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# Joint Sample Position Based Noise Filtering and Mean Shift Clustering for Imbalanced Classification Learning

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Abstract: The problem of imbalanced data classification learning has received much attention. Conventional classification algorithms are susceptible to data skew to favor majority samples and ignore minority samples. Majority weighted minority oversampling technique (MWMOTE) is an effective approach to solve this problem, however, it may suffer from the shortcomings of inadequate noise filtering and synthesizing the same samples as the original minority data. To this end, we propose an improved MWMOTE method named joint sample position based noise filtering and mean shift clustering (SPMSC) to solve these problems. Firstly, in order to effectively eliminate the effect of noisy samples, SPMSC uses a new noise filtering mechanism to determine whether a minority sample is noisy or not based on its position and distribution relative to the majority sample. Note that MWMOTE may generate duplicate samples, we then employ the mean shift algorithm to cluster minority samples to reduce synthetic replicate samples. Finally, data cleaning is performed on the processed data to further eliminate class overlap. Experiments on extensive benchmark datasets demonstrate the effectiveness of SPMSC compared with other sampling methods.

Key words: imbalanced data classification; oversampling; noise filtering; clustering

## 1 Introduction

Imbalanced data classification is an important research topic in machine learning<sup>[1]</sup>. Here, imbalanced data mean that the number of samples in some classes is far larger than the number of samples in others in a dataset. For example, in two classes of imbalanced data, the class with most samples is called majority class and the class with few samples is called minority class. In many practical applications, the minority class samples have significant research value because they contain important information, such as rare disease diagnosis<sup>[2–4]</sup>, fraudulent transaction detection<sup>[5]</sup>, DNA microarray data analysis<sup>[6]</sup>, text classification<sup>[7]</sup>, network intrusion detection<sup>[8, 9]</sup>, security management<sup>[10]</sup>, etc.

Traditional classification algorithms aim to improve the overall classification accuracy. However, in the imbalanced data scenario, since conventional classification algorithms usually favor majority class samples<sup>[11]</sup> and ignore minority class samples, directly applying classifiers may result in poor performance<sup>[12, 13]</sup>. In particular, even a high classification accuracy is obtained, it is not reliable. For example, in disease diagnosis, if the number of diseased samples is only 1% and the number of normal samples is 99% in a dataset, then the accuracy of classifying all samples as normal will be as high as 99%, but this accuracy is not reliable because the minority samples are not correctly identified and the cost of this misclassification is huge. Research has shown that not only between-class imbalance decreases classification performance, but also overlapping between classes<sup>[14, 15]</sup>, noisy samples<sup>[16]</sup>, small disjuncts<sup>[17, 18]</sup>, within-class

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imbalance<sup>[19]</sup>, and duplicate data<sup>[20]</sup> can also decrease classification performance.

To solve the above problems, many approaches have been proposed, which can be classified into four categories: data level methods<sup>[21, 22]</sup>, algorithm modification<sup>[23]</sup>, cost-sensitive learning<sup>[24, 25]</sup>, and ensemble learning<sup>[26]</sup>. Specifically, data level methods rebalance the dataset using oversampling, undersampling, or hybrid sampling methods. Algorithm modification is to improve the existing algorithms to recognize minority class samples more accurately. Costsensitive learning sets a larger cost for misclassifying minority classes, and ensemble learning improves classification performance by combining several basic classifiers. Among them, the data level method is favored by researchers because it only changes the distribution of the original data, is convenient to use, and can be directly applied to various classifiers.

The most common methods in the data level are random oversampling<sup>[17]</sup> and random undersampling<sup>[27]</sup>. The former randomly duplicates the minority class samples and the latter randomly reduces the majority class samples to rebalance the dataset. However, random oversampling is susceptible to overfitting<sup>[28]</sup> and random undersampling may remove samples containing important information<sup>[29]</sup>. Although both methods have their advantages, studies have shown that oversampling is preferred to undersampling in many practical applications<sup>[30, 31]</sup>. To overcome the drawbacks of oversampling methods, many modified oversampling methods have been proposed, such as synthetic minority oversampling technique (SMOTE)<sup>[32]</sup>, adaptive synthetic sampling approach (ADASYN)<sup>[33]</sup>, K-mean-SMOTE<sup>[34]</sup>, majority weighted minority oversampling technique (MWMOTE)<sup>[35]</sup>, borderline-SMOTE (B1-SMOTE and B2-SMOTE)<sup>[36]</sup>, density-based synthetic minority oversampling technique (DBSMOTE)<sup>[37]</sup>, etc. Although these methods improve the recognition accuracy of minority classes, they also have some problems such as insufficient noise filtering and generation of duplicates and outlier samples. To this end, we propose a joint sample position based noise filtering and mean shift clustering (SPMSC) method for imbalanced binary data in this paper. SPMSC can not only adequately filter the noise samples to alleviate their influence in the sample generation process, but also effectively reduce the production of duplicate samples and class overlap, so as to improve the recognition accuracy of samples.

The main contributions of this work can be summarized as follows.

• We propose a new noise filtering mechanism that can adequately filter the noise samples in the original dataset to weaken the effect of noise in the sample synthesis phase.

• We cluster minority samples using the mean shift method, which does not need to set the number of clusters in advance. Also, it does not lead to the generation of a large number of duplicate samples at the sample synthesis phase due to inappropriate distance thresholds as in the case of hierarchical clustering.

• We use the Tomek link data cleaning method after sample synthesis to further reduce class overlap in the processed dataset.

The remainder of this paper is organized as follows. We review some popular oversampling methods in Section 2. In Section 3, we describe the SPMSC method in detail. Experimental results and analysis are provided in Section 4. Finally, we conclude this paper in Section 5.

### 2 Related Work

The essence of sampling is to add or remove resamples to rebalance the originally imbalanced data. In this section, we briefly review some of the popular resampling methods.

SMOTE is one of the most representative oversampling methods. As shown in Fig. 1a, in Ref. [32], Chawla et al. randomly selected a sample y from the k nearest neighbors of minority sample x' to synthesize a new sample by Eq. (1):

synthetic = 
$$x' + \lambda \times (y - x')$$
 (1)

where  $\lambda$  is a random number between [0, 1]. This method is widely used because it is easy to understand and implement, however, it also has shortcomings. As shown in Fig. 1b, since this method performs sample synthesis for all minority samples without considering their distribution information relative to majority samples, it generates noisy samples and class overlap. Many approaches have been proposed to solve the problems generated by SMOTE. In Ref. [33], He et al. assigned sampling weights to each minority sample based on the number of majority samples in its nearest neighbors. However, it has the shortcomings of ignoring the effects of noisy samples and assigning unreasonable sampling weights. As shown in Fig. 2a, assuming k is 5, the noisy sample A will be assigned a larger weight, and the minority sample B will be assigned the same weight as the minority sample C although it is closer to the



(a) Random linear interpolation by using SMOTE

Sample synthesis. Fig. 1



(a) An example of the problems caused by ADASYN and borderline-SMOTE

(b) An example of the problems caused by safe-level-SMOTE



decision boundary. In Ref. [36], Han et al. proposed a method to divide minority samples into three classes: noise, safe, and danger, and to oversample only the samples in the danger. However, this method may not accurately identify the danger samples in some cases, as shown in Fig. 2a, sample B will be misclassified as safe class although it is a danger sample. Bunkhumpornpat et al.<sup>[38]</sup> assigned to each minority sample "safe-level" value, and then synthesized samples closer to the largest safe-level. However, as shown in Fig. 2b, in contrast to the minority sample F, the new samples synthesized by the minority samples D and E will be closer to themselves, which may lead to overfitting because these new samples are gathered around minority class samples with a large density and away from the decision boundary. In Ref. [22], Onan proposed a consensus clustering based on undersampling, which utilizes the consensus clustering mechanism to undersample the majority class samples and improves the classification accuracy. In Ref. [25], Jiang et al. proposed to change the class distribution of the training data by cloning minority class samples. Barua et al.<sup>[35]</sup> proposed a new method to identify the boundary minority samples and assign sampling weights, however, it has the disadvantages of inadequate noise filtering and duplication of generated samples with the original minority data. In Ref. [34], Douzas et al. proposed to firstly cluster the whole dataset by using K-means, then select the appropriate clusters according to the imbalance ratio, and finally use SMOTE to synthesize samples. However, the optimal number of clusters is difficult to find. Nekooeimehr and Lai-Yuen<sup>[39]</sup> proposed adaptive semi-unsupervised weighted oversampling (A-SUWO) method which uses a semi-unsupervised hierarchical clustering method to cluster minority samples, then uses misclassification errors and cross-validation to determine the number of samples to be synthesized in each subcluster, and finally assigns sampling weights based on the average distance of the minority samples from their nearest majority class neighbors. However, the method is more complex and may not be suitable for larger datasets.

