

# Prediction of Sleep Stages Based on Wearable Signals Using Machine Learning Techniques

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**Abstract:** Sleep's impact on mood and health is widely recognized by medical researchers with such understanding disseminating among average people in recent years. The main objective of this work was the development of machine learning algorithms for automatic sleep cycles detection. The features were selected based on the AASM manual, which is considered the gold standard for human technicians. For training the models we used MESA, a database containing 2056 full overnight unattended polysomnographies. With the goal of developing an algorithm that would only require a photoplethysmography (PPG) device to be able to accurately predict sleep stages and quality, the main channels used from this dataset were peripheral oxygen saturation and PPG. Testing the performance of Random forest, Gradient Boosting, Gaussian Naive-bayes, K-Nearest Neighbours, Support Vector Machine and Multilayer Perceptron classifiers, and using features extracted from the dataset, we achieved 80.50 % accuracy, 0.7586 Cohen's kappa, and 77.38% F1-score, for five sleep stages, using a Multilayer Perceptron. To assess its performance in a real-world scenario we acquired sleep data and compared the classifications attributed by a popular sleep stage classification android app and our algorithm, resulting in a strong level of agreement (90.96% agreement, 0.8663 Cohen's kappa), for four sleep stages.

## 1 INTRODUCTION

Sleep's impact on mood and health is widely recognized by medical researchers with such understanding disseminating among average people in recent years. While newer studies strengthen the suspected link between inadequate sleep and a wide range of infirmities (Minkel et al., 2012), the general population is not very conscious of their sleep quality. As a result, there is much interest in having proper means of studying sleep, given its importance and how difficult it is to accurately diagnose sleep disorders, considering how individuals are affected by sleep loss, and their ability to recover from said sleep loss, varies significantly (Tkachenko and Dinges, 2017). The discovery of the brain's electrical activity was the main contributor responsible for the development of the field of sleep medicine in the second half of the 20th century. The examination of the electroencephalogram (EEG) patterns that occur during sleep lead to the current division of the sleep period into different stages, thus

creating the basis of sleep medicine and the study of human sleep (Worley, 2018). One of the major discoveries was that sleep is much more restorative to both waking cognition and health when it occurs and goes through the appropriate physiological sequences. This is to say that, due to the way that sleep is structured into distinct stages, where each one has a certain set of characteristics and its own physiological role, the exclusive measurement of the amount of time slept is not enough for the quality of sleep to be determined. As such, sleep quality depends not only on total time slept but on many other factors such as fragmentation, amount of time spent in each sleep stage, and how the sleep cycles are structured.

Currently, polysomnography (PSG) is the most common technique used to study sleep disorders, being able to record multiple biosignals simultaneously (Rundo and Downey, 2019; Karlen et al., 2009). It has, however, the issue of being expensive and inconvenient (Kelly et al., 2012), with the fact that this type of exam is normally performed

in a clinic additionally raising the issue of negative bias, as people may behave differently than normal when they know they are being monitored. Furthermore, there is the matter of longitudinal data. Laboratory PSGs are usually a single-night snapshot, whereas sleep is a dynamic process that is affected by the existence and intensity of many other factors that vary from day to day.

Wearable sleep-trackers, on the other hand, are low-cost devices capable of measuring biosignals and, from this data, inferring information about certain behaviours, like sleep. They present many advantages over PSG, such as their convenience, ease of use, affordability and data accessibility, with the possibility of using some sort of cloud-based platform for storage of data, thus allowing the acquisition of an unprecedented amount of information about sleep and other behaviours or health parameters.

Because of the costs and time expenditures associated with PSG, there is much interest in the development of algorithms that can be deployed in a wearable device, being able to automatically and accurately classify sleep stages with a similar degree of accuracy as the current gold-standard. Ideally, such an algorithm should also strive to be as simple as possible (both in terms of signals used and model complexity).

### 1.1 State-of-the-Art

As sleep disorders are common in modern society, with the main difficulty of their treatment being detection and diagnosis (Pavlova and Latreille, 2019), and with the recent increase in popularity of using wearable devices in medicine (Akkaş et al., 2020), some studies have already been developed on the performance of models for sleep stage prediction using different biosignals, features or classifiers.

