

# VariantFiltering: filtering of coding and non-coding genetic variants

**Dei M. Elurbe<sup>1,2</sup> and Robert Castelo<sup>3,4</sup>**

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<sup>1</sup> CIBER de Enfermedades Raras (CIBERER), Barcelona, Spain

<sup>2</sup> Present address: CMBI, Radboud University Medical Centre, Nijmegen, The Netherlands.

<sup>3</sup> Department of Experimental and Health Sciences, Universitat Pompeu Fabra, Barcelona, Spain.

<sup>4</sup> Research Program on Biomedical Informatics (GRIB), Hospital del Mar Medical Research Institute, Barcelona, Spain.

## 1 Overview

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The aim of this software package is to facilitate the filtering and annotation of coding and non-coding genetic variants from a group of unrelated individuals, or a family of related ones among which at least one of them is affected by a genetic disorder. When working with related individuals, [VariantFiltering](#) can search for variants from the affected individuals that segregate according to a particular inheritance model acting on autosomes (dominant, recessive homozygous or recessive heterozygous -also known as compound heterozygous) or on allosomes (X-linked), or that occur *de novo*. When working with unrelated individuals, no mode of inheritance is used for filtering but it can be used to search for variants shared among individuals affected by a common genetic disorder.

The main input is a multisample Variant Call Format (VCF) file which is parsed using the R/Bioconductor infrastructure, and particularly the functionality from the [VariantAnnotation](#) package for that purpose. This infrastructure also allows [VariantFiltering](#) to stream over large VCF files to reduce the memory footprint and to annotate the input variants with diverse functional information.

A core set of functional data are annotated by [VariantFiltering](#) but this set can be modified and extended using Bioconductor annotation packages such as [MafDb.1Kgenomes.phase3.hs37d5](#), which stores and exposes to the user minor allele frequency (MAF) values frozen from the latest release of the 1000 Genomes project.

The package contains a toy data set to illustrate how it works through this vignette, and it consists of a multisample VCF file with variants from chromosomes 20, 21, 22 and allosomes X, Y from a trio of CEU individuals of the 1000 Genomes project. To further reduce the execution time of this vignette, only the code for the first analysis is actually evaluated and its results reported.

## 2 Setting up the analysis

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To start using [VariantFiltering](#) the user should consider installing the packages listed in the `Suggests:` field from its `DESCRIPTION` file. After loading [VariantFiltering](#) the first step is to build a parameter object, of class `VariantFilteringParam` which requires at least a character string with the input VCF filename, as follows:

## VariantFiltering: filtering of coding and non-coding genetic variants

```
> library(VariantFiltering)
> CEUvcf <- file.path(system.file("extdata", package="VariantFiltering"),
+                       "CEUtrio.vcf.bgz")
> vfpar <- VariantFilteringParam(CEUvcf)
> class(vfpar)

[1] "VariantFilteringParam"
attr(,"package")
[1] "VariantFiltering"

> vfpar

VariantFiltering parameter object

VCF file(s): CEUtrio.vcf.bgz
Genome version(s): hg19(NCBI)
Number of individuals: 3 (NA12878, NA12891, NA12892)
Genome-centric annotation package: BSgenome.Hsapiens.1000genomes.hs37d5 (1000genomes hs37d5)
Variant-centric annotation package: SNPlocs.Hsapiens.dbSNP144.GRCh37 (dbSNP Human BUILD 144)
Transcript-centric annotation package: TxDb.Hsapiens.UCSC.hg19.knownGene
Gene-centric annotation package: org.Hs.eg.db
Radical/Conservative AA changes: AA_chemical_properties_HanadaGojoboriLi2006.tsv
Codon usage table: humanCodonUsage.txt
Regions to annotate: CodingVariants, IntronVariants, FiveSpliceSiteVariants, ThreeSpliceSiteVariants, PromoterVariants
Other annotation pkg/obj: MafDb.1Kgenomes.phase1.hs37d5,
                        PolyPhen.Hsapiens.dbSNP131,
                        SIFT.Hsapiens.dbSNP137,
                        phastCons100way.UCSC.hg19,
                        humanGenesPhylostrata

All transcripts: FALSE
```

