strum package - examples

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This document contains the whole analysis process for the first three example models from the introduction document. Note that the input data used for these examples are not necessarily simulated to give a meaningful result for each analysis.

```
> library(strum)
```

1 Genetic association analysis

This is an example of a typical genetic association analysis model with a latent trait (similar to MIMIC model). Suppose that there are three measurements (P1, P2 and P3), and it is hypothesized that there is a single latent trait (L1) underlying the three measurements. The latent variable L1 is influenced by a SNP and a set of variance components, polygenic(p) and random environmental(e). Each trait is also influenced by its own random environmental factor. This is the model diagram.
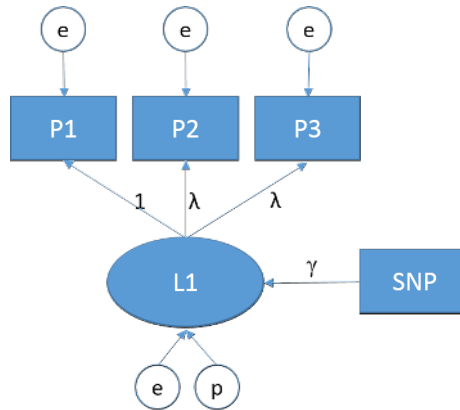

Figure 1: Genetic association analysis model

1.1 Construct model

The first step is to construct a **strumModel** object specifying the above model using *createStrumModel()* function.

```
> assoForm1 =
+    'L1 =~ P1 + P2 + P3 + <e>
+     L1 ~ aSNP + <p,e>
+     '
> myAssoModel = createStrumModel(formulas = assoForm1)

 Creating strumModel ........................... Done

> myAssoModel

Basic properties of the model:
        Model Class ................... strumModel
        Ascertainment ..................... FALSE

List of all variables:
       Obs Covariate InEita   InY Exogen.
L1    FALSE     FALSE   TRUE FALSE   FALSE
aSNP   TRUE      TRUE  FALSE FALSE      NA
P1     TRUE     FALSE  FALSE  TRUE      NA
P2     TRUE     FALSE  FALSE  TRUE      NA
P3     TRUE     FALSE  FALSE  TRUE      NA

Model formulas:
  L1 =~ P1 + P2 + P3 + <e>
  L1 ~ aSNP + <p,e>
```

1.2 Prepare data

The next step is to prepare data. In this example, the data must be a data.frame with 4 required fields - family, id, father, mother, since the model includes the polygenic variance component (p). To run a strum analysis, you need to construct a **strumData** object created by the *createStrumData()* function with a data.frame. The following code shows the step using the example input file "chr1Ped.csv".

```
> dName = system.file("extdata/example_ped.csv", package = "strum")
> dF = read.csv(dName, header=T)[,1:18]
> names(dF) = c("family","id", "father","mother",names(dF)[5:18])
> myAssoData = createStrumData(dF, "Pedigree")

 Creating strumData ............................. Done

> myAssoData
```

```
Data type: Pedigree
Data size: 477 entries, 18 variables

First 5 rows of data values:
  family id father mother sex disease proband        P1         P2         P3
1      1  1      0      0   0       0       0  0.4093955 0.44450079 -0.3867515
2      1  2      0      0   1       0       0 -1.5037814 1.52582608  0.8832360
3      1  3      1      2   0       0       0  1.5850090 0.08833692  0.9322619
4      1  4      1      2   1       0       0  1.6246356 0.60065352  1.0895325
5      1  5      1      2   1       0       0 -0.4111477 0.08588345 -0.6477336
    SBP     DBP     A1     A2      S1      S2 aSNP rs6040343
1  3.39  4.1800  1.300  1.630  1.9000 -0.0613    1         1
2 -3.49 -2.7200 -0.784 -3.550 -2.8900 -2.1300    0         0
3 -3.40  0.0815 -1.820 -4.390 -1.6700 -3.0900    1         0
4 -7.04 -3.6500 -0.183 -4.740 -2.9800 -2.3500    0         0
5  4.60  4.9900  2.440 -0.117 -0.0408 -0.4350    1         1

phi object contains  75  matrices:
First matrix:
$`1`
    1   2   3   4   5   6   7
1 1.0 0.0 0.5 0.5 0.5 0.5 0.5
2 0.0 1.0 0.5 0.5 0.5 0.5 0.5
3 0.5 0.5 1.0 0.5 0.5 0.5 0.5
4 0.5 0.5 0.5 1.0 0.5 0.5 0.5
5 0.5 0.5 0.5 0.5 1.0 0.5 0.5
6 0.5 0.5 0.5 0.5 0.5 1.0 0.5
7 0.5 0.5 0.5 0.5 0.5 0.5 1.0

Empty IBD object.
```