# **3** Proposed Method

In this section, we propose a new method for imbalanced data classification. Our method consists of three main steps: (1) noisy sample filtering, (2) mean shift clustering and sample synthesis, and (3) data cleaning.

# 3.1 Preliminary

Mean shift is a center-based nonparametric clustering method. The method works by calculating the average value of the distance between a candidate point  $x_i$  and the points within a given radius r and then updating the position of  $x_i$ , which forms a cluster with the points within its radius when  $x_i$  is not moving. The shift vector m is calculated by Eq. (2):

$$m(x_i) = \frac{\sum_{x_j \in N(x_i)} K(x_j - x_i) x_j}{\sum_{x_i \in N(x_i)} K(x_j - x_i)} - x_i \qquad (2)$$

where *m* points to the region with the largest density increase, and  $N(x_i)$  represents the neighborhood of  $x_i$  in a given *r* range. According to the shift vector *m*, the update process of a candidate point  $x_i$  is shown in Eq. (3).

$$x_i^{t+1} = x_i^t + m(x_i^t)$$
(3)

where t is the number of iterations needed. The algorithm does not require setting the number of clusters beforehand, which can be set automatically without relying on the parameter bandwidth that indicates the range of the region to be searched, and it can handle clusters of arbitrary shape. Therefore, the mean shift algorithm is suitable for the segmentation of data. For more information about the use of this algorithm, please refer to sklearn (https://scikit-learn.org/stable/).

# 3.2 Our method

MWMOTE<sup>[35]</sup> is a popular oversampling method for processing imbalanced data problems. Although it is specific for some problems such as within-class imbalance and class overlap, it also has the shortcomings of inadequate noise filtering and generating duplicate samples. Motivated by this, we propose a new approach to cope with these problems.

### 3.2.1 Noise sample filtering

At present, many algorithms remove noise based on K-NearestNeighbor (KNN) noise filtering criterion, i.e., if all k nearest neighbors of a minority sample are other classes, then the minority sample is considered as a noisy sample. However, this noise removal method is difficult to eliminate the most noisy samples. As shown in Fig. 3, only noisy sample L is removed using this method, while noisy samples M and N will still be retained. In order to

Majority sample

Fig. 3 Disadvantages of KNN-based noise filtering.

filter noise adequately, we propose a new noise filtering method. There are three main steps: Firstly, an input dataset Q is further divided into minority sample set  $S_{min}$  and majority sample set  $S_{maj}$  according to the labels, and then the Euclidean distance between each sample in  $S_{min}$  and each sample in Q is calculated to form a distance matrix.

dis = Euclidean<sub>$$x_i \in S_{\min}, y_i \in Q(x_i, y_j)$$
 (4)</sub>

Specifically, because an sample has zero Euclidean distance from itself, we set it to a large constant in order to avoid this situation influencing the next judgments. Then, for each minority sample, we find the nearest one and three instances to it in dis using Eqs. (5) and (6), respectively.

$$index_1 = smallest_{x_i \in S_{min}}(1, dis(x_i))$$
 (5)

$$index_3 = smallest_{x_i \in S_{min}}(3, dis(x_i))$$
 (6)

Finally, two conditional judgments are made, where if index<sub>1</sub> is not a minority class, then the number of count belonging to the majority class in index<sub>3</sub> is judged in turn, and if count is greater than or equal to 2, then the minority sample is considered as noise. To further compare our denoising method with the KNN-based denoising method above, we use the two-dimensional dataset paw02a-600-5-70-BI in keel (https://sci2s.ugr.es/keel/datasets.php#sub1) to visualize the denoising results. As shown in Figs. 4 and 5 compared to MWMOTE's KNN-based denoising method, our method can more adequately remove noise even though some of the noise samples are specially distributed.

# 3.2.2 Mean shift clustering and sampling weighting

The use of average-linkage agglomerative clustering to divide the minority class samples may result in a large number of class clusters or only one or a few minority samples in a class cluster because an optimal distance threshold cannot be found, which is likely



Fig. 4 KNN-based noise filtering.



Fig. 5 Proposed denoising method.

to generate many duplicate instances in the sample synthesis step. Motivated by this, we use a parameterfree mean shift clustering algorithm to divide minority samples. This algorithm does not require setting the number of clusters beforehand and does not generate too many class clusters, thus reducing the generation of duplicate samples in the sample synthesis step. We will give an experimental comparison in Section 4. Next, we assign sampling weights to the minority class samples that satisfy the requirements according to Eqs. (7)–(9):

$$S_w(x_i) = \sum_{y_j \in S_{\text{bmaj}}} I_w(y_j, x_i)$$
(8)

$$S_p(x_i) = \frac{S_w(x_i)}{\sum\limits_{z_i \in S_{\text{imin}}} S_w(z_i)}$$
(9)

Specifically, please refer to Ref. [35] for a detailed explanation of the boundary minority class samples  $S_{\text{imin}}$ , boundary majority class samples  $S_{\text{bmaj}}$ , closeness factor  $C_f$ , and density factor  $D_f$ .

#### 3.2.3 Data cleaning

After synthesizing instances for each boundary minority class sample in the cluster they belong to by using Eq. (1) according to the selection probability  $S_p$ , we use the Tomek link method to clean the processed dataset with the aim of further removing class overlap. Tomek link is a data cleaning method defined as follows: For two different classes of samples *x* and *y*, d(x, y) denotes the distance between them, and if there is no sample *z* such that d(x, y) < d(x, z) or d(x, y) < d(y, z), then (x, y) is a Tomek link and is removed.

In summary, the full steps of the proposed method are presented in Algorithm 1. For Algorithm 1, we have two remarks.

**Remark 1.** The purpose of first judging index $_1$  in the noise denoising process is to avoid misclassifying some minority samples as noisy samples because of their particular distribution.

Remark 2. When clustering minority class samples

# Algorithm 1 SPMSC

- 1: Input
- 2: Q: dataset,  $S_{min}$ : minority class sample set,  $S_{maj}$ : majority class sample set, denoise: storing noise samples, and num: the number of samples to be generated.
- 3:  $K_1$ : nearest neighbors for finding the boundary majority class and  $K_2$ : nearest neighbors for finding the boundary minority class.
- 4: Procedure begin
- 5: Calculate the distance matrix between the sample in  $S_{\min}$  and the sample in Q using Eq. (4).
- 6: for each  $x_i \in S_{\min}$ , obtain index<sub>1</sub> and index<sub>3</sub> by using Eqs. (5) and (6).
- 7: if the index<sub>1</sub> label is not a minority class, initialize m = 0, for each index  $\in$  index<sub>3</sub>, if the label of the index is the majority class, m + 1, if  $m \ge 2$ , then add that minority class to denoise.
- 8: Denoised minority class sample set  $T_{\min} = S_{\min}$  denoise.
- 9: for each  $x_i \in T_{\min}$ , find its  $K_1$  nearest majority class samples  $N_{\max}(x_i)$  to form boundary majority sample set  $S_{\max} = \bigcup_{x_i \in T_{\min}} N_{\max}(x_i).$
- 10: for each  $y_i \in S_{\text{bmaj}}$ , find its  $K_2$  nearest minority class samples  $N_{\min}(y_j)$  to form boundary minority sample set  $S_{\min} = \bigcup_{y_j \in S_{\text{bmaj}}} N_{\min}(y_i).$
- 11: Clustering of  $S_{\min}$  using the mean shift algorithm.
- 12: for each  $x_i \in S_{\text{imin}}$  and  $y_j \in S_{\text{bmaj}}$ , sampling weights are calculated by using Eq. (7).
- 13: for each  $x_i \in S_{\text{imin}}$ , the selection probability is calculated using Eq. (9).
- 14: Initialize the set  $S = S_{\min}$ .
- 15: **do for** i = 1, 2, ..., num
- 16: Select sample x according to  $S_p$  and find the cluster<sub>x</sub>
- 17: where x is located.
- 18: Select sample y randomly in cluster<sub>x</sub>.
- 19: A synthetic sample syn is generated by using Eq. (1) and
- adds syn to  $S: S = S \cup \text{syn}$ .
- 21: end loop
- 22: Obtain new dataset new: new =  $S \cup S_{maj}$
- 23: The dataset new is cleaned using Tomek link.
- 24: **End**

using the mean shift algorithm, no parameters need to be set, where the bandwidth is estimated by the provided estimate\_bandwidth function.