The display of the *VariantFilteringParam* object indicates a number of default values which can be overridden when calling the construction function. To quickly see all the available arguments and their default values we should type:

```
> args(VariantFilteringParam)

function (vcfFilename, pedFilename = NA_character_, bsgenome = "BSgenome.Hsapiens.1000genomes.hs37d5",
  orgdb = "org.Hs.eg.db", txdb = "TxDb.Hsapiens.UCSC.hg19.knownGene",
  snpdb = "SNPlocs.Hsapiens.dbSNP144.GRCh37", weightMatricesFileNames = NA,
  weightMatricesLocations = rep(list(variantLocations()), length(weightMatricesFileNames)),
  weightMatricesStrictLocations = rep(list(FALSE), length(weightMatricesFileNames)),
  radicalAAchangeFilename = file.path(system.file("extdata",
    package = "VariantFiltering"), "AA_chemical_properties_HanadaGojoboriLi2006.tsv"),
  codonusageFilename = file.path(system.file("extdata", package = "VariantFiltering"),
    "humanCodonUsage.txt"), geneticCode = getGeneticCode("SGC0"),
  allTranscripts = FALSE, regionAnnotations = list(CodingVariants(),
    IntronVariants(), FiveSpliceSiteVariants(), ThreeSpliceSiteVariants(),
    PromoterVariants(), FiveUTRVariants(), ThreeUTRVariants()),
  intergenic = FALSE, otherAnnotations = c("MafDb.1Kgenomes.phase1.hs37d5",
    "PolyPhen.Hsapiens.dbSNP131", "SIFT.Hsapiens.dbSNP137",
    "phastCons100way.UCSC.hg19", "humanGenesPhylostrata"),
  geneKeytype = NA_character_, yieldSize = NA_integer_)
```

## VariantFiltering: filtering of coding and non-coding genetic variants

NULL

The manual page of `VariantFilteringParam` contains more information about these arguments and their default values.

### 3 Annotating variants

After setting up the parameters object, the next step is to annotate variants. This can be done using upfront an inheritance model that will substantially filter and reduce the number of variants and annotations or, as we illustrate here below, calling the function `unrelatedIndividuals()` that just annotates the variants without filtering out any of them:

```
> uind <- unrelatedIndividuals(vfpar)
```

```
> class(uind)
```

```
[1] "VariantFilteringResults"
```

```
attr(,"package")
```

```
[1] "VariantFiltering"
```

```
> uind
```

VariantFiltering results object

Genome version(s): "hg19" (NCBI)

Number of individuals: 3 (NA12878, NA12891, NA12892)

Variants segregate according to a(n) unrelated individuals inheritance model

Quality filters

PASS	INDELQual	LowQual	SNPQual
TRUE	TRUE	TRUE	TRUE

Functional annotation filters

variantType	dbSNP	location	consequence
FALSE	FALSE	FALSE	FALSE
codonusageFC	OMIM	aaChangeType	maxMAF
FALSE	FALSE	FALSE	FALSE
genePhyloStratum	S0terms		
FALSE	FALSE		

The resulting object belongs to the *VariantFilteringResults* class of objects, defined within *VariantFiltering*, whose purpose is to ease the task of filtering and prioritizing the annotated variants. The display of the object just tells us the genome version of the input VCF file, the number of individuals, the inheritance model and what variant filters are activated.

To get a summary of the number of variants annotated to a particular feature we should use the function `summary()`:

```
> summary(uind)
```

	S0ID	Description	Nr. Variants	% Variants
1	S0:0002012	start_lost	1	0.23
2	S0:0001631	upstream_gene_variant	116	26.54
3	S0:0001624	3_prime_UTR_variant	23	5.26
4	S0:0001623	5_prime_UTR_variant	11	2.52

## VariantFiltering: filtering of coding and non-coding genetic variants

5	SO:0001589	frameshift_variant	2	0.46
6	SO:0001587	stop_gained	1	0.23
7	SO:0001583	missense_variant	18	4.12
8	SO:0001575	splice_donor_variant	4	0.92
9	SO:0001574	splice_acceptor_variant	8	1.83
10	SO:0001627	intron_variant	361	82.61
11	SO:0001819	synonymous_variant	17	3.89

The default setting of the `summary()` function is to provide feature annotations in terms of Sequence Ontology (SO) terms. The reported number of variants refer to the number of different variants in the input VCF file annotated to the feature while the percentage of variants refers to the fraction of this number over the total number of different variants in the input VCF file.

