Identifying interesting SNP interactions with logicFS

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1 Introduction

Logic regression proposed by Ruczinski et al. (2003) is a classification method that attempts to predict the case-control status based on Boolean combinations of binary variables. It has already been successfully applied to SNP data by Kooperberg et al. (2001).

This package contains functions that use logic regression as a wrapper to identify interesting combinations of binary variables and to measure the importance of these interactions. A description of the used methods is given in Schwender and Ickstadt (2006). Even though the intended purpose of this package is the identification of SNP interactions, it can also be applied to other types of binary (or categorical) data.

logicFS also contains a first basic Bagging version of the logic regression and a fast implementation of the Quine-McCluskey algorithm based on matrix algebra. The latter is described in Schwender (2006) and can be used to identify a minimum disjunctive normal form of a given truth table.

2 SNP Data and their Transformation into Dummy Variables

As always, the package has to be loaded.

> library(logicFS)

Loading required package: LogicReg

As example data set, we use the data set contained in logicFS.

```
> data(data.logicfs)
```

data(data.logicfs) consists of two objects: A simulated matrix, data.logicfs, containing the data of 15 variables (columns) and 400 observations (rows) and a vector, cl.logicfs, consisting of the class labels of the 400 observations.

Each of the variables in data.logicfs is categorical and can take the realizations 1, 2 and 3. The class label of the first 200 observations is 1 (for case) and the class label of the remaining observations is 0 (for control). If one of the following expression is TRUE, then the corresponding observation is a case:

```
SNP1 == 3

SNP2 == 1 \& SNP4 == 3

SNP3 == 3 \& SNP5 == 3 \& SNP6 == 1
```

where SNP1 is in the first column of data.logicfs, SNP2 in the second, and so on.

Even though the variables are called SNPs, these variables have not been generated by an algorithm for simulating SNP data. So most of the values of a variable can, e.g., be 3 even though 3 should actually code the homozygous variant genotype.

Logic regression and hence the functions in logicFS can only handle binary predictors. So the categorical SNP variables have to be transformed into two binary dummy variables. If the homozygous reference genotype of the SNPs is coded by 1, the heterozygous genotype by 2 and the homozygous variant type by 3, then the function make.snp.dummy can be used to code these SNPs. Thus, a binary representation of the variables in data.logicfs can be obtained by

> bin.snps <- make.snp.dummy(data.logicfs)

SNPs are coded as follows:

```
SNP SNP_1 SNP_2 Assumed Genotype
1 1 0 0 Homozygous Reference
2 2 1 0 Heterozygous
3 3 1 Homozygous Variant
```

3 Feature Selection Using Logic Regression

The binary dummy variables in bin.snps can now be used in logic.fs to identify potentially interesting combinations of these variables and to measure the importance of these interactions. Since the default version of the logic regression and thus the current version of logic.fs is based on simulated annealing, we set

```
> my.anneal <- logreg.anneal.control(start = 2, end = -2, iter = 10000)
```

and the number B of considered logic regression models to 20 to speed up the computation. The feature selection is then performed by

```
> log.out <- logic.fs(bin.snps, cl.logicfs, B = 20, nleaves = 10,
+ rand = 123, anneal.control = my.anneal)</pre>
```

where we allow that a maximum of nleaves = 10 variables is in the logic regression models and set rand to 123 to make the results of this vignette reproducible.

The output of logic.fs is given by

> log.out

Selection of Interactions Using Logic Regression

Number of Iterations: 20

Logic Regression Type: Classification

Max. Number of Leaves: 10

The 5 Most Important Interactions:

	${\tt Importance}$	${\tt Proportion}$	Expression
1	27.55	0.95	!SNP2_1 & SNP4_2
2	16.05	0.95	SNP1_2
3	10.20	0.35	SNP3_2 & SNP5_2 & !SNP6_1
4	0.70	0.20	SNP5_2 & !SNP6_1
5	0.60	0.05	SNP3_2 & SNP4_2 & SNP5_2

Not very surprisingly, only the interactions shown in Section 2 have a high importance, where, e.g., $!SNP2_1$ stands for NOT $SNP2_1$, i.e. $SNP2_1 = 0$.

