

IVAS : Identification of genetic Variants affecting Alternative Splicing

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Contents

1	Introduction	2
2	The input data set	2
2.1	The genotype data	2
2.2	The expression data	2
2.3	The SNP marker position data	3
2.4	The transcripts model data	3
3	The example dataset : data from Geuvadis RNA sequencing project of 1000 Genome samples	3
4	Loading data	3
5	The <code>ASdb</code> object	4
5.1	Searching alternatively spliced exons based on a reference transcript model.	4
5.2	Estimating expression ratio of AS exons with a data set including FPKM values of transcripts.	6
5.3	Finding SQTls	7
6	Identification of SQTls using multiple cores	8
7	Visualizing the result	8
8	Session Information	9

1 Introduction

Alternative splicing controls relative expression ratios of mature mRNA isoforms from a single gene. Mapping studies of Splicing Quantitative Trait Loci (SQTL), a genetic variant affecting the alternative splicing, are important steps to understand gene regulations and protein activity [1]. We present an effective and user-friendly computational tool to detect SQTLs using transcript expression data from RNA-seq and genotype data, both measured on the same sample. As RNA sequencing (RNA-seq) provides insight into relatively precise measurements of expression level of transcript isoforms from a gene, it is a useful tool to analyze complicated biological phenomenon of RNA transcripts including the alternative splicing [2]. The mapping analysis uses two statistical models : Linear regression model [3] and/or Generalized linear mixed model [5].

2 The input data set

The next subsection introduces the input data. To run this tool, two experimental data sets (an expression data frame from RNA-seq and a genotype data frame) are required. Moreover, we also need a data frame for positions of SNP markers and GTF file for transcript models. As any other genome-wide analyses, it is recommended to use as many samples as possible, usually of population scale, in order to guarantee a statistically significant result.

2.1 The genotype data

The genotype data should be prepared as a simple matrix data. Each column represents an individual and its name should match that of the expression matrix described below (2.2)

	ind1	ind2	ind3	ind4
SNP1	AA	AA	AT	TT
SNP2	CG	CC	GG	CG
SNP3	TT	TT	AT	TT

2.2 The expression data

The expression matrix must comprise expression values of transcripts from RNA-seq. We may obtain them by using alignment tools such as cufflinks. Each column represents an individual and its name should match that of the genotype matrix described above (2.1)

IVAS : Identification of genetic Variants affecting Alternative Splicing

	ind1	ind2	ind3	ind4
transcript1	10.5	15.4	6.7	12.4
transcript2	6.4	7.2	4.5	9.2
transcript3	15.4	14.5	13.2	17.8

2.3 The SNP marker position data

To search SNPs affecting alternative splicing, a data frame comprising genomic location of each SNP is required. It consists of following columns: SNP (SNP marker name), CHR(chromosome number), and locus(SNP position).

SNP	CHR	locus
SNP1	1	4964005
SNP2	1	23513047

2.4 The transcripts model data

We need a reference GTF (General Feature Format) file including information about gene structures such as the positions of exons, introns, and transcripts of genes. The GTF file must be `TxDb` object from the *GenomicFeatures* package [4].

3 The example dataset : data from Geuvadis RNA sequencing project of 1000 Genome samples

This example uses filtered data from an origin data generated by Geuvadis RNA sequencing project, available at <http://www.geuvadis.org/web/geuvadis/RNAseq-project> [6]. The example expression data includes transcripts of 11 randomly selected genes. The genotype data comprises SNPs in those genes.

4 Loading data

For this analysis, you need to load the *IVAS* package, SNP data, expression data, SNP position data, and `TxDb` object from GTF.

Loading *IVAS* package :

```
> library(IVAS)
```

Loading expression data :

```
> data(sampleexp)
```

IVAS : Identification of genetic Variants affecting Alternative Splicing

Loading SNP data :

```
> data(samplesnp)
```

Loading SNP position data :

```
> data(samplesnplocus)
```

Loading TxDb object :

```
> sampleDB <- system.file("extdata","sampleDB", package="IVAS")  
> sample.Txdb <- loadDb(sampleDB)
```

If you want to create the TxDb object from a GTF file, you need to use the `makeTxDbFromGFF` function in the `GenomicFeatures` package.