We can also obtain a summary based on the Bioconductor feature annotations provided by the functions `locateVariants()` and `predictCoding()` from the [VariantAnnotation](#) package, as follows:

```
> summary(uind, method="bioc")
```

	BIOCID	Nr. Variants	% Variants
2	intron	361	82.61
3	fiveUTR	11	2.52
4	threeUTR	23	5.26
5	coding	37	8.47
7	promoter	116	26.54
8	fiveSpliceSite	4	0.92
9	threeSpliceSite	8	1.83
10	frameshift	2	0.46
11	nonsense	1	0.23
12	nonsynonymous	18	4.12
13	synonymous	17	3.89

Since SO terms are organized hierarchically, we can use this structure to aggregate feature annotations into more coarse-grained SO terms using the argument `method="S0full"`:

```
> uindS0 <- summary(uind, method="S0full")
> uindS0
```

	SOID	Level	Description	Nr. Variants	% Variants
1	SO:0002012	9	start_lost	1	0.23
2	SO:0001582	8	initiator_codon_variant	1	0.23
3	SO:0001819	8	synonymous_variant	17	3.89
4	SO:0001631	5	upstream_gene_variant	116	26.54
5	SO:0001628	4	intergenic_variant	116	26.54
6	SO:0001624	8	3_prime_UTR_variant	23	5.26
7	SO:0001623	8	5_prime_UTR_variant	11	2.52
8	SO:0001622	7	UTR_variant	34	7.78
9	SO:0001589	9	frameshift_variant	2	0.46
10	SO:0001587	5	stop_gained	1	0.23
11	SO:0001906	4	feature_truncation	1	0.23
12	SO:0001583	11	missense_variant	18	4.12
13	SO:0001992	10	nonsynonymous_variant	19	4.35

## VariantFiltering: filtering of coding and non-coding genetic variants

14	SO:0001650	9	inframe_variant	19	4.35
15	SO:0001818	8	protein_altering_variant	21	4.81
16	SO:0001580	7	coding_sequence_variant	37	8.47
17	SO:0001968	6	coding_transcript_variant	61	13.96
18	SO:0001791	6	exon_variant	61	13.96
19	SO:0001575	8	splice_donor_variant	4	0.92
20	SO:0001574	8	splice_acceptor_variant	8	1.83
21	SO:0001629	7	splice_site_variant	12	2.75
22	SO:0001627	6	intron_variant	361	82.61
23	SO:0001568	6	splicing_variant	12	2.75
24	SO:0001576	5	transcript_variant	401	91.76
25	SO:0001564	4	gene_variant	401	91.76
26	SO:0001878	3	feature_variant	437	100.00
27	SO:0001537	2	structural_variant	437	100.00
28	SO:0001060	1	sequence_variant	437	100.00
29	SO:0002072	0	sequence_comparison	437	100.00

Here the `Level` column refers to the shortest-path distance to the most general SO term `sequence_variant` within the SO acyclic digraph. We can use this level value to interrogate the annotations on a specific granularity:

```
> uindSO[uindSO$Level == 6, ]
```

	SOID	Level	Description	Nr. Variants	% Variants
17	SO:0001968	6	coding_transcript_variant	61	13.96
18	SO:0001791	6	exon_variant	61	13.96
22	SO:0001627	6	intron_variant	361	82.61
23	SO:0001568	6	splicing_variant	12	2.75

Variants are stored internally in a *VRanges* object. We can retrieve the variants as a *VRanges* object with the function `allVariants()`:

```
> allVariants(uind)
```

```
VRangesList of length 3  
names(3): NA12878 NA12891 NA12892
```

This function in fact returns a *VRangesList* object with one element per sample by default. We can change the grouping of variants with the argument `groupBy` specifying the annotation column we want to use to group variants. If the specified column does not exist, then it will return a single *VRanges* object with all annotated variants.

Using the following code we can obtain a graphical display of a variant, including the aligned reads and the running coverage, to have a visual representation of its support. For this purpose we need to have the BAM files used to perform the variant calling. In this case we are using toy BAM files stored along with the *VariantFiltering* package, which for practical reasons only include a tiny subset of the aligned reads.

```
> path2bams <- file.path(system.file("extdata", package="VariantFiltering"),  
+                          paste0(samples(uind), ".subset.bam"))  
> bv <- BamViews(bamPaths=path2bams,  
+               bamSamples=DataFrame(row.names=samples(uind)))  
> bamFiles(uind) <- bv
```

## VariantFiltering: filtering of coding and non-coding genetic variants

```
> bamFiles(uind)

BamViews dim: 0 ranges x 3 samples
names: NA12878 NA12891 NA12892
detail: use bamPaths(), bamSamples(), bamRanges(), ...

> plot(uind, what="rs6130959", sampleName="NA12892")
```

