag{9}$$

  Since we normally have two kinds of clusters: batch and cell type, we use the F1 score of ASW to evaluate our model. The F1 score of ASW is defined as:

$$F1_{ASW} = \frac{2(1 - ASW_{batch})(ASW_{celltype})}{1 - ASW_{batch} + ASW_{celltype}} \tag{10}$$

  For the F1 score, since we intend to have different batches integrated in one reference dataset, and keep the cell type distribution to be the same, so the model with low ASW_{batch} rate and high $ASW_{celltype}$ rate will be considered as having good performance. Therefore, a higher F1 score suggests a better performing model.

- kBET: k-Nearest Neighbor (kNN) batch-effect test, as known as kBET, utilizes kNN-based algorithm to measure the results of batch effect removal. The algorithm randomly selects some data points and tests whether the local batch label distribution among them is similar to the global distribution. Ideally, the batch label distribution in the sample and the global data distribution should be very close to each other, so the rejection rate should be 0. In fact, there are some differences between the two, so we get an actual rejection rate between 0 and 1. To be consistent with the rest of the methods, we define the acceptance rate as:

$$AcceptanceRate = 1 - RejectionRate \quad (11)$$

A higher acceptance rate of one model suggests better performance of this model.

For large-scale datasets, we find the rejection rates are often too large to compare across different batch correction methods and the kNN searching process often fails because of memory issues, so we adopt two strategies: neighborhood size adjustment and subsampling to overcome this obstacle.

The parameter that controls neighborhood size is $heuristics$. If it is set to be $True$, kBET will return the optimal neighborhood size that can generate the maximal rejection rate. However, for large-scale datasets, this value is very large under $heuristics = True$ so that the differences between different methods' rejection rates are not significant. Therefore, we set this value to $False$ so that kBET can automatically choose 25% of the cells in the dataset to calculate the rejection rate, which amplifies the difference among different methods' rejection rates.

For subsampling, if we use n to represent the sample size and k to represent the neighborhood size, and if $n * k > 2^{31}$, the kNN algorithm will stop searching in R, which is often the case for large-scale datasets [24]. Therefore, for large-scale datasets, we split each of them into ten subsamples and calculate the overall rejection rate based on the average of the 10 rates from the subsamples.

- LISI: Local inverse Simpson's index (LISI) adopts similar principle to kBET, but the rule for choosing the neighbors in kNN is different from kBET. LISI chooses neighbors based on the local distribution of distances with an invariant perplexity. Moreover, the chosen neighbors will be used to calculate inverse Simpson's index, which is the number of accepting cells in neighbors. The definition of Inverse Simpson's rate is:

$$ISR = \frac{1}{\sum_{b=1}^{B} p(b)} \tag{12}$$

where b in the cell index and B is the cluster of one batch. ISR is the batch probabilities in local neighbors. Since LISI assigns a value for every different cell, we evaluate our model based on the average LISI rate. If it approaches the real batch number, the model with such LISI is better than the others. According to the definition of the LISI rate, we define two metrics: cell type LISI rate (cLISI) and batch LISI rate (bLISI). Our goal is to increase the bLISI rate while decreasing the cLISI rate. Therefore, we utilize the F1 score of LISI to evaluate different methods. The F1 score of LISI is defined as:

$$F1_{LISI} = \frac{2(1 - cLISI)(bLISI)}{1 - cLISI + bLISI} \tag{13}$$

- Graph Connectivity: This method is based on kNN and its target is to evaluate the conservation of cell type distribution after batch correction [5]. For the kNN graph G, of one set of integrated data, we evaluate whether cells with cell type label c are directly connected with each other across batches. Moreover, we could create the subset kNN graph $G(N_c,E_c)$ only for the cluster with cell type label c. Let T represent the set containing all cell type labels, we can use this equation to compute the graph connectivity score:

$$gc = \frac{1}{|T|} \sum_{c \in T} \frac{|LCC\left(G\left(N_c, E_c\right)\right)|}{|N_c|} \tag{14}$$

In this equation, $|LCC()|$ means the number of nodes in the largest connected component (LCC) of the given graph, and $|N_c|$ means the number of nodes with cell type label c. Based on the concept of LCC, higher graph connectivity score means better cell type optimization performance of a method.

5.5 Device and Software Package Information

The CPU of our device is Intel(R) Xeon(R), 2.30 GHz. The GPU of our device is NVIDA Tesla P100, the RAM is 25GB, and the hard disk size is 140GB. The version of Pytorch is 1.7.0 [42]. To run Liger and Seurat v4 in R, we also utilized a device with CPU: Intel(R) Core(TM) i5, 2.50GHz, RAM:12GB. To run iMAP on the human cell atlas dataset, iMAP, MNN, Harmony, Seurat v4 and Liger on mouse retina dataset, and all methods on mouse brain dataset, we used a HPC tool. To make life easier, the Colab platform under Google can meet most needs of running our code. The versions of software packages we used can be found in supplementary file 1 part 9.

5.6 Dataset Information

We considered eleven single-cell datasets to investigate the performance of AWGAN in comparisons with other methods, including seven small-scale datasets and four large-scale datasets. The Cell Lines dataset [43] was generated by 10x Genomics with three different batches. The DC dataset [44] is composed of two different batches based on the Smart-seq2 protocol. The Pancreas rm dataset [13] includes human pancreatic cells measured by 5 different platforms and is loaded from R package: SeuratData. The PBMC 3&68K dataset [43] was also generated by 10x Genomics, and PBMC 68K is the processed version. The MHSP dataset [45] [46] was acquired by researchers using the Mars-seq protocol and the Smart-seq2 protocol. The Mouse Cell Atlas dataset [47] [48] contains two batches from the Microwell-seq and the Smart-seq2 protocol. The Human PBMC dataset [43] was generated by 10x Genomics with two batches. The Human Cell Atlas dataset [49] is a large dataset with bone marrow cells in batch 1 and cord blood-derived cells in batch 2 based on the 10x Genomics protocol. The Mouse Retina dataset [50] [51] is composed of mouse retina data generated using the Drop-seq technology by two unassociated laboratories. The Mouse Brain dataset [52] [53] is composed of two datasets from two different research groups, which were generated by Drop-seq and SPLiT-seq protocols separately. The CRC dataset [26] is a scRNA-seq dataset

collected from 18 CRC patients using both Smart-seq2 and 10x Genomics. Moreover, the cell type composition of each dataset can be found in supplementary file 3.

5.7 Code Availability

An open-source implementation of the AWGAN and source code to reproduce the results can be downloaded from: https://github.com/HelloLTY/AWGAN.

5.8 Data Availability

Download procedures for the datasets used in this paper can be found in the supplementary file 4.

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