Thus, the logic expression !SNP2_1 & SNP4_2 means "SNP2 is of the homozygous reference genotype and SNP4 is of the homozygous variant genotype."

As displayed in Figure 1, the variable importance can be plotted by

> plot(log.out)

Since the names of SNPs are usually pretty long, coded names are used in the plot, where X1 codes the first variable in data, X2 the second, and so on. The original variable names are displayed in the plot if coded is set to FALSE in plot.

In the above analysis, the single tree approach of the logic regression is used. It is, however, also possible to perform a multiple tree analysis by changing the default of ntrees. Note that if this approach is applied to bin.snps one will get a lot of warnings (from glm) because of the structure of this data set (cases are unambiguously classified by the above logic expressions). With other data sets this usually will not happen.

Single Tree Measure

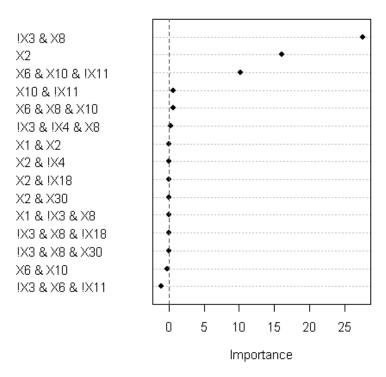


Figure 1: Variable Importance Plot

4 Bagged Logic Regression

A first basic Bagging version of the logic regression is implemented in the function logic.bagging. Similar to the analysis with logic.fs logic.bagging is applied to bin.snps by

```
> bag.out <- logic.bagging(bin.snps, cl.logicfs, B = 20, nleaves = 10,
+ rand = 123, anneal.control = my.anneal)
> bag.out
```

Bagged Logic Regression

Number of Iterations: 20

Logic Regression Type: Classification

Max. Number of Leaves: 10

OOB Error Rate: 2%

By default, both the out-of-bag error (obb=TRUE) and the importance of the variable combinations (importance=TRUE) are computed. The results of Section 3 can also be obtained by

> bag.out\$vim

The 5 Most Important Interactions:

	${\tt Importance}$	${\tt Proportion}$	Expression
1	27.55	0.95	!SNP2_1 & SNP4_2
2	16.05	0.95	SNP1_2
3	10.20	0.35	SNP3_2 & SNP5_2 & !SNP6_1
4	0.70	0.20	SNP5_2 & !SNP6_1
5	0.60	0.05	SNP3_2 & SNP4_2 & SNP5_2

and the importance can be plotted either by plot(bag.out\$vim) or simply by plot(bag.out).

The class of a new observation is predicted by majority voting and can be computed by

> cl.preds <- predict(bag.out, bin.snps)</pre>

where we here assume that bin.snps contains the new observations. The misclassification rate is then computed by

> mean(cl.preds != cl.logicfs)

[1] 0

Warning: There is currently no checking if the data set, newbin, containing the new observations and data in logic.bagging contain the same variables in the same order.

References

- Kooperberg, C., Ruczinski, I., LeBlanc, M., and Hsu, L. (2001), "Sequence Analysis Using Logic Regression," *Genetic Epidemiology*, 21, 626–631.
- Ruczinski, I., Kooperberg, C., and LeBlanc, M. (2003), "Logic Regression," Journal of the Computational and Graphical Statistics, 12, 475–511.
- Schwender, H. (2006), "A new Implementation of the Quine-McCluskey Algorithm Using Matrix Algebra," Technical Report, SFB 475, University of Dortmund, Germany, To appear.
- Schwender, H., and Ickstadt, K. (2006), "Identification of SNP Interactions Using Logic Regression," Technical Report, SFB 475, University of Dortmund, Germany. To appear soon.