In the literature about the performance of these kinds of wearables, it was possible to find information about *Fitbit Charge 2*, which records wrist activity through accelerometers and pulses through photoplethysmography (PPG). In Stucky et al. (2021), the authors compared this device against portable home PSG, displaying reasonably accurate mean values of sleep and heart rate (HR) estimates, should it follow careful data processing. One other device is the *Heally Recording System* which, through the combination of embedded sensors and electrodes in a shirt that measures respiratory and cardiac physiology, monitors sleep based on autonomic signals. It exhibited accuracy at approximately 80% agreement with manual scoring,

which is similar to accuracies obtained through actigraphy, considered an appropriate method for the assessment of sleep in patients with certain sleep disorders (McCall and McCall, 2012).

Other studies, relying on ML, have successfully developed algorithms for sleep stage prediction. For example, Tsinalis et al. (2020) managed to obtain sleep stage-specific characteristics with an average accuracy of 86% based on EEG data, while Yildirim et al. (2019), developed and applied a 19-layer 1D convolutional neural network model to EEG and EOG signals, achieved the highest classification accuracies for 5 of its 6 sleep classes as over 91%.

More specifically for studies using the same dataset (that will be described in the next section) that was chosen for this work, we have Kudo et al. (2022) that, using PPGs and accelerometers' information extracted from the public datasets Apple watch Sleep (Walch et al., 2019), and Multi-Ethnic Study of Atherosclerosis (MESA) (Zhang et al., 2018; Chen et al., 2015), achieved a macro F1 score of 0.655 and Cohen's kappa score of 0.527, using a recurrent neural network. Another similar study, published by Sridhar et al. (2020), using the ECG signal of both the Sleep Heart Health Study (Quan et al., 1997) and the MESA dataset for training, validation and testing of the developed algorithm, obtained an overall performance of 77% accuracy and 0.66 Cohen's kappa against the reference stages on a held-out portion of the datasets used for training.

All these studies suggest that the development of similar fully automatic recognition systems could serve as a suitable replacement for manual inspection of PSG signals, particularly for large-scale studies.

## 2 DATASET DESCRIPTION

Initially, the algorithm was trained through the use of a publicly available online database, selected from others such as the NCH Sleep DataBank (Lee et al., 2021), or the Sleep Heart Health Study. After a comparison between several of these databases, one was selected based on its size, sensor quality and quantity, detail of the scoring, and how recently collected was the data.

The set of PSG recordings used for this work was obtained from MESA. This dataset included a sleep exam with 2237 participants, consisting of full overnight unattended polysomnographies that were conducted between 2010-2012, and had the following demographics described in Table 1.

Table 1: Dataset demographics of the MESA database (adapted from (NSRR team, 2022)).

Characteristics	Value
Number of PSGs	2056
Number of Patients	2237
Age	(Years)
Mean	69.6
Median	69.0
Standard deviation	$\pm 9.2$
Minimum	54.0
Maximum	95.0
Gender	
Female	1198
Male	1039
Race/ethnicity	
White, Caucasian	830
Chinese American	265
Black, African-American	616
Hispanic	526

The information pertaining to the PSG studies in the MESA dataset are contained in two separate file formats for each study. The EDF files store 27 biosignal channels, from which we used only three, HR information, the PPG recording, and oxygen saturation. With the exception of the PPG signal that was sampled at 256 Hz, the channels were sampled at 1 Hz. On the other hand, the XML files contain annotations corresponding to the PSG recordings, such as sleep stages and their duration.

For the real-world validation of the developed models, 14 nights of sleep were acquired using a biosignalsPlux device on the posterior side of the left wrist (Plux Wireless Biosignals, 2022) that recorded both PPG and accelerometer data, and a smartwatch (Ticwatch E2) on the right wrist.

### 3 METHODOLOGY

In order to build the code developed in this work to analyse and process the dataset, as well as build the ML models, Python was used through the code editor Spyder. Several different libraries were used, including BeautifulSoup4, Pandas, NumPy, scikit-learn, Tensorflow, and hranalysis. After the development of the algorithms, to test them in a real-world scenario, they were used to classify the sleep stages of the acquired 14 nights of sleep. These classifications were then compared to the results

obtained from ‘‘Sleep as Android’’ (Chaudhry, 2017), which is one of the most reviewed android sleep analysis smartphone applications, using the measurements taken using the smartwatch.

