# Judges System for Classifiers Specialization 

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#### Abstract

In this paper we designed and implemented a new ensemble of classifiers based on a sequence of classifiers which were specialized in regions of the training dataset where errors of its trained homologous are concentrated. In order to separate this regions, and to determine the aptitude of each classifier to properly respond to a new case, it was used another set of classifiers built hierarchically. We explored a selection based variant to combine the base classifiers. We validated this model with different base classifiers using 37 training datasets. It was carried out a statistical comparison of these models with the well known Bagging and Boosting, obtaining significantly superior results with the hierarchical ensemble using Multilayer Perceptron as base classifier. Therefore, we demonstrated the efficacy of the proposed ensemble, as well as its applicability to general problems.


Keywords-classifiers, delegation, ensemble

## I. Introduction

EVERY new emerging classification problem on Bioinformatics applications need to be more accurate. In addition, every day becomes more difficult to achieve this with the well-known simple models. There are techniques arising based on combining multiple models into a single one to achieve better results in those problems that needs more accuracy in classification. These models are known as ensemble of classifiers. In Statistics, and mainly the Artificial Intelligence, it has been advanced enough in this way. There are several models on the literature, some ones that base their operation in diversifying the training dataset for each classifier [1-3], some others managing the parameters of the models [4, 5] or simply combining different models [6] to avoid errors correlation.

In this work a new multiclassifier is presented inspired by the necessity of specializing classifiers in different sectors of the training dataset to achieve an increase of its effectiveness. The problem resides in how to separate these sectors, what is a difficult task keeping in mind the great quantity of features that characterizes most of the problems, and the little information we had of them. For these reason, it becomes necessary to build these regions based on what we really know, the performance of classifiers on each training case.

## II. Datasets

In order to validate this model, 11 bases were chosen from

[^0]TABLE I
VALIDATION DATASETS (BIOINFORMATICS OR BIOMEDICAL PROBLEMS)

| Dataset | Nominal <br> features | Numeric <br> features | Class <br> labels | Cases |
| :--- | :--- | :--- | :--- | :--- |
| audiology | 69 | 0 | 24 | 226 |
| breast-cancer | 0 | 4 | 3 | 625 |
| Diabetes | 0 | 8 | 2 | 768 |
| Heart-c | 7 | 6 | 2 | 303 |
| Heart-h | 7 | 6 | 2 | 294 |
| Heart-statlog | 0 | 13 | 2 | 270 |
| Horse-colic | 4 | 23 | 2 | 300 |
| Hypothyroid | 22 | 7 | 4 | 3772 |
| Lung-cancer | 56 | 0 | 3 | 32 |
| promoters | 57 | 0 | 2 | 106 |
| Yeast | 0 | 8 | 10 | 1484 |

the UCI Repository [7] representing Bioinformatics or Biomedical problems. These datasets are shown on Table I.
This is a fewer population for any statistical comparison. More bases are needed to give the tests the potential to decide if any significant difference exists between the classical and the novel model. For this reason, 26 general bases were picked

TABLE II
GENERAL DATASETS FOR VALIDATION

|  | GENERAL DATASETS FOR VALIDATION |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Dataset | Nominal <br> features | Numeric <br> features | Class <br> labels | Cases |
| autos | 8 | 17 | 6 | 205 |
| balance-scale | 4 | 0 | 3 | 625 |
| balloons | 4 | 0 | 2 | 76 |
| cars | 6 | 0 | 4 | 1728 |
| colic | 15 | 7 | 2 | 368 |
| credit-a | 9 | 6 | 2 | 690 |
| credit-g | 13 | 7 | 2 | 1000 |
| glass | 0 | 9 | 6 | 214 |
| hayes-roth | 4 | 0 | 3 | 132 |
| hepatitis | 13 | 6 | 2 | 155 |
| ionosphere | 0 | 34 | 2 | 351 |
| Iris | 0 | 4 | 3 | 150 |
| kr-vs-kp | 36 | 0 | 2 | 3196 |
| labor | 8 | 8 | 2 | 57 |
| lenses | 4 | 0 | 3 | 23 |
| liver-disorders | 0 | 6 | 2 | 345 |
| lymph | 2 | 16 | 4 | 148 |
| monks | 6 | 0 | 2 | 415 |
| postoperative | 7 | 1 | 3 | 90 |
| segment | 0 | 19 | 7 | 2310 |
| shuttle-land | 5 | 1 | 2 | 15 |
| sick | 12 | 7 | 2 | 3772 |
| sonar | 0 | 60 | 2 | 208 |
| soybean | 35 | 0 | 19 | 683 |
| vote | 16 | 0 | 2 | 435 |
| wine | 0 | 13 | 3 | 178 |
|  |  |  | 2 |  |