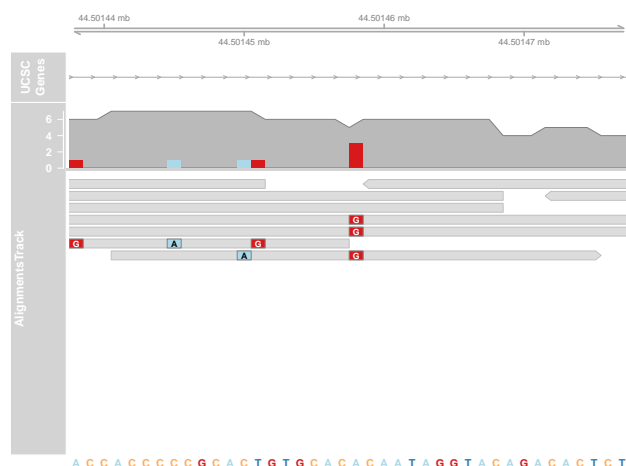


Figure 1: Browser-like display of a variant.

## 4 Filters and cutoffs

In the case of having run the `unrelatedIndividuals()` annotation function we can filter variants by restricting the samples involved in the analysis, as follows:

```
> samples(uind)

[1] "NA12878" "NA12891" "NA12892"

> samples(uind) <- c("NA12891", "NA12892")
> uind

VariantFiltering results object

Genome version(s): "hg19"(NCBI)
Number of individuals: 2 (NA12891, NA12892)
Variants segregate according to a(n) unrelated individuals inheritance model
Quality filters
  PASS INDELQual LowQual SNPQual
TRUE TRUE TRUE TRUE
Functional annotation filters
  variantType dbSNP location consequence
      FALSE FALSE FALSE FALSE
codonusageFC OMIM aaChangeType maxMAF
      FALSE FALSE FALSE FALSE
```

## VariantFiltering: filtering of coding and non-coding genetic variants

```
genePhyloStratum      S0terms
FALSE                 FALSE

> uindS02sam <- summary(uind, method="S0full")
> uindS02sam[uindS02sam$Level == 6, ]
```

	SOID	Level	Description	Nr. Variants	% Variants
17	S0:0001968	6	coding_transcript_variant	54	14.29
18	S0:0001791	6	exon_variant	54	14.29
22	S0:0001627	6	intron_variant	308	81.48
23	S0:0001568	6	splicing_variant	9	2.38

As we can see, restricting the samples for filtering variants results in fewer variants. We can set the original samples back with the function `resetSamples()`:

```
> uind <- resetSamples(uind)
> uind
```

VariantFiltering results object

Genome version(s): "hg19"(NCBI)  
Number of individuals: 3 (NA12878, NA12891, NA12892)  
Variants segregate according to a(n) unrelated individuals inheritance model

Quality filters

PASS	INDELQual	LowQual	SNPQual
TRUE	TRUE	TRUE	TRUE

Functional annotation filters

variantType	dbSNP	location	consequence
FALSE	FALSE	FALSE	FALSE
codonusageFC	OMIM	aaChangeType	maxMAF
FALSE	FALSE	FALSE	FALSE

genePhyloStratum S0terms  
FALSE FALSE

The rest of the filtering operations we can perform on a *VariantFilteringResults* objects are implemented through the *FilterRules* class which implements a general mechanism for generating logical masks to filter vector-like objects; consult its manual page at the [IRanges](#) package for full technical details.

The *Variantfiltering* package provides a number of default filters, which can be extended by the user. To see which are these filters we just have to use the `filters()` function:

```
> filters(uind)
```

FilterRules of length 15  
names(15): PASS INDELQual LowQual SNPQual ... maxMAF genePhyloStratum S0terms

Filters may be active or inactive. Only active filters will participate in the filtering process when we interrogate for variants. To know what filters are active we should use the `active()` function as follows:

```
> active(filters(uind))
```

PASS	INDELQual	LowQual	SNPQual
TRUE	TRUE	TRUE	TRUE

## VariantFiltering: filtering of coding and non-coding genetic variants

variantType	dbSNP	location	consequence
FALSE	FALSE	FALSE	FALSE
codonusageFC	OMIM	aaChangeType	maxMAF
FALSE	FALSE	FALSE	FALSE
maxMAF	genePhyloStratum	S0terms	
FALSE	FALSE	FALSE	