In the next Sub-Sections, the algorithms used to perform the classification of the data, the metrics upon which they are evaluated, and both which features and how they were extracted are described.

#### 3.1 Data Pre-Processing and Feature Extraction

The first step of the extraction of the data from the PPG records was the standardization of the signal, achieved through the subtraction of the signal’s mean followed by its division by its standard deviation. After that, the signal was segmented in short windows (half second interval) and the mean of each of these intervals was subtracted to minimize baseline drift. Subsequently a 4th order Chebyshev II bandpass filter (sampling frequency of 256 Hz and cut-off frequencies of 0.05 and 30 Hz) was used.

At this stage we segmented the signal according to the sleep stage annotations and began extracting features. These features, a total of 30, range from the maximum, mean and minimum values of oxygen saturation and HR, in this case also including its standard deviation, to features related to heart rate variation (HRV). The resulting analysis of HRV is grouped under time-domain and frequency-domain.

In the time-domain, 12 features were used, such as root mean square of successive differences between N-N intervals ( $RMSSD$ ), standard deviation of these differences ( $SDSD$ ), number of pairs of successive N-N intervals that differ by more than 50 ms and 20 ms ( $NN50$  and  $NN20$ ), total proportion of  $NN50$  and  $NN20$  in relation to the total number of N-N intervals, standard deviation of all N-N intervals (calculated over each 30 second interval), mean and median of the N-N intervals ( $Mean\_nni$  and  $Median\_nni$ ), coefficient of variation ( $SDNN$  divided by  $Mean\_nni$ ), coefficient of variation of successive differences ( $RMSSD$  divided by  $Mean\_nni$ ) and, finally, the difference between the longest and shortest N-N interval.

As for the frequency-domain, seven features were used, including total power spectral density (Golgouneh and Tarvirdizadeh, 2020), power in the very low ( $vlf$ ), low ( $lf$ ), and high ( $hf$ ) frequency bands (Salahuddin et al., 2007), normalised  $lf$  and  $hf$  power, and the ratio between these two powers.

Two additional features related to the PPG signal’s entropy (more specifically fuzzy (Chen et al., 2007) and dispersion entropy (Rostaghi and

Azami, 2016)) were extracted after averaging its value in windows of 32 samples, to minimize time spent for this step and the information loss resulting from the averaging.

Finally, despite only classifying sleep in 30 second intervals, the two preceding stage classifications were also used as features, so as to take into account the continuity of sleep.

### 3.2 Classification Models

To classify the sleep stages, we used both machine learning models (such as Random Forest, Gradient Boosting, Gaussian Naive-Bayes, K-Nearest Neighbours, and Support Vector Machine) and artificial neural networks (Multilayer Perceptrons).

The choice of these algorithms was based on both literature reviews done for other sleep stage classification studies, and trial and error.

#### 3.2.1 Random Forest

Random Forest is an ensemble learning method that constructs and uses numerous decision trees. Due to random variable selection and bootstrap aggregation leading to lower correlation across trees, the ensemble prediction is generally more accurate than any of its decision trees individual predictions.

#### 3.2.2 Gradient Boosting

Gradient Boosting is an ensemble learning method of weak prediction models, usually decision trees. With careful tuning of its parameters, it may result in better performance than Random Forest models.

#### 3.2.3 Gaussian Naive-Bayes

Gaussian Naive Bayes classifiers are based on applying Bayes’ theorem with a strong independence assumption to classify the data.

#### 3.2.4 K-Nearest Neighbours

K-Nearest Neighbours (KNN) classifiers utilise proximity to make predictions. For classification problems, a class label is assigned to a data element based on the vote of the K number of its nearest neighbours. It is possible to construct a weighted version using the distance between data points.

#### 3.2.5 Support Vector Machine

Support-Vector Machine (SVM) algorithms attribute classifications by finding a hyperplane in an N-

dimensional space that is able to separately classify the data points.

### 3.2.6 Multilayer Perceptron

Multilayer Perceptrons are a fully connected class of feedforward artificial NNs, consisting of at least three layers of nodes. With the exception of the nodes in the input layer, each node is a neuron that uses a nonlinear activation function.