up too, so a very varied group was collected. This quantity of bases constitutes an enough piece of data to carry out a correct
statistical analysis of the results that there will be shown. We will test over the whole set, and look for a similar behavior on the Bioinformatics and Biomedical bases. Table II shows the additional data.

## III. Ensemble Model

As it was said previously, we are presenting a model built by using training subsets. Therefore, each classifier can be specialized in the region of the base where all previous ones failed. Some techniques already use this method [2, 8]; they reinforce the learning of the classifiers in the cases that they have misclassified. Once carried out the above-mentioned, the challenge consists in how to be able to detect how much sure can we be each classifier can predict the objective feature of a new case correctly, and not to make a simple voting (like previous models) where it is rejected in what region of the training database it is expert each classifier. In order to explain our proposal, we should specify how to obtain the diversity of the base classifiers, avoiding correlation among their errors; and then, we should define the method used to combine their outputs.

## A. Diversity

The general idea of the algorithm we are proposing is based on going building a group of classifiers specialized in regions of the training dataset. Therefore, they will leave separating progressively the subsets of cases that have not been still well classified, or in fact, they are not probably well classified, for any classifier in the system. So that, at each step, get focus o the learning interest on the cases that it interests us to learn.

We need to divide the dataset in such a way that each classifier is trained to reach the highest accuracy taking into account its capabilities. Another external classifier will evaluate these capabilities or judge that suggests which classifier can be the more suitable or specific on each subset, or analysis level of the original dataset.

We ensure the diversity by changing the dataset for each classifier, so we decided to use an only one model for the base classifiers, that is, the same classifier is used iteratively in the learning of more and more restricted groups. So, when we speak of the first (second...) classifier, really speaks of the classifier it was trained in the first (second...) training subset. For a specific problem, it demands of course, that we should decide which classifier model will be used, keeping in mind the characteristics of the problem. We should also define the pattern of the simple classifier (external or judge) to evaluate the performance of the base classifiers on each iteration, that is, to separate the cases in the successive subsets of training.

The proposed multiclassifier system is constituted by levels. In the first level, we meet the most general classifier; it is able to classify any type of cases, but at the same time with fewer yields. The following one is limited to the cases that the first one did not probably classify well, therefore it is less general than the previous one, it is not able to classify any kinds of cases, but in those that were trained, it should obtain better results. This process repeats until arriving at the last level,
where we meet the classifier that trains with the most specific group, with highest performance in a much-reduced group of cases: those that have been more difficult for the rest of the classifiers. Therefore in this work, the level $i$ will be related with the training of the classifier $C_{\mathrm{i}}$ and of the judge-classifier $J_{\mathrm{i}}$, that evaluates which cases can be probably well classified by $C_{\mathrm{i}}$ and separates the remaining cases to constitute the base of training of the most specific classifier $C_{i+1}$.
Figure one shows three levels system, that is, three judges ( $J_{0}, J_{1}$ and $J_{2}$ ) and four classifiers $\left(C_{0}, C_{1}, C_{2}\right.$ and $C_{3}$ )


Fig. 1 Three levels ensemble. $D C_{0} \subseteq D C_{1} \subseteq D C_{2} \subseteq D C_{3}$.
The steps can be formalized as fallows:

1. Train classifier $C_{i}$ corresponding to level $i$.
2. Train judge-classifier $J_{i}$ in order to separate those misclassified cases $C_{i}$.
3. Construct a new training subset $D C_{i+1}$ with those cases $J_{i}$ decides will probably be misclassified by $C_{i}$.
It is not the same the cases that were misclassified by $C_{i}$ than the cases that $J_{i}$ decided as misclassified. If we neglect this difference, the system can be in a fiasco. The ideal conditions, of course, are that $J_{i}$ decides with $100 \%$ of truthfulness, but this is not probable for none of the existing classifiers up to now.
The process should be repeated until completing some stop condition and it can have many to prove. In this work, we will not be plentiful in this condition, we will simply fix the maximum number of levels and we will iterate until arriving to the same one. Therefore, we must add this final step to the procedure described above:
4. If $D C_{i+1}$ is not empty neither the maximum number of iterations is reached, go to the next level having $D C_{i+1}$ as training dataset.
Notice the stop condition can be reached by the last level, or if the classifier of a given level correctly classifies all cases.

## B. Output combination

Already trained the classifiers, in such a way there is no correlation in their errors, we just need to define how to combine their outputs to classify a new case. The classifiers $J_{i}$ have been used to define the limits of the training regions for each classifier $C_{i+1}$, therefore they are those who will take the leading roll in the process of combination of the outputs.

We should have present that is necessary to carry out the combination on the base of the contained knowledge at classifiers $J_{i}$. In figure 1 depicts how $J_{0}$ separates the training base in two subsets; let us call to whole training base $U$, and $B$ to the group formed to train $C_{l}$. Then the cases that should be classified by $C_{0}$ are those that belong to the subset $U \backslash B$, and another specialized classifier must classify $B$ subset: $C_{I}$. The same thing happens at each level. We can obtain for each $J_{i}$ how probable is that the classifier $C_{i}$ correctly classify a given case or not. Following this idea, the probability $P_{i}$ that the case x will be well classified by the classifier $C_{i}$ can be represented as:

$$
\begin{equation*}
P_{i}(x)=\left(1-P J_{f}(x)\right) \cdot\left(1-P J_{1}(x)\right) \cdots\left(1-P J_{i-1}(x)\right) \cdot P J_{i}(x) \tag{1}
\end{equation*}
$$

Where $P J_{i}(\mathrm{x})$ is the probability that the classifier $J_{i}$ decides that $C_{i}$ should classify the case x , for $i>0$. This expression represents the probability that $C_{i}$ contributes a good answer, conditioned to all the previous ones have possibly responded in wrong way. Clearly $P_{0}(\mathrm{x})=P J_{0}(\mathrm{x})$. Once calculated the conditional probabilities, we will have the system chooses the answer given by the classifier that will respond to each case with more certainty. The learning algorithms and classification are as they continue:

## Algorithm 1. Training of the model:

1. Initialize $i \leftarrow 0$.
2. Build $C_{i}$ instance $C$ and train it with $U$.
3. Build $B_{i}$ from $U$ changing the goal feature as fallows:
a. Right if $C_{i}$ well classified the case, or
b. Wrong in other case.
4. Build $S_{i}$ instance of $S$ and train it with $B_{i}$.
5. Suppress from $U$ those cases that $S_{i}$ classifies as Wrong.
6. If $U$ is empty, $N \leftarrow i$.
7. If $N>i+1, i \leftarrow i+1$ go to 2 .
8. if $U$ is not empty, build $C_{N+1}$ instance of $C$ and train it with $U$

## Algorithm 2. Classification process by mean of selection:

1. For each level $i$, compute by expression (1), the probability $P_{i}(x)$ that $x$ must be classified by Ci
2. Look for the level with the highest value of $P_{i}$, then classify x by $C_{i}$.

## IV. Results and discusion

In order to validate this method, we present a comparison between the accuracy obtained by our method and results achieved by Bagging and AdaBoost.M1 that are described as very effective and efficient methods in bases of general classification problems in the literature (Opitz and Maclin, 2000). The results are obtained by means of a 10 -fold crossvalidation. As it is evident, we cannot demonstrate the new model's superiority making a simple comparison of these results. It was used for the same one, the statistical tests of related populations' comparison, in particular non-parametric tests: Friedman (more than two populations) and Wilcoxon (two populations).