#### 4 Result and Discussion

In this section, we conduct experiments and analyze the experimental results from multiple perspectives to verify the effectiveness of the SPMSC method.

#### 4.1 Dataset description and comparison methods

We conduct experiments by using 21 datasets from KEEL, UCI<sup>§</sup>, UCI\_extended<sup>¶</sup>, and RCSMOTE<sup>‡</sup>. These datasets have different sample sizes, feature attributes, and degrees of imbalance, and the details are shown in Table 1. Among them, wdbc is breast cancer dataset and its labels "M" and "B" denote malignant and benign, respectively. To be consistent with the other 20 datasets, we consider malignant as the minority sample assigned label "1" and benign as the majority sample assigned label "0".

To verify the effectiveness of the proposed method, we compare SPMSC with eight popular sampling methods, namely random oversampling

§ https://archive.ics.uci.edu/ml/index.php.

https://github.com/felix-last/evaluate-kmeans-smote/releases/download/ v0.0.1/uci\_extended.tar.gz.

<sup>☎</sup> https://raw.githubusercontent.com/M-Hashemzadeh/RCSMOTE/master/ ImplementationSourceCodes.zip. (ROS)<sup>[17]</sup>, SMOTE<sup>[32]</sup>, SMOTE-Tomeklinks (STL)<sup>[40]</sup>, ADASYN<sup>[33]</sup>, B1-SMOTE<sup>[36]</sup>, B2-SMOTE<sup>[36]</sup>, safelevel-SMOTE (SLS)<sup>[38]</sup>, and MWMOTE<sup>[35]</sup> on three classifiers, KNN<sup>[41]</sup>, GaussianNB<sup>[42]</sup>, and SVM<sup>[43]</sup>.

#### 4.2 Evaluation measures

The method for evaluating classifier performance in machine learning is based on confusion matrix. As shown in Fig. 6, where TN represents the number of negative (majority) class samples correctly classified, FP represents the number of negative (majority) class samples misclassified as positive (minority) class samples, FN represents the number of positive (minority) class samples misclassified as negative (majority) class samples, and TP represents the number of positive (minority) class samples correctly classified. The traditional evaluation method accuracy is not applicable in imbalance scenarios because it only takes into account the overall accuracy and ignores the importance of minority class samples. Therefore, some evaluation methods for imbalance scenarios<sup>[44, 45]</sup> are proposed, and the specific definitions are shown as follows:

$$F\text{-measure} = \frac{2 \times \text{Sensitivity} \times \text{Precision}}{\text{Sensitivity} + \text{Precision}}$$
(10)

$$G$$
-mean =  $\sqrt{\text{Sensitivity} \times \text{Specificity}}$  (11)

$$AUC = \frac{1 + TPR - FPR}{2} \tag{12}$$

Dataset	Number of minority samples	Number of majority samples	Number of samples	Attribute	Degree of imbalance
Yeast1	429	1055	1484	8	1:2.46
Yeast3	163	1321	1484	8	1:8.10
Yeast5	44	1440	1484	8	1:32.73
Pima	268	500	768	8	1:1.87
Glass0	70	144	214	9	1:2.06
Haberman	81	225	306	3	1:2.78
Vehicle1	217	629	846	18	1:2.90
Vehicle3	212	634	846	18	1:2.99
Glass-0-1-2-3_vs_4-5-6	51	163	214	9	1:3.20
Vehicle0	199	647	846	18	1:3.25
Ecoli1	77	259	336	7	1:3.36
Ecoli3	35	301	336	7	1:8.60
Ilpd	165	414	579	10	1:2.51
Heart	120	150	270	13	1:1.25
Liver_disorders2	72	200	272	6	1:2.78
Liver_disorders4	36	200	236	6	1:5.56
Pima2	134	500	634	8	1:3.73
Segment	330	1980	2310	16	1:6.00
Tic-tac-toe	332	626	958	9	1:1.89
Winequality-red4	53	1546	1599	11	1:29.17
Wdbc	212	357	569	31	1:1.68

Table 1Description of the imbalanced datasets.

	Predicted negative	Predicted positive
Actual negative	TN	FP
Actual positive	FN	TP

Fig. 6 Confusion matrix.

$$TPR = \frac{TP}{TP + FN} \tag{13}$$

$$FPR = \frac{FP}{FP + TN} \tag{14}$$

## 4.3 Experimental setting

The three classifiers and eight sampling algorithms are used in the experimental comparison, where, for the SVM classifier we used the hinge loss function as the loss term, C = 0.1, max\_ite = 10000, and penalty is  $L_2$ . MWMOTE uses the parameters suggested in the original paper, i.e.,  $K_1 = 5$ ,  $K_2 = 3$ ,  $K_3 = |S_{\min}|/2$ ,  $C_p = 3$ ,  $C_f$  (th) = 5, and CMAX = 2. The rest of classifier and comparison algorithm parameters used are set to default values.

#### 4.4 Result comparison and analysis

# 4.4.1 Duplicate data comparison

To visualize the duplicate points in the processed dataset, three 2-dimensional datasets are selected from KEEL. Figure 7 shows the data distribution after using the MWMOTE method. The blue point represents the majority class, the red point represents the minority class, and the point in the black rectangular box is the synthetic duplicate sample. It can be observed that the edges of these duplicate sample points show a sawtooth shape. Table 2 shows the number of duplicates of the original data in the synthesized samples using our method compared to the MWMOTE method, from which it can be concluded that our method can effectively reduce the generation of duplicate samples in the sample synthesis phase compared to the MWMOTE method.

# 4.4.2 Contrastive analysis of sampling performance

In this subsection, we compare the performance of the SPMSC method with the comparison method.

Tables 3-5 show the results for SPMSC and other eight sampling methods obtained using three classifiers on 21 datasets, with the best measures in bold. Additionally, 4-fold stratified cross validation was used to maintain the proportion of classes in the original data, and each experiment was repeated three times in order to eliminate the effect of randomness. As shown in Table 3, SPMSC obtains the best results at least on one measure in 19 out of 21 datasets when using the KNN classifier. As shown in Table 4, SPMSC obtains the best results at least on one measure in 11 out of 21 datasets when using the NB classifier. As shown in Table 5, SPMSC obtains the best results at least on one measure in 13 out of 21 datasets when using the SVM classifier. In comparison with the other eight sampling methods, the SPMSC method obtains the highest number of best results.

Figure 8 shows the average of the results of SPMSC and the other eight sampling methods. For KNN and SVM classifiers, SPMSC outperforms the other methods in all three measures. For the NB classifier, SPMSC



(a) paw02a-600-5-70-BI

(b) paw02a-800-7-60-Bl

(c) 03subcl5-600-5-30-BI-full

Fig. 7 Visualization of duplicate data generated using the MWMOTE method.

Table 2	Comparison (	of dunlicate dat	a generated hy	the SPMSC method	l and the MWM	OTE method
Table 2	Comparison	VI UUDIICAIC UAI	a generateu dv	The SI MISC memory	i anu une ivi vvivi	OIL memou.

Dataset	Number of synthetic samples	Number of duplicate samples			
Dataset	Number of synthetic samples	MWMOTE	SPMSC		
paw02a-600-5-70-BI	400	90	5		
paw02a-800-7-60-BI	700	159	14		
03subcl5-600-5-30-BI-full	400	104	11		

Table 3 Experimental results obtained on 21 datasets by using KNN classifier (*F-M* is short for *F*-measure and *G-M* is short for *G*-mean).