### 3.3 Model’s Hyperparameters

The characteristics of machine learning algorithms are strongly tied to their hyperparameters, with their optimization and tuning being pivotal to a model’s performance (Feurer and Hutter, 2019). For the non-neural network models, the chosen method to tune these hyperparameters was grid search, which is a tuning technique that computes their optimum values through an exhaustive search in a manually introduced subset of values. Scikit-learn library’s implementation of this function was used, with the hyperparameters’ values being presented in table 2.

Table 2: Values for the different parameters to be optimized when utilizing scikit-learn’s grid search.

	Parameters	Values
Random Forest	n_estimators	10-100, 100-1000 (increasing by 10 and 100, respectively)
	Criterion	gini, entropy
	max_depth	None, 10-100 (increasing by 10)
Gradient Boosting	n_estimators	10-100, 100-1000 (increasing by 10 and 100, respectively)
	Criterion	friedman_mse, squared_error, mse
	max_depth	1,3,5, 10-100(increasing by 10)
KNN	n_neighbours	1, 3, 5, 7, 9, 11
	Weights	uniform, distance
	Metric	manhattan, Euclidean
SVM	C	0.0001, 0.01, 0.05, 0.1, 0.5, 1.0, 5, 10
	Kernel	linear, poly, rbf
	Gamma	scale, 0.0001, 0.01, 0.05, 0.1, 0.5, 1.0, 5, 10

For the neural network models, their hyperparameters were chosen to be tuned through trial and error due to their increased complexity.

### 3.4 Model Evaluation

After training the algorithms, it is necessary to evaluate their performance. To do this, the MESA dataset was first split into testing and training sets, so as to permit an assessment and minimization of the impact of the model’s overfitting to the data, which would otherwise lead to an imprecise estimation of the model’s capabilities. Additionally, as sleep stages’ distribution is naturally unbalanced (Worley, 2018), to promote a more even learning process, these sets were balanced.

Finally, some metrics were calculated to evaluate their performance. These metrics were accuracy, Cohen’s kappa and macro average F1-score.

## 4 RESULTS

For the non-neural network models, after optimizing their hyperparameters through grid search, the evaluated metrics for the best performing models of each type obtained are presented in Table 3.

Table 3: Values of the chosen metrics for the highest performance non-neural network models of each type.

	Accuracy (%)	Cohen's kappa
Random Forest	79.30	0.7412
Gradient Boosting	82.34	0.7792
Gaussian Naive-Bayes	68.41	0.6052
KNN	21.99	0.0249
SVM	25.03	0.0628

As can be observed in Table 3, the best performing models are Random Forest and Gradient Boosting, with this last model presenting an overall more balanced performance for all of its classifications when compared to the other models and presenting the highest accuracy and Cohen’s kappa for the balanced test dataset.

For the neural network models, the first step of tuning its architecture was selecting the number of layers and neurons per layer. Accordingly, starting by the hidden layer number, it was discovered that models with three layers are optimal (Figure 1).

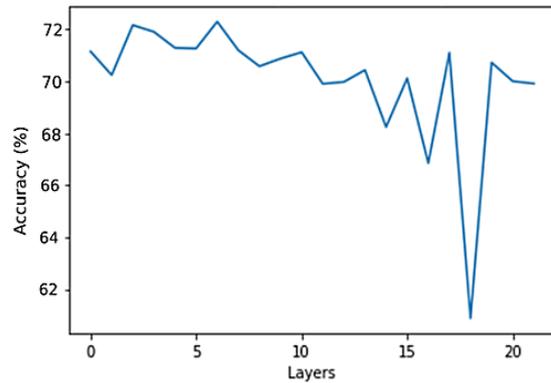


Figure 1: Model accuracy per number of layers.

Following this, the optimal neuron count per layer for this 3-layered model was found (Figure 2).

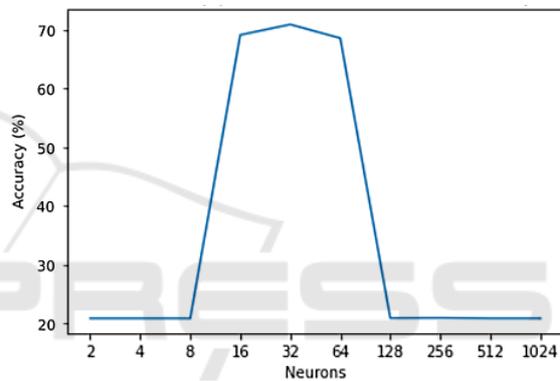


Figure 2: Model accuracy per number of neurons in each layer.

