Model selection based on combined penalties for biomarker identification

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ABSTRACT

The growing role of targeted medicine has led to an increased focus on the development of actionable biomarkers. Current penalized selection methods that are used to identify biomarker panels for classification in high dimensional data, often result in highly complex panels that need careful pruning for practical use. We propose a stepwise forward variable selection method which combines the L0, with L1 or L2 norms. The penalized likelihood criterion that is used in the stepwise selection procedure results in more parsimonious models, keeping only the relevant features. Simulation results and a real application show that our approach exhibits a comparable performance with common selection methods with respect to the prediction performance whilst minimizing the number of variables in the selected model resulting in a more parsimonious model as desired.

INTRODUCTION

In the biomarker research area, penalization techniques provide an attractive approach to their variable selection and are rather appealing due to their sparse representation and good predictive accuracy. In genomic research, an L1 penalty [1] is routinely used due to its convexity and optimization simplicity. However, the result of the L1 type regularization may not be sparse enough for a good interpretation. The development of methods to obtain sparser solutions is becoming essential part in the classification and feature selection area. We proposed a stepwise forward approach for model selection in the framework of penalized regression using a penalty that combines the L0 norm, which is based on the number of coefficients, with L1 norm which is based on the size of coefficients or L2 norm which take into account the grouping effect (i.e. keep in the model groups of variables that are correlated).

The aim is to find a model that includes as few relevant variables as possible with good predictive performance. This is an important consideration for classification and screening applications where the goal is to develop a test using as few features as possible.

METHODS

We consider the logistic regression model with logit link function (logit(p) = xβ). The vector of n binary responses is y = (y1, y2, ..., yn) and X is an n x d matrix of predictors. Under the regularization framework, the estimated coefficients β̂ = (β̂1, β̂2, ..., β̂d) are found by minimizing the objective function $\hat{\beta} = \text{argmin}_{\beta \in \mathbb{R}^d} L(\beta) + \lambda P(\beta)$

where P(β) is a regularization term and $\lambda$ a tuning parameter.

The combined L0 + L1 penalty

Following Liu and Wu [2] the penalization term is defined as $CL_{d}^\alpha(\beta) = \sum_{j=1}^{d} \max(0, \alpha - |\beta_j|^{\alpha})$

where $L_0^\alpha(\beta) = 1$, $|\beta_j| < \alpha$ is a continuous approximation to L0 and $0 \leq \alpha \leq 1$

The estimated coefficients are obtained by minimizing the objective function $\hat{\beta} = \text{argmin}_{\beta \in \mathbb{R}^d} L(\beta) + \sum_{j=1}^{d} \max(0, \alpha - |\beta_j|^{\alpha})$

However, the applicability of the proposed $CL_{d}^\alpha$ penalty was restricted to moderate datetimes due to the nonconvexity of $CL_{d}^\alpha$.

We consider a stepwise heuristic approach where at each step we obtain the estimated coefficients by:

$$\hat{\beta}_j = \text{argmin}_{\beta_j} \{-\log L(\beta) + \lambda \sum_{j=1}^{d} \max(0, \alpha - \beta_j^{\alpha})\}$$

The selected model is obtained on the criterion that minimizes $\hat{\beta}$

$\hat{\beta}$

Algorithm:

Given a set of $d$ standardized predictors $X = X_1, ..., X_d$, and a binary response $y \in \{0,1\}$:

Step 1: Consider all the univariate models $M_1: Y = \beta_1 X_1$, $M_2: Y = \beta_2 X_2$, ..., $M_d: Y = \beta_d X_d$

Keep $M_j \in \{1, ..., d\}$ that minimizes (1)

Step 2: With the model chosen in step 1, e.g. $M_2$, and in an additive way we consider all the $d-1$ models ($M''$) by adding the remaining $d-1$ variables one at a time to the model $M_2$:

$M_1': Y = \beta_2 X_2 + \beta_1 X_1$

$M_2': Y = \beta_2 X_2 + \beta_1 X_1$

Keep $M_j^{(2)}$ of the models that minimizes the function in (1)

Step 3: Continue until the value of the function (1) in the current step is bigger than the previous step.

The combined L0 + L1 penalty

We also consider another penalty combination, the L10 (variable selection) with L2 norms (grouping effect) and the penalization term is the following:

$CL_{d}^\alpha(\beta) = \sum_{j=1}^{d} \max(0, \alpha - |\beta_j|^{\alpha}) + \lambda |\beta_j|$.

Figure 1: Two-dimensional contour plot of CL03. L1, L2, CL2, CL0, with $\alpha = 0.1$

RESULTS

We examine the performance of the proposed stepwise method with the above introduced combined penalties (stepCL, stepCL2) via simulations and a real data application. We include the global minimization (CL, CL2) for a comparison, and we also consider the results from Lasso and the adaptive Lasso. We compare the different methods • in terms of model complexity • the classification performance

All the functions that were used for the combined penalty approach can be found in the R-package “stepPenal”, available on CRAN.

Tuning of parameters $\alpha$, $\lambda$: 5-fold Cross Validation was used and the Area Under the ROC curve on the validation set is estimated over a grid of values of $\alpha$ and $\lambda$. The $(\lambda, \alpha)$ that yields the 1sd from the maximum AUC is selected.

In this poster we will only present results from the real data application. Simulation results can be found in [2]

The real data come from a biomarker study with baseline measurements of d=187 proteins from n=53 patients. The objective is to extract potential candidate markers discriminating responders from non-responders.

In order to evaluate the performance of the models and in the absence of an external validation dataset we use bootstrapping. We applied all the methods on B=1000 bootstrapped datasets of the protein data, by sampling with replacement. Figure 2 shows that the classification performance of the stepwise methods stepCL and stepCL2 is as good as the other variable selection methods (Lasso and adaptive Lasso), albeit including the least predictors.

Figure 2: Boxplots of the total Variables selected by the models (top panel) and the AUC of the ROC curves (bottom panel) over the B=1000 bootstrapped data of the protein data

CONCLUSION

We have proposed a stepwise forward approach for model selection in the framework of penalized regression using a penalty that combines the L0 with L1 or L2 norms. The aim is to find a model that includes as few relevant variables as possible on the one hand, and have good predictive performance on the other hand. The combined penalization term $CL_{d}^\alpha(\beta)$ that was introduced by Liu and Wu [3] was limited to moderate datasets due to limitations of the optimization algorithm. Considering the heuristic stepwise forward approach, we can apply the penalization $CL_{d}^\alpha(\beta)$ and $CL_{d}^2(\beta)$ to high-dimensional data by using the BFGS algorithm in a stepwise approach which is found to work well in practice for nonconvex and non-smooth functions [4]. As future work we also consider to apply our method to regression problems for variable selection with a continuous response as well as time-to-event data.

REFERENCES


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