1.3   Run analysis

Now, run the association analysis by the function call *strum()* with two previously constructed objects as the arguments.

```
> myAssoResult = strum(myAssoModel, myAssoData)

 Start STRUM analysis ...
   Fitting model step 1 ......................... Done
   Fitting model step 2 ......................... Done
   Testing model fit ............................ Done

Analysis completed!
```

1.4 Result

The result object contains the model description and two result tables. The first table contains the fitted parameter values with standard errors, confidence intervals, and p-values. The second table contains the information on the model fit from four different measures. For association analysis, you would test H_0: $\gamma = 0$ versus H_1: $\gamma \neq 0$. In this model, γ is the parameter L1 $\sim$ aSNP, which equals to 0.9445929 with the pvalue = 1.591018e-21.

```
> myAssoResult

=========
  Model
=========

Basic properties of the model:
        Model Class ............. strumFittedModel
        Ascertainment ..................... FALSE

List of all variables:
      Obs Covariate InEita   InY Exogen.
L1   FALSE     FALSE    TRUE FALSE    FALSE
aSNP  TRUE      TRUE   FALSE FALSE       NA
P1    TRUE     FALSE   FALSE  TRUE       NA
P2    TRUE     FALSE   FALSE  TRUE       NA
P3    TRUE     FALSE   FALSE  TRUE       NA

Model formulas:
  L1 =~ P1 + P2 + P3 + <e>
  L1 ~ aSNP + <p,e>


==========
  Result
==========

Parameter estimates:
                estimate    stdError      lowerCI    upperCI       pValue
L1=~P2         0.9885029  0.05314867   0.88433341  1.0926724  3.284230e-77
L1=~P3         1.0105280  0.05822325   0.89641252  1.1246435  1.773988e-67
L1~aSNP        0.9445929  0.09913024   0.75030117  1.1388846  1.591018e-21
P1~[intercept] 0.2880181  0.13404076   0.02530298  0.5507331  3.165543e-02
P2~[intercept] 0.1829933  0.13133455  -0.07441771  0.4404043  1.635179e-01
P3~[intercept] 0.3505190  0.12747443   0.10067375  0.6003643  5.964614e-03
P1~~P1<e>      1.1318519  0.18199793   0.77514251  1.4885613  2.501075e-10
P2~~P2<e>      1.2400615  0.14775629   0.95046454  1.5296585  2.377245e-17
P3~~P3<e>      0.6908993  0.19387808   0.31090526  1.0708934  1.829182e-04
```

```
L1~~L1<p>      0.9445110 0.26947683  0.41634613 1.4726759 2.283205e-04
L1~~L1<e>      1.1478948 0.20752417  0.74115493 1.5546347 1.588615e-08

Chi-square fit statistics:
                           kappa   chiStat df     pValue
Un-adjusted            1.0000000  2.719114  7 0.90971460
Mean adjusted          0.2057025 13.218676  7 0.06695628
Mean-Variance adjusted 0.1599908 16.995440  9 0.04878743
Theoretically corrected       NA 11.586595  7 0.11500000

Model fit indices:
                             value
Comparative Fit Index (CFI)      1
```

2 Genetic linkage analysis

In this section, we show an example of a typical genetic linkage analysis model with a latent trait using IBD information. Suppose again that there are three measurements as above (P1, P2 and p3) and a single latent trait (L1) underlying the three measurements. The latent variable L1 is influenced by a set of genetic and random variance components. Each trait is also influenced by its own random environmental factor. The model diagram looks like following.
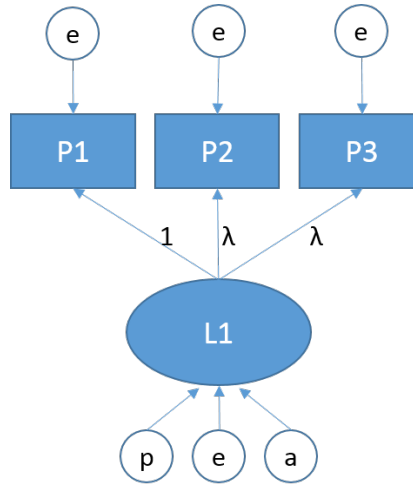
Figure 2: Genetic linkage analysis model