TABLE III
Classification accuracy results for Bagging and AdaBoost.M1 MODELS

| Datasets | Bagging |  |  | AdaBoost.M1 |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
|  | J48 | SMO | MLP | J48 | SMO | MLP |
| audiology | 0.801 | 0.783 | 0.819 | 0.841 | 0.796 | 0.845 |
| breast-cancer | 0.814 | 0.872 | 0.938 | 0.770 | 0.869 | 0.925 |
| diabetes | 0.751 | 0.775 | 0.763 | 0.717 | 0.766 | 0.759 |
| heart-c | 0.776 | 0.848 | 0.825 | 0.815 | 0.828 | 0.799 |
| heart-h | 0.793 | 0.844 | 0.806 | 0.782 | 0.847 | 0.793 |
| heart-statlog | 0.804 | 0.837 | 0.837 | 0.785 | 0.840 | 0.796 |
| horse-colic | 0.710 | 0.673 | 0.707 | 0.717 | 0.697 | 0.710 |
| hypothyroid | 0.995 | 0.936 | 0.952 | 0.996 | 0.947 | 0.950 |
| lung-cancer | 0.531 | 0.438 | 0.438 | 0.531 | 0.344 | 0.375 |
| promoters | 0.840 | 0.915 | 0.906 | 0.915 | 0.943 | 0.906 |
| yeast | 0.590 | 0.571 | 0.596 | 0.563 | 0.571 | 0.577 |

Tables III and IV sample the results obtained for the accuracy in the classification on Bioinformatics and

TABLEIV
CLASSIFICATION ACCURACY RESULTS FOR THE PROPOSED MODEL

| CLASSIFICATION ACCURACY RESULTS FOR THE PROPOSED MODEL |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| Datasets |  | Novel model |  |  |
|  | J48 |  | SMO |  |
| audiology | 0.721 | 0.819 | 0.845 |  |
| breast-cancer | 0.959 | 0.969 | 0.963 |  |
| diabetes | 0.764 | 0.775 | 0.764 |  |
| heart-c | 0.785 | 0.835 | 0.828 |  |
| heart-h | 0.813 | 0.857 | 0.820 |  |
| heart-statlog | 0.807 | 0.841 | 0.815 |  |
| horse-colic | 0.683 | 0.707 | 0.743 |  |
| hypothyroid | 0.995 | 0.936 | 0.953 |  |
| lung-cancer | 0.777 | 0.892 | 0.858 |  |
| promoters | 0.858 | 0.934 | 0.943 |  |
| yeast | 0.570 | 0.524 | 0.594 |  |

Biomedical bases, using the models mentioned with a decision tree (J48), Support Vector Machine (SMO) and Multilayer Perceptron (MLP). The statistical validation shows that the results of the three models, using J48 and SMO are, at least comparable.

TABLE V
Classification Accuracy Results for Bagging and AdaBoost.M1 MODELS IN THE REST OF DATASETS