Dataset	Measure	ROS	SMOTE	STL	ADASYN	<b>B1-SMOTE</b>	<b>B2-SMOTE</b>	SLS	MWMOTE	SPMSC
	F-M	0.522743	0.536692	0.530 222	0.531450	0.526050	0.528 153	0.535 802	0.547 124	0.553 061
Yeast1	G-M	0.655 485	0.666418	0.661 095	0.662 405	0.657 654	0.657 407	0.665758	0.675 997	0.681 259
	AUC	0.717969	0.722232	0.720031	0.721 376	0.709 195	0.709 174	0.727 138	0.739 181	0.746 506
	F-M	0.676411	0.660401	0.681 125	0.644 022	0.681 272	0.587 982	0.654 146	0.676332	0.693 541
Yeast3	G-M	0.891676	0.883111	0.890240	0.886152	0.869750	0.863 613	0.883747	0.898 834	0.902797
	AUC	0.915 162	0.920008	0.921 669	0.914774	0.917 576	0.918 808	0.933 057	0.928 967	0.933179
-	F-M	0.617 09	0.581 992	0.586 894	0.576608	0.594 626	0.531 439	0.573 519	0.584 642	0.586 229
Yeast5	G-M	0.901115	0.920686	0.921 039	0.920406	0.921 821	0.940 573	0.955 442	0.933 357	0.933 667
	AUC	0.943 845	0.954293	0.954 104	0.954 451	0.942 487	0.960 038	0.963 699	0.965 878	0.966 162
	F-M	0.630745	0.636319	0.632 69	0.631 125	0.635 532	0.638 106	0.648711	0.639 191	0.666 279
Pima	G-M	0.709 839	0.713319	0.711135	0.706979	0.713 088	0.711956	0.724 938	0.716678	0.737 819
	AUC	0.759433	0.767970	0.771 224	0.767731	0.764 239	0.761 179	0.777 657	0.778224	0.783 507
	F-M	0.684 341	0.649 884	0.666 602	0.653 333	0.700 074	0.675 992	0.677 002	0.691 898	0.687 228
Glass0	G-M	0.760 103	0.724730	0.743 947	0.721762	0.770 171	0.739 385	0.753216	0.762917	0.758 680
	AUC	0.854666	0.842 320	0.843 387	0.848 051	0.859 307	0.843 273	0.843 035	0.842218	0.851 841
	F-M	0.421019	0.44088	0.357 196	0.405 857	0.374 330	0.407 483	0.387785	0.459 331	0.440 081
Haberman	G-M	0.581717	0.600795	0.525 155	0.568 410	0.542283	0.569476	0.553118	0.615 404	0.597 123
	AUC	0.629310	0.633 590	0.605 722	0.623 992	0.616953	0.604 969	0.635 230	0.639 898	0.657 138
	F-M	0.607.019	0.640211	0.618 508	0.621 596	0.613285	0.611 922	0.636 870	0.623796	0.641 375
Vehicle1	G-M	0.662310	0.670 863	0.667 071	0.665134	0.654 432	0.658 299	0.675 294	0.658 596	0.690 055
	AUC	0.787 609	0.795734	0.788 960	0.788 365	0.785 827	0.788 863	0.783 594	0.791720	0.794212
	F-M	0.605 194	0.637 168	0.623961	0.620 988	0.617 169	0.617.063	0.614 278	0.623436	0.651 422
Vehicle3	G-M	0.687 533	0.710.690	0.692.633	0.677 971	0.694 288	0.695 193	0.693.086	0.680.048	0.712 928
venicies	AUC	0.797 793	0.805 608	0.806233	0.812.216	0.807 651	0.797 736	0.810702	0.792.435	0.812 696
	F-M	0 747 578	0 712 082	0 748 193	0 764 255	0 749 501	0.715.580	0 741 925	0731710	0.738.032
Glass-0-1-2-3 vs 4-5-6	G-M	0.823.241	0.789.808	0.818.214	0.837.035	0.823 587	0.814.925	0.821.308	0.811.880	0.814.975
01035-0-1-2-5_v3_4-5-0	AUC	0.874.293	0.901 798	0.881.623	0.914.888	0.883417	0.011723	0.874.070	0.880.470	0.916.891
		0.815 186	0.901790	0.841 176	0.911000	0.819.156	0.782.420	0.825 707	0.823.652	0.853790
Vehicle()	$G_{-M}$	0.010 100	0.023700	0.041170	0.031 920	0.019150	0.782 420	0.023707	0.023032	0.033770
venicieo		0.067.80/	0.974.264	0.923 140	0.920393	0.970.341	0.057.626	0.074.061	0.921975	0.942 015
	$\frac{AUC}{F_{-}M}$	0.725 720	0.777.888	$\frac{0.973171}{0.711010}$	0.774.080	0.730.820	0.741 560	0.717.686	0.748132	0.752 647
Egoli1	C M	0.723720	0.727 888	0.711.717	0.823 722	0.750.829	0.820.512	0.717000	0.824.080	0.831 520
Ecolifi		0.810 407	0.875 104	0.862 338	0.823722	0.873.270	0.868 328	0.820 320	0.827 312	0.880 111
		0.049427	0.075194	0.802 558	0.583.611	0.679 270	0.603 328	0.039107	0.582168	0.624.126
Ecoli3	$G_{-M}$	0.240.313	0.394 937	0.399079	0.365.011	0.870.285	0.017 845	0.349430	0.382108	0.024 120
Ecolis		0.021250	0.009171	0.012 880	0.800.037	0.013.061	0.072338	0.004 / 00	0.004 661	0.007 040
		0.094700	0.903730	0.912 889	0.097 443	0.913 901	0.907 403	0.914 497	0.904 001	0.905 155
Und	$\Gamma - M$	0.448 / /0	0.447750	0.433270	0.439 848	0.417290	0.431 180	0.405 025	0.433 049	0.404 337
npa		0.575177	0.501 525	0.574 445	0.579 525	0.546 505	0.571 092	0.362 367	0.579452	0.505 000
	F M	0.034 329	0.020 803	0.055 555	0.020 397	0.020137	0.027 383	0.020710	0.022.023	0.030 209
Haant	$\Gamma - M$	0.797113	0.019.000	0.013209	0.790.303	0.829203	0.809 292	0.024 390	0.020400	0.032421
пеан		0.810010	0.034 391	0.033414	0.010/90	0.841 005	0.820 900	0.041 0.00	0.843 383	0.878.024
		0.007 020	0.072.004	0.070701	0.302.011	0.809 905	0.303 040	0.210.201	0.882 278	0.070 924
Liver disorders?	G M	0.290402	0.278 902	0.330.081	0.313 037	0.294 030	0.314720	0.319 391	0.314 014	0.550 204
Liver_disorders2		0.441491	0.457 244	0.404 934	0.403 420	0.449433	0.400712	0.407 938	0.438493	0.504.004
		0.409 444	0.407 005	0.339.028	0.343472	0.472778	0.432 301	0.320 389	0.490 972	0.326 194
Liven disenders4	C M	0.292 010	0.557410	0.293 031	0.299 300	0.201 300	0.247 324	0.272 800	0.502 /81	0.542 519
Liver_disorders4		0.505755	0.624.444	0.500.505	0.577422	0.539.520	0.510.599	0.559 049	0.573478	0.619 222
	F M	0.004 107	0.024 444	0.559444	0.021 007	0.010278	0.332778	0.360 276	0.393 333	0.020 333
D:2	$\Gamma - M$	0.44/001	0.475739	0.471 140	0.400 204	0.470.020	0.430 403	0.403 203	0.495 545	0.304 809
Pima2		0.030070	0.070277	0.072732	0.0/11/0	0.009 098	0.039 291	0.005 252	0.094 374	0.701012
	EM	0.702.007	0.719742	0.717297	0.714 878	0.734 442	0.732310	0.755 552	0.740740	0.746 090
Comment	г-М С М	0.055 407	0.054.072	0.055 126	0.009 283	0.070928	0.7890/9	0.001 333	0.056.044	0.057501
Segment		0.93309/	0.9340/3	0.933 130	0.070 705	0.931423	0.930 239	0.949 39/	0.930 944	0.93/301
	AUC E M	0.970831	0.9/802/	0.911032	0.918 /83	0.9//318	0.978783	0.9/9043	0.979799	0.203 243
The A	г-М С М	0.02/109	0.000 931	0.038492	0.003 388	0.013.011	0.382030	0.040 /01	0.00/021	0.080490
11c-tac-toe		0.090.342	0.769 570	0.764.450	0.750.429	0.0737570	0.031/2/	0.707028	0.737490	0.128 908
	AUC E M	0.131/09	0.100319	0.104430	0.139428	0.134312	0.701204	0.130.909	0.126.520	0.190982
Wincoustites and 4	г-М С М	0.139 100	0.1303/0	0.13/932	0.141 328	0.148 952	0.1/0018	0.124 273	0.130 338	0.13/043
winequanty-red-4		0.483 383	0.39/033	0.39/314	0.39998/	0.441 000	0.30/331	0.333 908	0.003 333	0.005 299
	AUC E M	0.39/243	0.030400	0.038203	0.04/844	0.013 032	0.020332	0.0433/1	0.032011	0.073 094
<b>W7_11_</b> -	r-M CM	0.930.043	0.949 103	0.950.982	0.923 382	0.930 092	0.900051	0.9334/1	0.949 109	0.950 015
wabc		0.900 807	0.901 342	0.901/4/	0.944 23/	0.932027	0.928013	0.903 038	0.9012//	0.000 352
	AUC	0.770/0/	U.771 /77	0.770.021	0.7041/9	0.983 202	0.713 830	0.770 380	0.989 200	0.990 290