2.1 Construct model

The above linkage model can be constrcuted as a **strumModel** object using *createStrumModel()* function.

```
> linkForm1 =
+    'L1 =~ P1 + P2 + P3 + <e>
+     L1 ~ <a,p,e>
+    '
> myLinkModel = createStrumModel(formulas = linkForm1)

 Creating strumModel ............................ Done

> myLinkModel

Basic properties of the model:
        Model Class ................... strumModel
```

```
        Ascertainment ..................... FALSE
```

```
List of all variables:
     Obs Covariate InEita   InY Exogen.
L1 FALSE     FALSE    TRUE FALSE   FALSE
P1  TRUE     FALSE   FALSE  TRUE      NA
P2  TRUE     FALSE   FALSE  TRUE      NA
P3  TRUE     FALSE   FALSE  TRUE      NA

Model formulas:
  L1 =~ P1 + P2 + P3 + <e>
  L1 ~ <a,p,e>
```

2.2 Prepare data

From the above linkage analysis model, *a* represents the major gene variance components, which requires the ibd information to be imported. The ibd information for the family data can be imported by specifying the name of ibd file into *ibdFileName* argument for *createStrumData()*. The use of the example ibd file "GENIBD.chr1Ped.ibd", which contains the ibd information of family data in "chr1Ped.csv", is shown in the following code. We use the data.frame, dF, created for the previous association analysis model.

```
> iName = system.file("extdata/GENIBD.chr1Ped.ibd", package = "strum")
> myLinkData = createStrumData(dF, "Pedigree", ibdFileName=iName)

 Importing S.A.G.E. IBD file ..................... Done
 Creating strumData ............................. Done
```

2.3 Run analysis

Now, run the linkage analysis by the function call *strum()*. If you want to perform the linkage analysis on all markers exist in the IBD file, you don't need to specify the marker name as an argument for *strum()* function. In this case, each marker will be analysed one by one, and the result object will contain a list of the linkage analysis results for all markers.

```
> myLinkResultAll = strum(myLinkModel, myLinkData)
```

To analyze a subset of IBD markers, then you can specify the names of them as follows;

```
> mNames = c("chr1marker1", "chr1marker2")
> myLinkResult = strum(myLinkModel, myLinkData, ibdMarkers=mNames)

 Start STRUM analysis ...
```

```
  chr1marker1:
   Fitting model step 1 .......................... Done
   Fitting model step 2 ......................... Done
   Testing model fit ........................... Done

  chr1marker2:
   Fitting model step 1 ......................... Done
   Fitting model step 2 ......................... Done
   Testing model fit ........................... Done

Analysis completed!
```

2.4 Result

The result object again contains the model description and result tables. The first table contains the fitted parameter values with standard errors, confidence intervals, and p-values. The second table contains the information on the model fit from four different measures. For linkage analysis, you would test H_0: $\alpha = 0$ versus H_1: $\alpha \neq 0$. In this model, α is the parameter L1 $\sim\sim$ L1<a>, which equals to 0.2787365 with the pvalue = 2.594076e-01.

```
> myLinkResult[[1]]

=========
  Model
=========

Basic properties of the model:
          Model Class ............. strumFittedModel
          Ascertainment ...................... FALSE

List of all variables:
      Obs Covariate InEita   InY Exogen.
L1 FALSE       FALSE   TRUE FALSE   FALSE
P1  TRUE       FALSE  FALSE  TRUE      NA
P2  TRUE       FALSE  FALSE  TRUE      NA
P3  TRUE       FALSE  FALSE  TRUE      NA

Model formulas:
  L1 =~ P1 + P2 + P3 + <e>
  L1 ~ <a,p,e>


==========
  Result
==========
```

```
Parameter estimates:
                  estimate   stdError    lowerCI   upperCI        pValue
L1=~P2           0.8678181 0.07587710  0.7191018 1.016535 2.726799e-30
L1=~P3           0.8914807 0.09331546  0.7085858 1.074376 1.254982e-21
P1~[intercept]   1.1238196 0.11825192  0.8920501 1.355589 2.027470e-21
P2~[intercept]   1.0432875 0.11808091  0.8118532 1.274722 9.977489e-19
P3~[intercept]   1.1673092 0.12331581  0.9256147 1.409004 2.907264e-21
P1~~P1<e>        0.8806093 0.30655087  0.2797806 1.481438 2.035310e-03
P2~~P2<e>        1.3941225 0.19786635  1.0063116 1.781933 9.221350e-13
P3~~P3<e>        0.6740625 0.21180676  0.2589289 1.089196 7.301963e-04
L1~~L1<p>        1.3112519 0.52164084  0.2888546 2.333649 5.973493e-03
L1~~L1<e>        1.4094234 0.35748785  0.7087601 2.110087 4.030537e-05
L1~~L1<a>        0.2788592 0.43189124 -0.5676320 1.125351 2.592465e-01


Chi-square fit statistics:
                           kappa   chiStat df    pValue
Un-adjusted            1.0000000 3.246504 10 0.9750134
Mean adjusted          0.4180031 7.766697 10 0.6516137
Mean-Variance adjusted 0.3483359 9.320036 12 0.6753792
Theoretically corrected       NA 9.066546 10 0.5258000


Model fit indices:
                             value
Comparative Fit Index (CFI)      1
```