| Datasets |  |  |  |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | J48 |  |  | Sagging | AdaBoost.M1 |  |  |  |
|  | SMO | MLP | J48 | SMO | MLP |  |  |  |
| Autos | 0.800 | 0.712 | 0.751 | 0.873 | 0.737 | 0.761 |  |  |
| balance-scale | 0.814 | 0.872 | 0.938 | 0.770 | 0.869 | 0.925 |  |  |
| balloons | 0.724 | 0.763 | 0.763 | 0.776 | 0.737 | 0.724 |  |  |
| Cars | 0.933 | 0.936 | 0.997 | 0.955 | 0.942 | 0.994 |  |  |
| Colic | 0.856 | 0.832 | 0.834 | 0.823 | 0.804 | 0.799 |  |  |
| Credit-a | 0.859 | 0.848 | 0.864 | 0.826 | 0.826 | 0.833 |  |  |
| Credit-g | 0.740 | 0.748 | 0.754 | 0.691 | 0.747 | 0.706 |  |  |
| Glass | 0.734 | 0.542 | 0.687 | 0.776 | 0.598 | 0.701 |  |  |
| Hayes-roth | 0.712 | 0.803 | 0.795 | 0.720 | 0.818 | 0.826 |  |  |
| hepatitis | 0.813 | 0.858 | 0.865 | 0.852 | 0.832 | 0.781 |  |  |
| ionosphere | 0.912 | 0.877 | 0.912 | 0.934 | 0.889 | 0.912 |  |  |
| Iris | 0.953 | 0.960 | 0.947 | 0.953 | 0.980 | 0.940 |  |  |
| kr-vs-kp | 0.993 | 0.960 | 0.995 | 0.996 | 0.973 | 0.992 |  |  |
| Labor | 0.807 | 0.895 | 0.912 | 0.877 | 0.930 | 0.912 |  |  |
| lenses | 0.783 | 0.696 | 0.696 | 0.652 | 0.826 | 0.696 |  |  |
| liver-disorders | 0.722 | 0.591 | 0.728 | 0.661 | 0.655 | 0.687 |  |  |
| lymph | 0.777 | 0.858 | 0.838 | 0.784 | 0.858 | 0.838 |  |  |
| monks | 0.590 | 0.636 | 0.547 | 0.561 | 0.627 | 0.564 |  |  |
| postoperative | 0.689 | 0.700 | 0.622 | 0.578 | 0.656 | 0.656 |  |  |
| segment | 0.976 | 0.928 | 0.970 | 0.982 | 0.928 | 0.966 |  |  |
| shuttle-land | 0.600 | 0.533 | 0.600 | 0.733 | 0.667 | 0.667 |  |  |
| Sick | 0.988 | 0.939 | 0.976 | 0.990 | 0.952 | 0.968 |  |  |
| Sonar | 0.774 | 0.774 | 0.841 | 0.793 | 0.793 | 0.846 |  |  |
| soybean | 0.925 | 0.931 | 0.934 | 0.921 | 0.922 | 0.939 |  |  |
| Vote | 0.970 | 0.959 | 0.954 | 0.954 | 0.952 | 0.954 |  |  |
| wine | 0.933 | 0.989 | 0.978 | 0.949 | 0.978 | 0.983 |  |  |

TABLE VI
CLASSIFICATION ACCURACY RESULTS FOR THE PROPOSED MODEL IN THE REST OF DATASETS

| Datasets | Novel model |  |  |
| :---: | :---: | :---: | :---: |
|  | J48 | SMO | MLP |
| autos | 0.829 | 0.737 | 0.771 |
| balance-scale | 0.803 | 0.878 | 0.920 |
| balloons | 0.697 | 0.763 | 0.789 |
| cars | 0.918 | 0.933 | 0.994 |
| colic | 0.856 | 0.845 | 0.845 |
| credit-a | 0.858 | 0.855 | 0.843 |
| credit-g | 0.717 | 0.750 | 0.728 |
| glass | 0.692 | 0.561 | 0.701 |
| hayes-roth | 0.705 | 0.826 | 0.818 |
| hepatitis | 0.819 | 0.871 | 0.832 |
| ionosphere | 0.900 | 0.889 | 0.920 |
| Iris | 0.960 | 0.967 | 0.973 |
| kr-vs-kp | 0.995 | 0.962 | 0.995 |
| labor | 0.842 | 0.982 | 0.930 |
| lenses | 0.848 | 0.739 | 0.783 |
| liver-disorders | 0.672 | 0.583 | 0.707 |
| lymph | 0.777 | 0.892 | 0.858 |
| monks | 0.614 | 0.627 | 0.576 |
| postoperative | 0.711 | 0.689 | 0.589 |
| segment | 0.970 | 0.923 | 0.968 |
| shuttle-land | 0.600 | 0.600 | 0.689 |
| sick | 0.989 | 0.939 | 0.973 |
| sonar | 0.764 | 0.779 | 0.841 |
| soybean | 0.922 | 0.934 | 0.939 |
| vote | 0.970 | 0.959 | 0.959 |
| wine | 0.923 | 0.987 | 0.983 |

Tables V and VI show the other results. Statistical tests showed that significant difference exists in favor of the proposed ensemble using MLP when being also compared with their homologous with MLP. Finally, this last one was compared with the previous models, using the three classification models, as base classifiers, and the differences were significant.

## V. Conclusions

In this paper, we presented a new ensemble of classifiers applicable to general problems, and in particular to Bioinformatics and Biomedical problems. It was built based on the specialization of the base classifiers on regions on the training Dataset divided by judges taking into account the local performance of the classifiers. Finally, it was statistically proved the novelty model is useful for different sort of problems, offering better results than Bagging and AdaBoost.M1.

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