	Table 4	Experin	nental resu	ılts obtain	ed on 21 d	latasets by ı	using NB cla	ssifier.		
Dataset	Measure	ROS	SMOTE	STL	ADASYN	B1-SMOTE	E B2-SMOTE	SLS	MWMOTE	SPMSC
	F-M	0.456 125	0.458 645	0.457 923	0.458730	0.457 897	0.455 935	0.457 501	0.460728	0.462 404
Yeast1	G-M	0.196 169	0.233 243	0.225 275	0.227735	0.227 596	0.209 021	0.216 401	0.243 092	0.261 612
	AUC	0.516403	0.522 134	0.520714	0.521 882	0.520712	0.516920	0.519 509	0.525 670	0.529 242
	F-M	0.219601	0.233 448	0.236 745	0.218 333	0.218 486	0.206 591	0.227 033	0.231 906	0.228 198
Yeast3	G-M	0.346482	0.437 491	0.452 808	0.338814	0.354 067	0.237 156	0.399 952	0.427 586	0.412470
	AUC	0.560114	0.591 879	0.598 693	0.557084	0.558 186	0.526 028	0.577 891	0.588 095	0.580 502
	F-M	0.114764	0.169 266	0.175 865	0.164 341	0.185 158	0.124 036	0.141 530	0.174 980	0.178614
Yeast5	G-M	0.725 108	0.822667	0.831 483	0.815 966	0.841 108	0.709 541	0.789 541	0.830 068	0.833 602
	AUC	0.763 194	0.832 828	0.840 120	0.826 926	0.851136	0.752 525	0.807734	0.838731	0.841 509
	F-M	0.645 945	0.646 374	0.637 224	0.640 895	0.632705	0.634 009	0.646 661	0.655 554	0.647 528
Pima	G-M	0.721 980	0.723 317	0.715 240	0.718 069	0.711247	0.712415	0.723 570	0.731 123	0.724 135
	AUC	0.725 552	0.726.687	$\frac{0.111552}{0.00575}$	0.720149	0./13418	0./14/284	0./26.68/	0.734 284	0.726418
C10	F-M C M	0.648 094	0.652.034	0.6493/3	0.626.997	0.612366	0.60/52/	0.648 094	0.622.321	0.651941
Glasso		0.031987	0.038 323	0.032 578	0.034 393	0.019109	0.000 333	0.031987	0.03/9//	0.004 927
	E M	0.703 882	0.709 555	0.703802 0.421200	0.067 290	0.072 998	0.000.034	0.703 882	0.000.007	0./12410
Habarman	G M	0.597 960	0.409.007	0.451 200	0.595 054	0.400.893	0.423 304	0.410 009	0.414 0.51	0.440 270
Haberman		0.558 918	0.613.518	0.508245 0.623410	0.549.944	0.608.647	0.616.267	0.549.591	0.603 791	0.571 050
	$\frac{F-M}{F}$	0 548 641	0 549 270	$\frac{0.023 + 10}{0.549 440}$	0 544 596	0.520.670	0.516.226	0.549 500	0.547 146	0 552 504
Vehicle1	G-M	0.658 387	0.655.214	0.659.961	0.622.060	0.624 599	0 593 398	0.659.831	0.652.197	0.664 494
veniere r	AUC	0.680 977	0.678 799	0.680387	0.675 831	0.662731	0.656 929	0.681 183	0.678 045	0.683 493
	F-M	0.538.009	0.540.603	0.543 196	0.541 562	0.556 529	0.546 873	0.546298	0.537.867	0.539489
Vehicle3	G-M	0.660 356	0.662 850	0.669 427	0.657 586	0.660412	0.652 023	0.670 367	0.663 493	0.664351
	AUC	0.671 981	0.675 110	0.679 031	0.680 509	0.690 769	0.684 460	0.680 609	0.674314	0.675110
	F-M	0.704 351	0.697 455	0.708 851	0.731 197	0.712 044	0.757 099	0.715747	0.697 455	0.714076
Glass-0-1-2-3_vs_4-5-6	G-M	0.786217	0.783 029	0.793611	0.810372	0.800491	0.839 872	0.796 798	0.783 029	0.801 492
	AUC	0.813759	0.810634	0.821 050	0.829864	0.828418	0.854 216	0.824 175	0.810634	0.827617
	F-M	0.550449	0.556710	0.555 883	0.495 115	0.491 435	0.488 522	0.551 346	0.555 603	0.559914
Vehicle0	G-M	0.719673	0.726740	0.725 349	0.672182	0.670231	0.666 693	0.720795	0.725 440	0.729619
	AUC	0.738 867	0.745 127	0.744 286	0.677 976	0.674 297	0.670 531	0.739 582	0.743 334	0.747 437
	F-M	0.500 329	0.560 281	0.592 680	0.503 932	0.644 887	0.512 575	0.538 384	0.533 907	0.425 556
Ecoli1	G-M	0.630874	0.711 160	0.740 656	0.608 356	0.772 606	0.612 207	0.685 275	0.674 823	0.417 278
	AUC	0.691 833	0.744 060	0.766 958	0.683 812	0.800 580	0.697 389	0.725 070	0.721943	0.594 925
<b>T</b> 110	F-M	0.436 652	0.505 780	0.538 696	0.474 945	0.575 198	0.519686	0.513131	0.552 585	0.534 887
Ecoli3	G-M	0.778663	0.846 429	0.868974	0.820 141	0.886 917	0.865 521	0.836 960	0.876219	0.852733
	AUC	0.799525	0.852 683	0.8/2039	0.831038	0.889 262	0.869 306	0.843 947	0.879284	0.85/105
Ilmd	F-M CM	0.550 /90	0.624 785	0.500.052	0.500.207	0.505 104	0.555209	0.558 290	0.500.085	0.304 203
npa		0.670356	0.034 / 83	0.031428	0.623 440	0.633.230	0.015727	0.619012	0.022717	0.688.030
	$\frac{AUC}{F_{-}M}$	$\frac{0.079}{0.820}$	0.811.612	0.090 220	0.004104	0.000 933	0.820.968	0.001 740	0.00+173	0.000 950
Heart	G-M	0.820140	0.830 104	0.820.031	0.819730	0.827 334	0.820 908	0.834707	0.815 035	0.842.185
ficart	AUC	0.838 276	0.830731	0.838 276	0.836.652	0.837 530	0.837 530	0.834 987	0.835 864	0.842 443
	F-M	0.409 303	0.404 563	0.424 769	0.415 193	0.401 602	0.400 243	0.426 514	0.397 700	0.413.093
Liver disorders2	G-M	0.427 289	0.453 305	0.489 620	0.450 340	0.444 083	0.449 047	0.458 026	0.449 125	0.469 329
	AUC	0.520278	0.519444	0.548 333	0.530278	0.514 444	0.515 000	0.544 167	0.512 500	0.531 389
	F-M	0.240762	0.278 479	0.260 661	0.264 386	0.273 366	0.264 478	0.244 441	0.269 838	0.283 549
Liver_disorders4	G-M	0.407 270	0.484 687	0.470 993	0.473 079	0.523 571	0.497 473	0.442 441	0.487 275	0.519918
	AUC	0.477222	0.535000	0.512 222	0.512222	0.536944	0.520 833	0.485 833	0.526111	0.548 611
	F-M	0.547 570	0.550409	0.556 164	0.539910	0.550126	0.541 306	0.555 083	0.551 332	0.563 640
Pima2	G-M	0.729 104	0.732 563	0.734 107	0.730484	0.734 527	0.733 970	0.733 161	0.732 823	0.742 681
	AUC	0.732 461	0.735 360	0.737 684	0.732724	0.736 824	0.735 188	0.736 572	0.735 360	0.745 936
	F-M	0.481 369	0.482 577	0.482 894	0.472 092	0.436 648	0.435 348	0.479 383	0.480721	0.480 994
Segment	G-M	0.794 299	0.795 966	0.795 843	0.783 563	0.742 439	0.744 212	0.793 147	0.791 630	0.791 519
	AUC	0.805 256	0.807 510	0.807014	0.792216	0.744712	0.747 931	0.804 760	0.800 494	0.799 998
<b>T</b> ' · · ·	F-M	0.437 575	0.435 789	0.431762	0.434 024	0.432149	0.450 205	0.430968	0.420 254	0.445 793
Tic-tac-toe	G-M	0.416158	0.403 033	0.408 068	0.4/2304	0.51/304	0.500.811	0.4/0358	0.48/6/3	0.520611
	AUC E M	0.498 893	0.320 /42	0.321133	0.323 /20	0.329 920	0.329 303	0.330 232	0.3290/0	0.115.049
Winaguality and 4	г-М С М	0.627 724	0.104 330	0.104413	0.102.08/	0.121130	0.113 323	0.1023/3	0.100 328	0.113 048
winequality-red-4		0.027730	0.621 100	0.021 322	0.020203	0.505 097	0.007 344	0.01/203	0.027008 0.637041	0.043 30/
	F-M	0 902 657	0 907 750	0.002.042	0.029782	0.029070	0.000.000	0.029.038	0.037.041	0 911 947
Wdbc	G-M	0.919784	0.924 505	0.921 029	0.936 901	0.945 502	0.942.120	0.919784	0.927 363	0.927 287
	AUC	0.920 599	0.925 316	0.922 003	0.937 161	0.945 625	0.942 382	0.920 599	0.928 109	0.928 125