5 The ASdb object

The ASdb object is a `s4` type class object, and the object is used by the `IVAS` package to store the results from functions in this `IVAS` package. The functions of `IVAS` will save their results by adding a slot. Each slot contains a list object consisting of three elements named as "ES", "ASS", and "IR" for each alternatively splicing pattern type (i.e. ES, ASS, and IR means exon skipping, alternative splice site, and intron retention, respectively).

5.1 Searching alternatively spliced exons based on a reference transcript model.

The `Splicingfinder` function tabulates patterns of alternatively spliced exons. This needs the TxDb object from `makeTxDbFromGFF` by reading a reference GTF file for reference transcript models. The `Splicingfinder` function categorizes alternatively spliced exons into four types of AS patterns (i.e. exon skipping, alternative 3-prime splice site, alternative 5-prime splice site, and intron retention). The result will be saved in the "SplicingModel" slot of ASdb.

To use this function :

```
> ASdb <- Splicingfinder(GTFdb=sample.Txdb,calGene=NULL,Ncor=1,out.dir=NULL)  
  
[1] "-----Processing : chr 2 -----"  
[1] "-----Processing : chr 3 -----"  
[1] "-----Processing : chr 6 -----"  
[1] "-----Processing : chr 8 -----"  
[1] "-----Processing : chr 9 -----"  
[1] "-----Processing : chr 11 -----"
```

IVAS : Identification of genetic Variants affecting Alternative Splicing

```

[1] "-----Processing : chr 17 -----"
[1] "-----Processing : chr 19 -----"

> ASdb

Splicing Models : ES = 182 Rows & ASS = 11 Rows & IR = 2 Rows
#ASdb object with SplicingModel

> head(slot(ASdb,"SplicingModel")$"ASS")

  Index  EnsID          Nchr Strand ShortEX
1 "ASS1" "ENSG00000186001" "3"  "+"   "197562545-197562609"
2 "ASS2" "ENSG00000183826" "6"  "-"   "38565686-38565833"
3 "ASS3" "ENSG00000183826" "6"  "-"   "38565686-38565833"
4 "ASS4" "ENSG00000172728" "8"  "-"   "33319006-33319245"
5 "ASS5" "ENSG00000172728" "8"  "-"   "33318930-33319243"
6 "ASS6" "ENSG00000166263" "17" "+"   "53076993-53077203"
  LongEX          ShortNeighborEX      LongNeighborEX
1 "197562545-197562693" "197566192-197566268" "197566192-197566268"
2 "38565686-38565897" "38607576-38607924" "38607576-38607700"
3 "38565686-38565897" "38607576-38607924" "38580610-38580809"
4 "33318890-33319243" "33310734-33311028" "33310734-33311028"
5 "33318890-33319243" "33310734-33311028" "33310734-33311028"
6 "53076987-53077203" "53076706-53076812" "53076706-53076812"
  Short_des          Long_des          ShortNeighbor_des
1 "197562545-197562609" "197562545-197562693" "197566192-197566268"
2 "38565686-38565833" "38565686-38565897" "38607576-38607924"
3 "38565686-38565833" "38565686-38565897" "38607576-38607924"
4 "33319006-33319245" "33318890-33319243" "33310734-33311028"
5 "33318930-33319243" "33318890-33319243" "33310734-33311028"
6 "53076993-53077203" "53076987-53077203" "53076706-53076812"
  LongNeighbor_des          Types
1 "197566192-197566268" "A5SS"
2 "38580610-38580809,38607576-38607700" "A5SS"
3 "38580610-38580809,38607576-38607700" "A5SS"
4 "33310734-33311028" "A3SS"
5 "33310734-33311028" "A3SS"
6 "53076706-53076812" "A3SS"

```

You are able to define only a single gene if the single gene is inputted. The first column, named by "Index", is a generally used as an identifier and commonly used in other functions of IVAS.