3 Structural Equation Model

This is an example of a SEM model with latent variables and polygeneic effect. Suppose that there are six measurements and three underline latent variables. anger is a latent variable which underlies the two measurements (A1, A2), bp is a latent variable which underlies the two measurements (SBP, DBP) and stress is a latent variable which underlies the two measurements (S1, S2). bp is caused by anger and stress, and stress is caused by anger and a SNP (rs6040343). All traits and latent variables are also influenced by their own polygenic and random variance components except stress, which the variance is fixed at 0.1 for both polygenic and random componants. The model diagram looks like following.
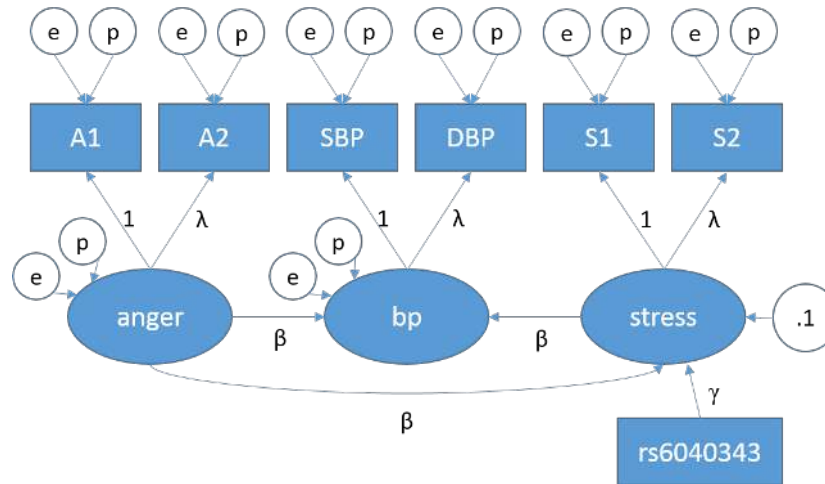

Figure 3: Structural equation model

3.1 Construct model

The above SEM model can be constrcuted as a **strumModel** object using *createStrumModel()* function.

```
> semForm1 =
+    'bp =~ SBP + DBP
+     anger =~ A1 + A2
+     stress =~ S1 + S2
+     bp ~ anger + stress
+     stress ~ anger + rs6040343
+     var(stress) = .1
+     '
> mySemModel = createStrumModel(formulas = semForm1)
```

```
  Creating strumModel ............................ Done

> mySemModel

Basic properties of the model:
        Model Class ................. strumModel
        Ascertainment ..................... FALSE

List of all variables:
          Obs Covariate InEita  InY Exogen.
bp       FALSE    FALSE   TRUE FALSE   FALSE
anger    FALSE    FALSE   TRUE FALSE    TRUE
stress   FALSE    FALSE   TRUE FALSE   FALSE
rs6040343 TRUE     TRUE  FALSE FALSE     NA
SBP       TRUE    FALSE  FALSE  TRUE     NA
DBP       TRUE    FALSE  FALSE  TRUE     NA
A1        TRUE    FALSE  FALSE  TRUE     NA
A2        TRUE    FALSE  FALSE  TRUE     NA
S1        TRUE    FALSE  FALSE  TRUE     NA
S2        TRUE    FALSE  FALSE  TRUE     NA

Model formulas:
  bp =~ SBP + DBP
  anger =~ A1 + A2
  stress =~ S1 + S2
  bp ~ anger + stress
  stress ~ anger + rs6040343
  var(stress) = .1
```