	Table 5	Experim	ental resul	lts obtaine	ed on 21 da	atasets by us	ing SVM cla	assifier.		
Dataset	Measure	ROS	SMOTE	STL	ADASYN	<b>B1-SMOTE</b>	<b>B2-SMOTE</b>	SLS	MWMOTE	SPMSC
	F-M	0.592 368	0.592131	0.581 181	0.588 900	0.584 872	0.576488	0.593 156	0.590 803	0.596 054
Yeast1	G-M	0.713 346	0.713843	0.705 003	0.709820	0.701 247	0.692685	0.714714	0.712832	0.717 396
	AUC	0.714 461	0.715 402	0.705 799	0.713 628	0.710064	0.701 789	0.716 163	0.714 406	0.718 483
	F-M	0.671 487	0.661 126	0.671 183	0.614 135	0.626 888	0.575 565	0.665 148	0.668 794	0.677 518
Yeast3	G-M	0.906 631	0.890 026	0.899 324	0.905 097	0.900 683	0.893 496	0.897 850	0.901 446	0.898 952
	AUC	0.907 212	0.890 506	0.899 807	0.906 600	0.901 438	0.896 096	0.898 138	0.901 794	0.899 401
	F-M	0.487 323	0.495 966	0.493 059	0.492 150	0.501 350	0.433 877	0.467 972	0.498 103	0.503 425
Yeast5	G-M	0.967164	0.968 239	0.967881	0.967 885	0.968 957	0.957730	0.964 281	0.968 600	0.969 319
	AUC	0.967708	0.968 / 50	0.968403	0.968 403	0.969 444	0.958 681	0.964 931	0.969.097	0.969 /92
D'	F-M C M	0.0/0214	0.001 902	0.003 338	0.034 307	0.0/1824	0.058 444	0.003 327	0.072284	0.0/1155
Pillia		0.740410	0.730703	0.730.510	0.731.075	0.744.003	0.732 103	0.730494	0.745400	0.744 304
	F M	0.749 940	0.758.012	0.739478	0.751075	0.744 800	0.753 134	0.739478	0.740403	0.745072
Glass	$G_{-M}$	0.636.695	0.649333	0.648 893	0.617.743	0.615 245	0.622.282	0.647 167	0.649333	0.000137
Glasso	AUC	0.030.093	0.049.555	0.040.095	0.711.397	0.707.925	0.715.278	0.718342	0.721.814	0.732.230
	F-M	0 441 240	0433730	0 430 352	0.416.802	0.458.551	0 459 494	0 435 317	0.457.352	0.468 581
Haberman	G-M	0.579 542	0.573780	0.572 188	0.559 309	0.599428	0.596414	0.574 643	0.597 680	0.610 503
muoonnun	AUC	0.626 982	0.623 042	0.620 849	0.614 482	0.636 842	0.639 482	0.624 530	0.636 803	0.642 685
	F-M	0.692 512	0.696358	0.696 701	0.688 481	0.707 012	0.705 141	0.698 989	0.696 667	0.695 970
Vehicle1	G-M	0.745 186	0.745 669	0.760113	0.753 700	0.773 932	0.779 834	0.746416	0.752276	0.754 094
	AUC	0.777 648	0.782176	0.786 267	0.781 495	0.801 638	0.805 376	0.779 998	0.783 062	0.784 554
	F-M	0.645 242	0.645 182	0.656 978	0.651763	0.661 518	0.681 340	0.669 321	0.645 753	0.655 155
Vehicle3	G-M	0.713410	0.719744	0.723 238	0.716800	0.720371	0.729 039	0.733 066	0.712 103	0.727605
	AUC	0.751473	0.756941	0.762 479	0.760101	0.760917	0.775 859	0.770351	0.753 811	0.764818
	F-M	0.739 797	0.781917	0.782 399	0.744 398	0.775 010	0.821 882	0.768 846	0.743 954	0.799 921
Glass-0-1-2-3_vs_4-5-6	G-M	0.821 315	0.851 598	0.851 892	0.830478	0.848 660	0.896 149	0.839 201	0.823 035	0.863 703
	AUC	0.834 909	0.860313	0.860313	0.841 551	0.857 264	0.899 167	0.850698	0.838 034	0.873 053
	F-M	0.907 931	0.911645	0.911645	0.903 730	0.908 021	0.848 079	0.894 209	0.911437	0.914718
Vehicle0	G-M	0.963 438	0.963 495	0.963 495	0.962 004	0.963 574	0.941 985	0.955 549	0.963 497	0.967 576
	AUC	0.963 801	0.963 671	0.963 671	0.962 308	0.963 852	0.943 401	0.955 765	0.963 671	0.967 900
	F-M	0.771 700	0.747 271	0.755 952	0.761 996	0.790773	0.784675	0.765 418	0.772630	0.769 159
Ecoli1	G-M	0.839 339	0.822935	0.832 183	0.842 877	0.869705	0.864 350	0.835 094	0.840279	0.841 206
	AUC	0.853680	0.837304	0.845 477	0.857 193	0.879936	0.872243	0.849 353	0.854338	0.854 308
E 1'2	F-M C M	0.561 027	0.603 521	0.599.024	0.5536/4	0.605 588	0.589680	0.5860/9	0.614 486	0.010453
Ecol13	G-M	0.8/3043	0.889.055	0.889 100	0.881 495	0.888 942	0.894011	0.883 088	0.892.399	0.805.262
	AUC E M	0.877018	0.890 203	0.890 283	0.884 202	0.890 283	0.552.462	0.0009/4	0.893 018	0.893 203
IInd	G M	0.502 500	0.500 082	0.570.520	0.571459	0.536 329	0.555402	0.572508	0.502 151	0.574120
npa		0.039392	0.047703	0.037 334 0 700 966	0.047780	0.634284	0.627 249	0.049217	0.635 958	0.699.845
	$\frac{F-M}{F}$	0.818.424	0.811 556	0.813.251	0.090.090	0.808 144	0.827 710	0.803.223	0.808 167	0.821 585
Heart	G-M	0.836 829	0.830491	0.831 536	0.819464	0.825719	0.844 010	0.822.917	0.827 216	0.838 914
licuit	AUC	0.837 577	0.830 909	0.831 697	0.819 986	0.825 818	0.844 109	0.823 364	0.827 531	0.839.065
	F-M	0.450 104	0.446 553	0.431 227	0.421 169	0.449 695	0.398 329	0.404 643	0.430 548	0.467 003
Liver_disorders2	G-M	0.602 826	0.589 149	0.551726	0.560439	0.593 400	0.531 377	0.533 357	0.569 691	0.594 609
	AUC	0.607 500	0.595 833	0.572 222	0.569 444	0.598 333	0.545000	0.549 444	0.576944	0.611 111
	F-M	0.368 846	0.357 452	0.381 826	0.378776	0.333 571	0.242 191	0.299 029	0.351667	0.387 227
Liver_disorders4	G-M	0.615953	0.610874	0.646 806	0.639 850	0.566946	0.465 268	0.521 900	0.617 224	0.650 434
	AUC	0.634722	0.627222	0.656111	0.658889	0.598 056	0.501 389	0.559 444	0.629722	0.670 278
	F-M	0.550264	0.518469	0.549 494	0.523 274	0.527 983	0.535752	0.557 247	0.542 289	0.548 858
Pima2	G-M	0.736702	0.708 609	0.736 243	0.722478	0.719777	0.730171	0.738 545	0.732 339	0.736923
	AUC	0.738 389	0.712784	0.738 289	0.723 642	0.721 814	0.731642	0.741 490	0.734 289	0.738 177
	$F-\overline{M}$	0.641 972	0.642 479	0.641 955	0.631 502	0.632 087	0.612 918	0.642 782	0.646 316	0.647 629
Segment	G-M	0.899 813	0.898 413	0.898 123	0.895 871	0.895 336	0.886 025	0.899 237	0.900 075	0.900 622
	AUC	0.904 031	0.902 263	0.902 011	0.900739	0.899 991	0.891 657	0.903 283	0.903 778	0.904 283
<b>T</b> .	F-M	0.447 417	0.450 643	0.441 412	0.451 107	0.450734	0.441 919	0.467 160	0.454 993	0.462 661
Tic-tac-toe	G-M	0.375374	0.416403	0.388171	0.386118	0.484 603	0.469514	0.466 263	0.406388	0.388 969
	AUC	0.491412	0.124.062	0.480 441	0.122.280	0.013803	0.496 946	0.522 584	0.499.097	0.50/446
Winaguelity and 4	г-М С М	0.139 223	0.134903	0.141 801	0.132 388	0.204 039	0.184 /80	0.205 321	0.100 59/	0.1392/3
winequality-red-4		0.703 /1/	0.00/038	0.090.039	0.003.211	0.041 303	0.032034	0.711038	0.702 043	0.717.002
	$F_{-M}$	0.710033	0.055.010	0.705 105	0.072 7.54	0.004 233	0.003974	0.06/ 207	0.700 100	0.717992
Wdbc	$G_{-M}$	0.202.340	0 974 925	0 971 117	0.961.968	0.963.518	0.092277	0.970.061	0.974.870	0.978 706
TT UUC	AUC	0.970 881	0.975.095	0.971 332	0.962.454	0.963 859	0.921 833	0.970362	0.975 079	0.978 842