IVAS : Identification of genetic Variants affecting Alternative Splicing

5.2 Estimating expression ratio of AS exons with a data set including FPKM values of transcripts

The `RatioFromFPKM` function calculates expression ratio between transcripts with and without alternatively spliced exons. First, `RatioFromFPKM` divides the isoforms from a single gene into two groups: transcripts with and without an alternatively spliced exon. Then, the ratio of the group totals of transcript FPKM values is calculated. The `RatioFromFPKM` requires expression data set of transcript FPKM values and `ASdb` with the "SplicingModel" slot. The result will be saved in the "Ratio" slot of `ASdb`

```
> ASdb <- RatioFromFPKM(GTFdb=sample.Txdb,ASdb=ASdb,Total.expdata=sampleexp,  
+ CalIndex="ASS7",Ncor=1,out.dir=NULL)  
> ASdb
```

```
Splicing Models : ES = 182 Rows & ASS = 11 Rows & IR = 2 Rows  
Ratio : ES = 0 Rows by 0 samples & ASS = 1 Rows by 78 samples & IR = 0 Rows by 0 samples  
#ASdb object with SplicingModel & Ratio
```

```
> head(slot(ASdb,"Ratio")$"ASS")
```

	Index	EnsID	Nchr	Strand	ShortEX	LongEX
1	"ASS7"	"ENSG00000170889"	"19"	"+"	"54704610-54704756"	"54704740-54704829"
		ShortNeighborEX			LongNeighborEX	Short_TX
1	"54705028-54705149"	"54705028-54705149"			"ENST00000302907 ENST00000391751"	
		Long_TX			Types	NA06984
1	"ENST00000391752 ENST00000402367"	"A5SS"			"0.0370610175563808"	
		NA06986			NA06989	NA06994
1	"0.0754673755080699"	"0.431995041306961"			"0.352248098956179"	
		NA07037			NA07048	NA07051
						NA07056
1	"0.615508066951179"	"0.2535297934717"			"0.396359920018477"	"0.229019337579839"
		NA07347			NA07357	NA10847
1	"0.147021679772774"	"0.294091318766693"			"0.0835188716083212"	
		NA10851			NA11829	NA11830
1	"0.030954840680335"	"0.0174246902189581"			"0.030532429762246"	
		NA11831			NA11843	NA11892
						NA11893
1	"0.15728432880497"	"0.215269984759597"			"0.136324142619792"	"0.38436403201718"
		NA11894			NA11920	NA11930
1	"0.212453507751045"	"0.0333217262293684"			"0.0681235360867984"	
		NA11931			NA11992	NA11993
1	"0.0248643687301508"	"0.245183066925427"			"0.276368360584032"	
		NA11994			NA11995	NA12004
1	"0.114340505887804"	"0.0688073456257966"			"0.0218694795539099"	
		NA12006			NA12043	NA12044
1	"0.719903020532971"	"0.0253480818915533"			"0.0691011133998759"	
		NA12045			NA12058	NA12144
1	"0.269579049697507"	"0.35466877311412"			"0.495392792194793"	
		NA12154			NA12155	NA12249

IVAS : Identification of genetic Variants affecting Alternative Splicing

```
1 "0.0353058516847381" "0.0356549500182518" "0.332527122764556"
  NA12272          NA12273          NA12275
1 "0.547392066663861" "0.0461822327489977" "0.134086715517285"
  NA12282          NA12283          NA12286
1 "0.584161781407799" "0.0376982756006002" "0.242375101101388"
  NA12287          NA12340          NA12341
1 "0.298063167714008" "0.344188773640231" "0.0398630023057284"
  NA12342          NA12347          NA12348
1 "0.189423547621436" "0.304069583466665" "0.0175489426208136"
  NA12383          NA12399          NA12400
1 "0.0285158376928488" "0.0208852172856115" "0.162129766403219"
  NA12413          NA12489          NA12546
1 "0.272617060489197" "0.102893668902972" "0.0480414433339426"
  NA12716          NA12717          NA12718
1 "0.38349497995995" "0.180658216895047" "0.269490808164129"
  NA12749          NA12750          NA12751
1 "0.0967290881103072" "0.226219189907337" "0.024698616842959"
  NA12761          NA12763          NA12775
1 "0.167469444660537" "0.0288808382631836" "0.475715991162855"
  NA12777          NA12778          NA12812
1 "0.281683152810493" "0.00965356629010742" "0.0351200909069428"
  NA12814          NA12815          NA12827
1 "0.583521251701744" "0.500254757772165" "0.313010002548281"
  NA12829          NA12830          NA12842
1 "0.035631624411982" "0.0102342527143945" "0.684792301681123"
  NA12843          NA12872          NA12873
1 "0.251585485678078" "0.220474614626964" "0.353762604584194"
  NA12874          NA12889          NA12890
1 "0.0362672467194004" "0.222764959132477" "0.27775528737905"
```

In this example, we will estimate ratio in the "ASS7" index among splicing models in ASdb.