By default, all filters are always inactive. To activate all of them, we can simply type:

```
> active(filters(uind)) <- TRUE
> active(filters(uind))
```

PASS	INDELQual	LowQual	SNPQual
TRUE	TRUE	TRUE	TRUE
variantType	dbSNP	location	consequence
TRUE	TRUE	TRUE	TRUE
codonusageFC	OMIM	aaChangeType	maxMAF
TRUE	TRUE	TRUE	TRUE
maxMAF	genePhyloStratum	S0terms	
TRUE	TRUE	TRUE	

```
> summary(uind)
```

	SOID	Description	Nr. Variants	% Variants
1	S0:0001631	upstream_gene_variant	3	20.0
2	S0:0001624	3_prime_UTR_variant	2	13.3
3	S0:0001583	missense_variant	1	6.7
4	S0:0001575	splice_donor_variant	1	6.7
5	S0:0001627	intron_variant	4	26.7
6	S0:0001819	synonymous_variant	15	100.0

To deactivate all of them back and selectively activate one of them, we should use the bracket `[]` notation, as follows:

```
> active(filters(uind)) <- FALSE
> active(filters(uind))["dbSNP"] <- TRUE
> summary(uind)
```

	SOID	Description	Nr. Variants	% Variants
1	S0:0002012	start_lost	1	0.20
2	S0:0001631	upstream_gene_variant	122	23.87
3	S0:0001624	3_prime_UTR_variant	27	5.28
4	S0:0001623	5_prime_UTR_variant	10	1.96
5	S0:0001587	stop_gained	1	0.20
6	S0:0001583	missense_variant	21	4.11
7	S0:0001575	splice_donor_variant	3	0.59
8	S0:0001574	splice_acceptor_variant	7	1.37
9	S0:0001627	intron_variant	427	83.56
10	S0:0001819	synonymous_variant	18	3.52



## VariantFiltering: filtering of coding and non-coding genetic variants

The previous filter just selects variants with an annotated dbSNP identifier. However, other filters may require cutoff values to decide what variants pass the filter. To set those values we can use the function `cutoffs()`. For instance, in the case of the `S0terms` filter, we should use set the cutoff values to select variants annotated to specific SO terms. Here we select, for instance, those annotated in UTR regions:

```
> change(cutoffs(uind), "S0terms") <- "UTR_variant"
> active(filters(uind))["S0terms"] <- TRUE
> summary(uind)

[1] S0ID      Description  Nr. Variants % Variants
<0 rows> (or 0-length row.names)

> summary(uind, method="S0full")

      S0ID Level      Description Nr. Variants % Variants
1  S0:0001624      8      3_prime_UTR_variant          27          75
2  S0:0001623      8      5_prime_UTR_variant          10          28
3  S0:0001622      7          UTR_variant             36         100
4  S0:0001968      6 coding_transcript_variant          36         100
5  S0:0001791      6          exon_variant             36         100
6  S0:0001576      5      transcript_variant           36         100
7  S0:0001564      4          gene_variant             36         100
8  S0:0001878      3          feature_variant           36         100
9  S0:0001537      2      structural_variant           36         100
10 S0:0001060      1      sequence_variant            36         100
11 S0:0002072      0      sequence_comparison          36         100
```

Note that the first call to `summary()` did not report any variant since there are no variants annotated to the SO term `UTR_variant`. However, when using the argument `method="S0full"`, all variants annotated to more specific SO terms in the hierarchy will be reported.

The methods `filters()` and `cutoffs()` can be employed to extend the filtering functionality. Here we show a simple example in which we add a filter to detect the loss of the codon that initiates translation. This constitutes already a feature annotated by [VariantFiltering](#) so that we can verify that it works:

```
> startLost <- function(x) {
+   mask <- start(allVariants(x, groupBy="nothing")$CDSLOC) == 1 &
+     as.character(allVariants(x, groupBy="nothing")$REFCODON) == "ATG" &
+     as.character(allVariants(x, groupBy="nothing")$VARCODON) != "ATG"
+   mask
+ }
> filters(uind)$startLost <- startLost
> active(filters(uind)) <- FALSE
> active(filters(uind))["startLost"] <- TRUE
> active(filters(uind))

      PASS      INDELQual      LowQual      SNPQual
      FALSE      FALSE      FALSE      FALSE
variantType      dbSNP      location      consequence
      FALSE      FALSE      FALSE      FALSE
codonusageFC      OMIM      aaChangeType      maxMAF
      FALSE      FALSE      FALSE      FALSE
maxMAF genePhyloStratum      S0terms      startLost
```