3.2 Prepare data

The next step is to prepare data. Note again that the data must be a data.frame with 4 required fields - family, id, father, mother, since the model includes the polygenic variance component (p) by default. A **strumData** object is created by *createStrumData()* function with a data.frame. Again, we use the data.frame, dF, created above using the example input file "chr1Ped.csv".

```
> mySemData = createStrumData(dF, "Pedigree")

  Creating strumData ............................. Done

> mySemData

Data type: Pedigree
Data size: 477 entries, 18 variables

First 5 rows of data values:
```

```
   family id father mother sex disease proband        P1         P2         P3
1      1  1      0      0   0       0       0   0.4093955 0.44450079 -0.3867515
2      1  2      0      0   1       0       0  -1.5037814 1.52582608  0.8832360
3      1  3      1      2   0       0       0   1.5850090 0.08833692  0.9322619
4      1  4      1      2   1       0       0   1.6246356 0.60065352  1.0895325
5      1  5      1      2   1       0       0  -0.4111477 0.08588345 -0.6477336
     SBP      DBP     A1      A2       S1       S2 aSNP rs6040343
1   3.39   4.1800   1.300   1.630   1.9000  -0.0613    1         1
2  -3.49  -2.7200  -0.784  -3.550  -2.8900  -2.1300    0         0
3  -3.40   0.0815  -1.820  -4.390  -1.6700  -3.0900    1         0
4  -7.04  -3.6500  -0.183  -4.740  -2.9800  -2.3500    0         0
5   4.60   4.9900   2.440  -0.117  -0.0408  -0.4350    1         1

phi object contains  75  matrices:
First matrix:
$`1`
    1   2   3   4   5   6   7
1 1.0 0.0 0.5 0.5 0.5 0.5 0.5
2 0.0 1.0 0.5 0.5 0.5 0.5 0.5
3 0.5 0.5 1.0 0.5 0.5 0.5 0.5
4 0.5 0.5 0.5 1.0 0.5 0.5 0.5
5 0.5 0.5 0.5 0.5 1.0 0.5 0.5
6 0.5 0.5 0.5 0.5 0.5 1.0 0.5
7 0.5 0.5 0.5 0.5 0.5 0.5 1.0

Empty IBD object.
```

3.3   Run analysis

Now, run the analysis by the function call *strum()* with two previously con-
structed objects as the arguments.

```
> mySemResult = strum(mySemModel, mySemData)

 Start STRUM analysis ...
    Fitting model step 1 ......................... Done
    Fitting model step 2 ......................... Done
    Testing model fit ............................ Done

Analysis completed!
```

3.4   Result

The result object again contains the model description and result tables. To test
the SNP effect to stress, you would test H_0: $\gamma = 0$ versus H_1: $\gamma \neq 0$. In this
model, γ is the parameter `stress` $\sim$ `rs6040343`, which equals to 1.013427436
with the pvalue = 2.000891e-12.

```
> mySemResult

=========
  Model
=========

Basic properties of the model:
        Model Class ............. strumFittedModel
        Ascertainment ..................... FALSE

List of all variables:
            Obs Covariate InEita   InY Exogen.
bp        FALSE    FALSE    TRUE FALSE   FALSE
anger     FALSE    FALSE    TRUE FALSE    TRUE
stress    FALSE    FALSE    TRUE FALSE   FALSE
rs6040343  TRUE     TRUE   FALSE FALSE      NA
SBP        TRUE    FALSE   FALSE  TRUE      NA
DBP        TRUE    FALSE   FALSE  TRUE      NA
A1         TRUE    FALSE   FALSE  TRUE      NA
A2         TRUE    FALSE   FALSE  TRUE      NA
S1         TRUE    FALSE   FALSE  TRUE      NA
S2         TRUE    FALSE   FALSE  TRUE      NA

Model formulas:
  bp =~ SBP + DBP
  anger =~ A1 + A2
  stress =~ S1 + S2
  bp ~ anger + stress
  stress ~ anger + rs6040343
  var(stress) = .1


==========
  Result
==========

Parameter estimates:
                     estimate    stdError      lowerCI   upperCI          pValue
bp=~DBP            1.034984664  0.04257819   0.95153295 1.1184364 1.618860e-130
anger=~A2          1.003405013  0.08030019   0.84601953 1.1607905  7.882569e-36
stress=~S2         1.054993930  0.08276525   0.89277702 1.2172108  3.247140e-37
bp~anger           0.823535846  0.19648258   0.43843707 1.2086346  2.772460e-05
stress~anger       0.942734335  0.07702719   0.79176382 1.0937049  1.924329e-34
bp~stress          1.052784855  0.16354335   0.73224578 1.3733239  1.215819e-10
stress~rs6040343   1.013517766  0.14609692   0.72717307 1.2998625  3.996716e-12
SBP~[intercept]   -0.117910157  0.30223817  -0.71028608 0.4744658  6.964454e-01
```