5.3 Finding SQTls

Using "SplicingModel" and "Ratio" slots in ASdb from `Splicingfinder` and `RatioFromPKM`, respectively, the `sQTlsFinder` function can identify significant SNPs associated with alternative splicing rate (ratio). The result will be saved in the "sQTls" slot of ASdb

```
> ASdb <- sQTlsFinder(ASdb=ASdb, Total.snpdata=samplesnp,
+   Total.snplocus=samplesnplocus, method="lm", Ncor=1)
> ASdb
```

IVAS : Identification of genetic Variants affecting Alternative Splicing

```
Splicing Models : ES = 182 Rows & ASS = 11 Rows & IR = 2 Rows
Ratio : ES = 0 Rows by 0 samples & ASS = 1 Rows by 78 samples & IR = 0 Rows by 0 samples
sQTLs : ES = 0 Rows & ASS = 1 Rows & IR = 0 Rows
#ASdb object with SplicingModel & Ratio & sQTLs
> head(slot(ASdb,"sQTLs")$"ASS")

      SNP      Index  EnsID      Strand Nchr Types
[1,] "rs3810232" "ASS7" "ENSG00000170889" "+"    "19" "A5SS"
      ShortEX      LongEX      ShortNeighborEX
[1,] "54704610-54704756" "54704740-54704829" "54705028-54705149"
      LongNeighborEX      pByGeno      FdrByGeno      diff
[1,] "54705028-54705149" "3.98508717225347e-13" "3.98508717225347e-13" "diff"
      met
[1,] "lm"
```

In this example, we will run the function with the linear regression model. `sQTLs Finder` shows chromosome numbers during mapping analysis.

6 Identification of SQTLs using multiple cores

`Splicingfinder`, `RatioFromFPKM`, and `sQTLsFinder` functions provide to use multi-thread through `foreach` function. The last argument "Ncor" of the functions denotes the number of threads.

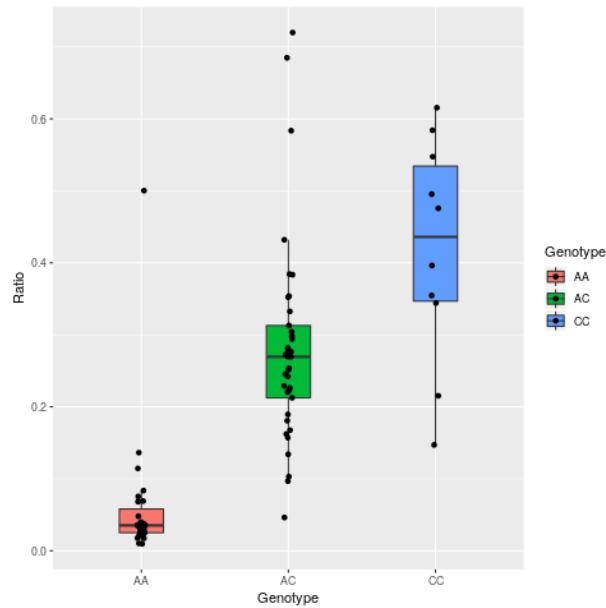
```
> ASdb <- Splicingfinder(GTFdb=sample.Txdb,calGene=NULL,Ncor=4)
> ASdb <- RatioFromFPKM(GTFdb=sample.Txdb,ASdb=ASdb,Total.expdata=sampleexp,Ncor=4)
> ASdb <- sQTLsFinder(ASdb=ASdb,Total.snpdata=samplesnp,
+   Total.snplocus=samplesnplocus,method="lm",Ncor = 4)
> ASdb
```

7 Visualizing the result

To visualize the results into boxplot, the `IVAS` package provides the `saveBplot` function. Using the data frame from the output of `sQTLsFinder` function, `saveBplot` can make the boxplot.

```
> saveBplot(ASdb=ASdb,Total.snpdata=samplesnp,Total.snplocus=samplesnplocus,
+   CalIndex="ASS7",out.dir="./result")
```


IVAS : Identification of genetic Variants affecting Alternative Splicing



The output png files are saved in "result" folder.