Finally, to reduce overfitting the influence of several regularization methods such as L1, L2 and dropout were tested, with only L2 regularization having a positive influence in the performance of the developed models (Figure 3).

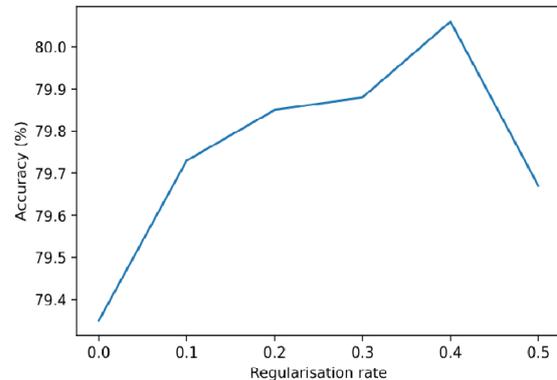


Figure 3: Correlation between maximum model accuracy and L2 regularisation rate.

In this figure, it is possible to observe that the best result is achieved when a L2 regularisation rate of 0.4 is used, with this model presenting an accuracy of 80.50%, Cohen’s kappa of 0.7563 and F1 score of 77.38 % (in the unbalanced test set). An example of the resulting classification can be seen in figure 4, where the orange lines are the true sleep stages and the blue lines are the model’s predictions, with the blue lines disappearing when they match.

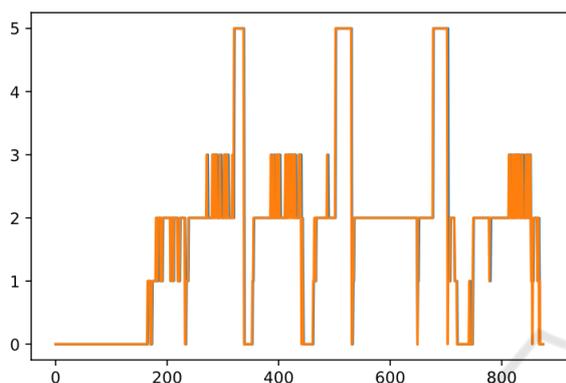


Figure 4: Sleep stages for each interval of a randomly chosen MESA file, classified by the developed algorithm.

Using this final model to classify the sleep night data that was recorded during this work, an agreement of 90.96%, Cohen’s kappa of 0.8663, and macro average F1-score of 90.52% was achieved when compared to the classifications attributed by the Android app, with these results being displayed in figure 5.

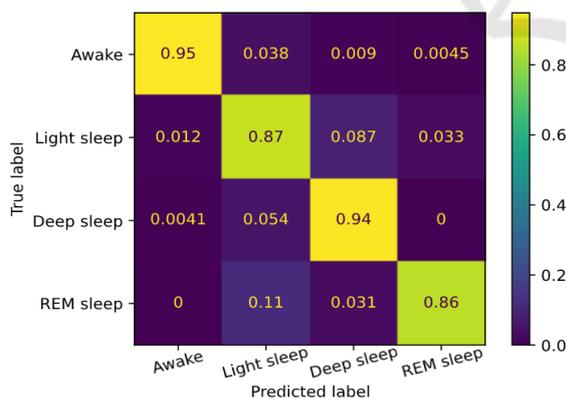


Figure 5: Normalized confusion matrix of the results obtained from the classification of real-world data.

## 5 DISCUSSION

First it is important to notice that, usually, models are stochastically trained, meaning that two models

with the matching architecture being trained with identical data in the same manner, might perform differently after training, which may further complicate the study and understanding of the training process. To solve this issue several identical models with the same characteristics were developed, at which point their average performance was evaluated, and then compared with the average performance of other models with different architectures.

As mentioned previously, some commonly used regularisation methods, such as L1, L2, or dropout, were tested. In the case of the latter, despite usually being described as improving model performance (Baldi and Sadowski, 2014; Srivastava et al., 2014), it failed to do so in this case, instead leading to a decrease in performance (even only 5% dropout lowers average accuracy to 41.84%). This decrease seems tied to dropout probability, where the higher the probability, the worse the performance is, until a plateau is reached at approximately 37.35% accuracy. The addition of L1 regularisation also seems to be detrimental to model development, with the higher the rate, the worse its impact on the model’s accuracy. On the other hand, L2 regularisation seems to improve the effectiveness of the models, and, while we found a regularisation rate of 0.4 to be optimal, there seems to be a wide range of values (from 0.1 to 2) where the model still benefits from its addition.