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```
DBP~[intercept]    -0.302734202 0.30931966 -0.90898960 0.3035212  3.277234e-01
A1~[intercept]      0.018375775 0.22433007 -0.42130308 0.4580546  9.347151e-01
A2~[intercept]     -0.030596804 0.21819352 -0.45824825 0.3970546  8.884800e-01
S1~[intercept]      0.118914921 0.21176180 -0.29613059 0.5339604  5.744224e-01
S2~[intercept]      0.009989619 0.19519661 -0.37258870 0.3925679  9.591843e-01
SBP~~SBP<p>         0.476876861 0.51400045 -0.53054551 1.4842992  1.767621e-01
DBP~~DBP<p>         1.345964880 0.60228069  0.16551643 2.5264133  1.271590e-02
A1~~A1<p>           1.133270488 0.32775409  0.49088427 1.7756567  2.724188e-04
A2~~A2<p>           0.807378677 0.38682325  0.04921904 1.5655383  1.843489e-02
S1~~S1<p>           1.097072908 0.39554259  0.32182368 1.8723221  2.772076e-03
S2~~S2<p>           0.680992694 0.35439049 -0.01359989 1.3755853  2.732878e-02
SBP~~SBP<e>         1.259598679 0.30646269  0.65894285 1.8602545  1.977261e-05
DBP~~DBP<e>         0.624480268 0.36310715 -0.08719666 1.3361572  4.273225e-02
A1~~A1<e>           0.702416998 0.20541629  0.29980847 1.1050255  3.137041e-04
A2~~A2<e>           1.224986062 0.27512020  0.68576039 1.7642117  4.242850e-06
S1~~S1<e>           1.273526304 0.25231863  0.77899087 1.7680617  2.240559e-07
S2~~S2<e>           1.234374633 0.25196340  0.74053544 1.7282138  4.815705e-07
bp~~bp<p>           0.587268608 0.60016699 -0.58903707 1.7635743  1.639114e-01
anger~~anger<p>     1.154519946 0.25287980  0.65888464 1.6501553  2.491657e-06
bp~~bp<e>           1.078159808 0.40991675  0.27473774 1.8815819  4.266831e-03
anger~~anger<e>     0.886105506 0.19373164  0.50639847 1.2658125  2.393857e-06


Chi-square fit statistics:
                          kappa    chiStat df    pValue
Un-adjusted           1.0000000   7.315514 25 0.9997751
Mean adjusted         0.5440466  13.446483 25 0.9705453
Mean-Variance adjusted  0.4857559  15.060060 28 0.9777947
Theoretically corrected      NA  17.950139 25 0.8445000


Model fit indices:
                            value
Comparative Fit Index (CFI)     1
```

4   SessionInfo

```
> sessionInfo();

R version 3.1.2 (2014-10-31)
Platform: x86_64-w64-mingw32/x64 (64-bit)

locale:
[1] LC_COLLATE=English_United States.1252
[2] LC_CTYPE=English_United States.1252
[3] LC_MONETARY=English_United States.1252
[4] LC_NUMERIC=C
[5] LC_TIME=English_United States.1252

attached base packages:
[1] grid       stats     graphics  grDevices utils     datasets  methods
[8] base

other attached packages:
[1] strum_0.6       Rgraphviz_2.10.0 graph_1.44.1     pedigree_1.4
[5] reshape_0.8.5   HaploSim_1.8.4   Matrix_1.1-4

loaded via a namespace (and not attached):
[1] BiocGenerics_0.12.1 lattice_0.20-29    MASS_7.3-35
[4] parallel_3.1.2      plyr_1.8.1         Rcpp_0.11.3
[7] stats4_3.1.2        tools_3.1.2
```

References

Morris, N.J., Elston, R.C., & Stein, C.M. (2010). A framework for structural equation models in general pedigrees. *Human heredity*, 70, 278–286.