8 Session Information

```
R version 3.6.0 (2019-04-26)
Platform: x86_64-pc-linux-gnu (64-bit)
Running under: Ubuntu 18.04.2 LTS

Matrix products: default
BLAS: /home/biocbuild/bbs-3.9-bioc/R/lib/libRblas.so
LAPACK: /home/biocbuild/bbs-3.9-bioc/R/lib/libRlapack.so

locale:
 [1] LC_CTYPE=en_US.UTF-8      LC_NUMERIC=C
 [3] LC_TIME=en_US.UTF-8      LC_COLLATE=C
 [5] LC_MONETARY=en_US.UTF-8  LC_MESSAGES=en_US.UTF-8
 [7] LC_PAPER=en_US.UTF-8     LC_NAME=C
 [9] LC_ADDRESS=C             LC_TELEPHONE=C
[11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C

attached base packages:
[1] stats4    parallel  stats     graphics  grDevices  utils      datasets
[8] methods  base

other attached packages:
```

IVAS : Identification of genetic Variants affecting Alternative Splicing

```
[1] IVAS_2.4.0          ggplot2_3.1.1      GenomicFeatures_1.36.0
[4] AnnotationDbi_1.46.0 Biobase_2.44.0     GenomicRanges_1.36.0
[7] GenomeInfoDb_1.20.0 IRanges_2.18.0     S4Vectors_0.22.0
[10] BiocGenerics_0.30.0
```

loaded via a namespace (and not attached):

```
[1] httr_1.4.0          tidyr_0.8.3
[3] bit64_0.9-7         splines_3.6.0
[5] foreach_1.4.4       assertthat_0.2.1
[7] BiocManager_1.30.4 blob_1.1.1
[9] GenomeInfoDbData_1.2.1 Rsamtools_2.0.0
[11] yaml_2.2.0          progress_1.2.0
[13] pillar_1.3.1        RSQLite_2.1.1
[15] lattice_0.20-38     glue_1.3.1
[17] digest_0.6.18       XVector_0.24.0
[19] minqa_1.2.4         colorspace_1.4-1
[21] htmltools_0.3.6     Matrix_1.2-17
[23] plyr_1.8.4          XML_3.98-1.19
[25] pkgconfig_2.0.2     biomaRt_2.40.0
[27] zlibbioc_1.30.0     purrr_0.3.2
[29] scales_1.0.0        BiocParallel_1.18.0
[31] lme4_1.1-21         tibble_2.1.1
[33] withr_2.1.2         SummarizedExperiment_1.14.0
[35] lazyeval_0.2.2      magrittr_1.5
[37] crayon_1.3.4        memoise_1.1.0
[39] evaluate_0.13       nlme_3.1-139
[41] MASS_7.3-51.4       doParallel_1.0.14
[43] tools_3.6.0         prettyunits_1.0.2
[45] hms_0.4.2           BiocStyle_2.12.0
[47] matrixStats_0.54.0 stringr_1.4.0
[49] munsell_0.5.0       DelayedArray_0.10.0
[51] Biostrings_2.52.0   compiler_3.6.0
[53] rlang_0.3.4         nloptr_1.2.1
[55] grid_3.6.0          RCurl_1.95-4.12
[57] iterators_1.0.10    labeling_0.3
[59] bitops_1.0-6        rmarkdown_1.12
[61] boot_1.3-22         gtable_0.3.0
[63] codetools_0.2-16   DBI_1.0.0
[65] R6_2.4.0            GenomicAlignments_1.20.0
[67] gridExtra_2.3       knitr_1.22
[69] dplyr_0.8.0.1       rtracklayer_1.44.0
[71] bit_1.1-14          ggfortify_0.4.6
[73] stringi_1.4.3       Rcpp_1.0.1
[75] tidyselect_0.2.5    xfun_0.6
```

References

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