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Fig. 8 Average results of nine sampling methods on 21 datasets achieved by (a) KNN classifier, (b) NB classifier, and (c) SVM classifier.

is only lower than the STL method in the *F*-measure metric, lower than the SMOTE, STL, and MWMOTE methods in the *G*-mean metric, and lower than the SMOTE and STL methods in the *AUC* metric, but still ranked in the top from an overall perspective.

To more intuitively compare the performance of SPMSC with the other eight sampling methods, the average rank of each method on 21 datasets is calculated. As shown in Fig. 9, the lower average rank values represent a higher rank, and from Fig. 9, we can get that SPMSC method ranks the highest on three classifiers.



(c) SVM

Fig. 9 Average rank results of nine sampling methods on 21 datasets achieved by (a) KNN classifier, (b) NB classifier, and (c) SVM classifier.

# 4.4.3 Contrastive analysis of performance improvement

Further, to compare the classification performance improvement of the data processed using the sampling method compared to the original data, we select the AUC metric from the results obtained on KNN, NB, and SVM classifiers for comparison. Table 6 shows the AUC results obtained by directly classifying 21 datasets with the three classifiers. Figure 10 describes the difference between the AUC values of each dataset in Tables 3-5 and the AUC values of each dataset in Table 6. As shown in Fig. 10, a positive value indicates that the data processed by the sampling method on the same classifier have better classification performance than the original data, and a negative value indicates that the data processed by the sampling method on the same classifier have lower classification performance than the original data. It is not difficult to find that the SPMSC method obtains the highest number of maximum differences, which indicates that the SPMSC outperforms the other eight comparison sampling methods. It is also observed that when KNN and NB classifiers are used, the AUC differences between the data processed using sampling methods and the original data are positive and negative. Yet when SVM

Table 6AUC results obtained for the original dataset onKNN, NB, and SVM classifiers.

Detect	AUC result							
Dataset	KNN classifer	NB classifer	SVM classifer					
Yeast1	0.735 060	0.517 142	0.594 049					
Yeast3	0.922 960	0.583 555	0.809 665					
Yeast5	0.947 159	0.804 167	0.500 000					
Pima	0.767 299	0.709 642	0.713 388					
Glass0	0.829 668	0.705 882	0.601 307					
Haberman	0.630 394	0.566 447	0.504 762					
Vehicle1	0.787 479	0.687 085	0.575 065					
Vehicle3	0.811 017	0.679 902	0.516 509					
Glass-0-1-2-3_vs_4-5-6	0.876 073	0.824 175	0.834 199					
Vehicle0	0.981 349	0.723 781	0.948 378					
Ecoli1	0.863 790	0.725 909	0.808 614					
Ecoli3	0.921 444	0.774 415	0.498 333					
Ilpd	0.626 836	0.680 558	0.500 000					
Heart	0.877 359	0.845 733	0.836 000					
Liver_disorders2	0.521 250	0.508 056	0.500 000					
Liver_disorders4	0.585 833	0.511 389	0.500 000					
Pima2	0.730 111	0.686 971	0.611 125					
Segment	0.979 645	0.804 507	0.743 940					
Tic-tac-toe	0.717 654	0.525 865	0.500 000					
Winequality-red-4	0.600 756	0.527 612	0.500 000					
Wdbc	0.991 241	0.920 599	0.970 813					

classifier is used, almost all the differences are positive. It shows that the performance of the same sampling method varies for different classifiers.

#### 4.4.4 Wilcoxon signed rank test

In this subsection, from the perspective of statistical analysis, we use a nonparametric test called the Wilcoxon signed rank test<sup>[46]</sup> to verify the statistical significance of the proposed method with the other eight sampling methods. The results are shown in Table 7. When using the KNN classifier, the *p*-values of all three measures are below the significance level  $\alpha = 0.05$ . Therefore, all null hypotheses are rejected, which indicates a significant improvement of SPMSC compared to the other eight sampling methods. When using the NB classifier, the null hypothesis cannot be rejected as follows: STL and B1-SMOTE under Fmeasure, STL under G-mean, and STL, B1-SMOTE, and MWMOTE under AUC, indicating that SPMSC does not have a significant improvement in comparison with these methods. Except for these cases, all null hypotheses are rejected, indicating that SPMSC has a significant improvement in comparison with other methods. When using the SVM classifier, SPMSC has significant improvement compared to other methods except for B2-SMOTE under G-mean measure.