With this said, for the NNs, the best performing model presented 80.50% accuracy, 0.7563 Cohen’s kappa on the balanced test set, and a macro average F1-Score of 77.38% on the complete, unbalanced test set. After an extensive search for the optimal configuration of hyperparameters, we found that the model consistently performed better in a 3 hidden layer, 32 neurons per layer, structure, with all hidden layers having a L2 regularisation rate of 0.4. Overall, we found that performance tends to be highest for models with 3 or 7 layers, with it dropping sharply outside these limits. Similarly for neuron count, accuracy dropped to around 20% for any number of neurons per layer outside of the interval between 16 and 64, whereas it seems mostly stable at around 80% accuracy and optimal at 32 neurons per layer.

The results obtained are promising as, while some models are able to achieve higher accuracy (Tsinalis et al., 2016; Yildirim et al., 2019), they do so while using more signals (usually EEG, EOG, or ECG), which significantly restricts their usability for everyday applications. Conversely, we reached better performance than many other models, including recently published studies that make use of

more signals or features (Sun et al., 2020), or employ the same dataset (Kudo et al., 2022; Sridhar et al., 2020).

For the classification of the real-world data that was acquired, a neural network was chosen over the other Gradient Boosting model, as even though its performance on the balanced test dataset was slightly inferior to the best performing non-neural network model, its performance on stage 2 classification (one of the most common stages for naturally-occurring unbalanced sleep) is substantially improved, which leads to this model being superior for real-world stage classifications (0.7586 Cohen's kappa in the complete, unbalanced test dataset, in contrast to 0.6967) without being as deleterious to lowest class accuracy (52.95% compared to 56.32% accuracy). Additionally, the increase in misclassifications by this model tends to be between physiologically similar stages (such as between stage 1 and stage 2, which are both usually considered light sleep, for example), which lowers the importance of such errors. This neural network being the most accurate algorithm developed is in line with the current state-of-the-art, as the model's increased complexity theoretically allows it to more accurately classify the different sleep stages.

After this selection, the device's data was scored by our algorithm, and then compared with the classifications by the Android application, at which point a strong level of agreement (McHugh, 2012) (90.96% accuracy, 0.8663 Cohen's kappa and a macro average F1-Score of 90.52%) was observed.

## 6 CONCLUSION

This work's main objective was the development of a ML algorithm that detects and classifies sleep cycles. For this end, both NN and non-NN models were developed.

The performance achieved for the final NN model was higher than many other studies, despite generally using a lesser amount of features or signals and the same or similar datasets.

Another goal of this work was to test the developed model's performance in a real-world scenario. To achieve this, we simultaneously recorded 14 nights of sleep using a biosignalsPlux device with PPG and accelerometer sensors and a widely used Android sleep scoring application paired with a commercially available wearable device. After comparing the resulting classifications we obtained a strong level of agreement. This leads

us to believe in the potential of the developed algorithm to be used in real-world scenarios.

While the main goals of this work were fulfilled, it still presents some limitations that could be improved, namely in terms of feature acquisition and extraction.

Future studies should attempt to integrate these algorithms into devices. This way, not only is it possible to increase the similarity between the devices and algorithms being compared, but it should also be easier to acquire a larger amount of data, ideally, from a larger set of individuals as well.

The recording of more data itself would also likely lead to improvements in the determination of the real-world performance of the models, besides the potential use of this data for model training. In this regard, the recording and comparison of results with a PSG study would be optimal.

Additionally, during feature extraction, we chose to reduce the number and quality of the entropies used as features, due to time and computation constraints. As, even after this, these were some of the most relevant features, the extraction and use of them without averaging the signal beforehand could lead to some performance improvements.

Finally, throughout this work several models were created, some of them having similar levels of accuracy and other selected metrics to the final model developed. Due to this, a complementary study that could be done is the creation of another ensemble model that utilises the output of these models as inputs, as these types of models tend to have a better performance than the sum of their parts (Zhang and Ma, 2012).

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