The mgsa package

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1 Introduction

Model-based Gene Set Analysis (MGSA, Bauer et al. [1]) is a Bayesian modeling approach for gene set enrichment. The package *mgsa* implements MGSA and tools to use MGSA together with the Gene Ontology [2].

2 Quick start

We start with a small simulated dataset which contains example_go, a random subset of yeast gene ontology annotations with 20 terms and example_o, a simulated set of observed positive genes. These genes could for example be the "hits" of some screen or a set of differentially expressed genes. In the simulation, the terms GO:0006109 and GO:0030663 were active, implying that genes annotated to these terms were more likely to be observed positives than other genes.

```
> library(mgsa)
[1] "/project/mgsa.Rcheck mgsa"
[1] "Package mgsa initialized"
> data("example")
> example_go
Object of class MgsaSets
10 sets over 158 unique items.
Set annotations:
                               term
                                                 definition
GO:0046292 formaldehyde metabol... The chemical reactio...
GO:0006109 regulation of carboh... Any process that mod...
GO:0008113 peptide-methionine-(... Catalysis of the rea...
GO:0016849 phosphorus-oxygen ly... Catalysis of the cle...
{\tt G0:0046527} glucosyltransferase ... Catalysis of the tra...
... and 5 other sets.
Item annotations:
                            name
SFA1
        Bifunctional enzyme ...
YJL068C Non-essential intrac...
```

```
ADR1
        Carbon source-respon...
CAT8
        Zinc cluster transcr...
FYV10
        Protein of unknown f...
... and 153 other items.
> example_o
 [1] "SFA1"
                "ADR1"
                           "CAT8"
                                      "FYV10"
                                                 "GCR1"
                                                            "GCR2"
                                                                       "GID7"
 [8] "HAP2"
                "HAP3"
                           "HAP4"
                                      "HAP5"
                                                            "PCL6"
                                                                       "PCL7"
                                                 "PCL10"
[15] "PCL8"
                "PFK26"
                           "PFK27"
                                      "PH085"
                                                 "PIG1"
                                                            "PIG2"
                                                                       "REG1"
[22] "SIP4"
                "SNF1"
                                      "TYE7"
                                                            "UBP14"
                           "SNF4"
                                                 "UBC8"
                                                                       "VID28"
[29] "YLR345W" "GSC2"
                           "CCT5"
                                      "CPR6"
                                                 "CPR7"
                                                            "HSC82"
                                                                       "PET100"
[36] "TIM9"
                "COP1"
                           "GL03"
                                      "RET2"
                                                 "RET3"
                                                            "SEC21"
                                                                       "SEC26"
[43] "SEC27"
```

The method mgsa fits the MGSA model. It returns a MgsaMcmcResults object whose print method displays the most likely active terms. On this example, mgsa correctly reports largest posterior probabilities for the terms GO:0006109 and GO:0030663. The call to set.seed(), which sets the seed of the random number generator, simply ensures the example of this vignette to be reproducible. It is not required for mgsa() to work.

```
> set.seed(0)
> fit = mgsa(example_o, example_go)
> fit

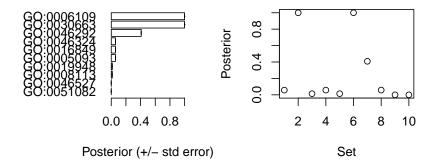
Object of class MgsaMcmcResults
158 unique elements in population.
43 unique elements both in study set and in population.
'data.frame': 10 obs. of 4 variables:
$ inPopulation: int 1 34 2 1 2 8 2 1 21 86
$ inStudySet : int 0 28 0 0 0 7 1 0 1 6
$ estimate : num 0.057 1 0.0138 0.0576 0.0138 ...
$ std.error : num 6.77e-04 6.96e-06 1.90e-04 6.01e-04 2.79e-04 ...
NULL
```

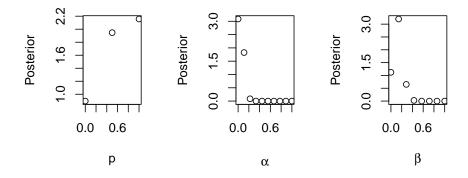
Posterior on set activity (decreasing order):

```
inPopulation inStudySet estimate
                                                 std.error
GD:0006109
                                28 0.9999794 6.961322e-06
GD:0030663
                      8
                                 7 0.9999464 1.683033e-05
GO:0046292
                      2
                                 1 0.4086602 1.695734e-03
GO:0046324
                      1
                                 0 0.0579632 6.940961e-04
GO:0016849
                                 0 0.0576434 6.006926e-04
                      1
GD:0005093
                      1
                                 0 0.0570154 6.773733e-04
GO:0019948
                      2
                                 0 0.0138470 2.788480e-04
                      2
GO:0008113
                                 0 0.0138426 1.898030e-04
GO:0046527
                     21
                                 1 0.0000000 0.000000e+00
                                 6 0.0000000 0.000000e+00
GO:0051082
```

The method plot provides a graphical visualization of the fit.

```
> plot(fit)
```





3 Using the Gene Ontology

The Gene Ontology [2]

References

- [1] S. Bauer, J. Gagneur and P. N. Robinson. GOing Bayesian: model-based gene set analysis of genome-scale data. *Nucleic acids research*, 2010.
- [2] The Gene Ontology Consortium. Gene Ontology: tool for the unification of biology. *Nature Genetics*, 25:25–29,2000.