# 4.4.5 Running time comparison

In this subsection, from the perspective of running time, we compare the time cost of the SPMSC method with the comparison method, and the results are shown in Table 8. From the results, it can be found that random oversampling has the shortest running time and SMOTE ranks the second due to their simple implementation mechanism. Compared to the original method SMOTE, some SMOTE variants such as ADASYN, B1-SMOTE, B2-SMOTE, etc., require some additional time for weighting the samples and deciding the boundary samples. Specifically, the SPMSC method requires some additional running time compared to the comparison method due to the computation of the distance matrix and the discrimination of the boundary minority samples. But, this extra computing time of a few or tens of seconds is acceptable, especially for offline computing.

In summary, the comparison and analysis of the experimental results can be concluded that the SPMSC method outperforms the comparison method on most datasets. Furthermore, a statistical analysis method called the Wilcoxon sign rank test is used to further demonstrate a significant difference between the SPMSC method and the comparison method.



Fig. 10 Performance difference between the dataset processed using the sampling method and the original dataset on the three classifiers.

# 5 Conclusion

In this paper, we propose a joint sample position based noise filtering and mean shift clustering (SPMSC) method to deal with imbalanced data. The advantages of SPMSC are that it can adequately filter noisy samples by utilizing information about the position and distribution of minority samples relative to the majority; it uses a mean shift algorithm to cluster minority samples to prevent duplicate data from being generated at the sample synthesis stage due to the creation of inappropriate class clusters; and it uses a data cleaning method to further eliminate class overlap in the processed dataset. For evaluating the proposed method, 21 datasets with different imbalance ratios and eight popular sampling algorithms are used, and the experimental results show the effectiveness of SPMSC.

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Classifor		F-measure			G-mean		AUC			
Algorithr		<i>p</i> -value	$\alpha = 0.05$	Algorithm	<i>p</i> -value	$\alpha = 0.05$	Algorithm	<i>p</i> -value	$\alpha = 0.05$	
	ROS	0.001 021	Rejected	ROS	0.000 123	Rejected	ROS	0.000 106	Rejected	
	SMOTE	0.000214	Rejected	SMOTE	0.000 141	Rejected	SMOTE	0.000 123	Rejected	
	STL	0.000 419	Rejected	STL	0.000080	Rejected	STL	0.000702	Rejected	
KNN	ADASYN	0.000 281	Rejected	ADASYN	0.000 321	Rejected	ADASYN	0.000 321	Rejected	
	B1-SMOTE	0.002 642	Rejected	B1-SMOTE	0.000 162	Rejected	B1-SMOTE	0.000 321	Rejected	
	<b>B2-SMOTE</b>	0.000 281	Rejected	<b>B2-SMOTE</b>	0.000080	Rejected	<b>B2-SMOTE</b>	0.000 069	Rejected	
	SLS	0.000 106	Rejected	SLS	0.000702	Rejected	SLS	0.001 155	Rejected	
	MWMOTE	0.007066	Rejected	MWMOTE	0.004 615	Rejected	MWMOTE	0.000 321	Rejected	
	ROS	0.006 363	Rejected	ROS	0.001 304	Rejected	ROS	0.002 098	Rejected	
	SMOTE	0.017270	Rejected	SMOTE	0.011738	Rejected	SMOTE	0.030 365	Rejected	
	STL	0.098 741	Not rejected	STL	0.169775	Not rejected	STL	0.305 198	Not rejected	
ND	ADASYN	0.012 949	Rejected	ADASYN	0.003 705	Rejected	ADASYN	0.006 363	Rejected	
IND	<b>B1-SMOTE</b>	0.180 841	Not rejected	B1-SMOTE	0.035 480	Rejected	B1-SMOTE	0.091 845	Not rejected	
	<b>B2-SMOTE</b>	0.027 306	Rejected	<b>B2-SMOTE</b>	0.014 269	Rejected	<b>B2-SMOTE</b>	0.029 829	Rejected	
	SLS	0.020 812	Rejected	SLS	0.002356	Rejected	SLS	0.017270	Rejected	
	MWMOTE	0.038 632	Rejected	MWMOTE	0.049 552	Rejected	MWMOTE	0.058 186	Not rejected	
	ROS	0.000 214	Rejected	ROS	0.002356	Rejected	ROS	0.001 021	Rejected	
	SMOTE	0.000 069	Rejected	SMOTE	0.000702	Rejected	SMOTE	0.000 060	Rejected	
	STL	0.000 245	Rejected	STL	0.000 367	Rejected	STL	0.000 281	Rejected	
SVM	ADASYN	0.000 060	Rejected	ADASYN	0.000 245	Rejected	ADASYN	0.000 321	Rejected	
<b>3</b> V IVI	B1-SMOTE	0.015 707	Rejected	B1-SMOTE	0.042020	Rejected	B1-SMOTE	0.017270	Rejected	
	<b>B2-SMOTE</b>	0.017270	Rejected	<b>B2-SMOTE</b>	0.068 035	Not rejected	<b>B2-SMOTE</b>	0.038 632	Rejected	
	SLS	0.017 270	Rejected	SLS	0.002961	Rejected	SLS	0.011738	Rejected	
	MWMOTE	0.000214	Rejected	MWMOTE	0.002 356	Rejected	MWMOTE	0.000 321	Rejected	

 Table 7
 Wilcoxon signed rank test between the SPMSC method and the comparison method.

 Table 8
 Runtime of the SPMSC method and the comparison method.

Detect					Runtime (	s)			
Dataset	ROS	SMOTE	STL	ADASYN	B1-SMOTE	<b>B2-SMOTE</b>	SLS	MWMOTE	SPMSC
Yeast1	0.0008	0.0041	0.0309	0.0135	0.0206	0.0226	0.0405	23.1532	11.7985
Yeast3	0.0009	0.0035	0.0368	0.0075	0.0102	0.0117	0.0575	3.6535	3.3682
Yeast5	0.0009	0.0034	0.0328	0.0051	0.0059	0.0069	0.0656	1.4152	1.6774
Pima	0.0006	0.0025	0.0088	0.0071	0.0120	0.0132	0.0139	8.0749	4.1992
Glass0	0.0005	0.0012	0.0028	0.0023	0.0038	0.0043	0.0040	0.5362	0.3733
Haberman	0.0005	0.0014	0.0028	0.0025	0.0040	0.0040	0.0054	0.8721	0.6220
Vehicle1	0.0008	0.0042	0.0411	0.0126	0.0168	0.0181	0.0473	6.5939	4.3879
Vehicle3	0.0008	0.0043	0.0400	0.0125	0.0168	0.0175	0.0496	5.9462	3.9708
Glass-0-1-2-3_vs_4-5-6	0.0005	0.0012	0.0028	0.0021	0.0078	0.0080	0.0091	0.3630	0.3203
Vehicle0	0.0008	0.0043	0.0427	0.0122	0.0160	0.0159	0.0489	4.8383	3.6301
Ecoli1	0.0005	0.0014	0.0043	0.0028	0.0043	0.0050	0.0080	0.7334	0.5603
Ecoli3	0.0006	0.0015	0.0049	0.0023	0.0030	0.0031	0.0107	0.3465	0.3728
Ilpd	0.0006	0.0020	0.0073	0.0050	0.0080	0.0088	0.0139	3.6298	2.2631
Heart	0.0005	0.0014	0.0035	0.0041	0.0056	0.0059	0.0035	1.4638	0.8093
Liver_disorders2	0.0006	0.0015	0.0034	0.0026	0.0038	0.0039	0.0062	0.7293	0.4516
Liver_disorders4	0.0006	0.0014	0.0030	0.0020	0.0025	0.0027	0.0070	0.3012	0.2630
Pima2	0.0007	0.0021	0.0076	0.0044	0.0063	0.0075	0.0163	2.0576	1.5813
Segment	0.0013	0.0093	0.3011	0.0372	0.0425	0.0430	0.1790	13.4986	11.6032
Tic-tac-toe	0.0007	0.0030	0.0144	0.0090	0.0133	0.0136	0.0169	11.6969	7.2688
Winequality-red-4	0.0010	0.0035	0.0355	0.0060	0.0069	0.0076	0.0746	1.3710	1.9317
Wdbc	0.0007	0.0034	0.0175	0.0105	0.0152	0.0153	0.0208	4.1194	2.7673

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