## VariantFiltering: filtering of coding and non-coding genetic variants

```

FALSE          FALSE          FALSE          TRUE

> summary(uind)

      SOID      Description Nr. Variants % Variants
1 S0:0002012      start_lost           1         100
2 S0:0001631 upstream_gene_variant       1         100
3 S0:0001623  5_prime_UTR_variant       1         100
4 S0:0001583      missense_variant       1         100
5 S0:0001575 splice_donor_variant       1         100
6 S0:0001627      intron_variant       1         100

```

As we can see, our filter works as expected and selects the only variant which was annotated with the SO term `start_lost`. Note that there is also an additional annotation indicating this variant belongs to an UTR region resulting from an alternative CDS.

Properly updating cutoff values may be problematic if we do not know how exactly are they employed by the corresponding filters. To facilitate setting the right cutoff values we use the `change()` method and we illustrate it on the filter of minor allele frequency (MAF) values:

```

> active(filters(uind)) <- FALSE
> active(filters(uind))["maxMAF"] <- TRUE
> cutoffs(uind)$maxMAF

CutoffsList of length 2
names(2): popmask maxvalue

> cutoffs(uind)$maxMAF$popmask

      AFKGp1 AFR_AFKGp1 AMR_AFKGp1 ASN_AFKGp1 EUR_AFKGp1
      TRUE      TRUE      TRUE      TRUE      TRUE

> cutoffs(uind)$maxMAF$maxvalue

[1] 0.5

> change(cutoffs(uind)$maxMAF, "popmask") <- FALSE
> cutoffs(uind)$maxMAF$popmask

      AFKGp1 AFR_AFKGp1 AMR_AFKGp1 ASN_AFKGp1 EUR_AFKGp1
      FALSE      FALSE      FALSE      FALSE      FALSE

> change(cutoffs(uind)$maxMAF, "popmask") <- c(AFKGp1=TRUE)
> cutoffs(uind)$maxMAF$popmask

      AFKGp1 AFR_AFKGp1 AMR_AFKGp1 ASN_AFKGp1 EUR_AFKGp1
      TRUE      FALSE      FALSE      FALSE      FALSE

> change(cutoffs(uind)$maxMAF, "maxvalue") <- 0.01
> summary(uind)

      SOID      Description Nr. Variants % Variants
1 S0:0001631 upstream_gene_variant       55      30.05
2 S0:0001624  3_prime_UTR_variant        6       3.28
3 S0:0001623  5_prime_UTR_variant        3       1.64
4 S0:0001589 frameshift_variant          1       0.55
5 S0:0001587      stop_gained            1       0.55
6 S0:0001583      missense_variant        8       4.37

```

## VariantFiltering: filtering of coding and non-coding genetic variants

7	S0:0001575	splice_donor_variant	2	1.09
8	S0:0001574	splice_acceptor_variant	3	1.64
9	S0:0001627	intron_variant	157	85.79
10	S0:0001819	synonymous_variant	4	2.19

In this case we selected variants with  $MAF < 0.01$  in the global population of the 1000 Genomes project. If we are interested in retrieving the actual set of filtered variants, we can do it using the function `filteredVariants()`:

```
> filteredVariants(uind)
VRangesList of length 3
names(3): NA12878 NA12891 NA12892
```

To further understand how to manipulate *Vranges* and *VRangesList* objects, please consult the package [VariantAnnotation](#).

## 5 Inheritance models

We can filter upfront variants that do not segregate according to a given inheritance model. In such a case, we also need to provide a PED file at the time we build the parameter object, as follows:

```
> CEUped <- file.path(system.file("extdata", package="VariantFiltering"),
+                       "CEUtrio.ped")
> param <- VariantFilteringParam(vcfFileNames=CEUvcf, pedFilename=CEUped)
```

Here we are using a PED file included in the [VariantFiltering](#) package and specifying information about the CEU trio employed in this vignette.

To use an inheritance model we need to replace the previous call to `unrelatedIndividuals()` by one specific to the inheritance model. The [VariantFiltering](#) package offers 5 possible ones:

- **Autosomal recessive inheritance analysis: Homozygous.**

Homozygous variants responsible for a recessive trait in the affected individuals can be identified calling the `autosomalRecessiveHomozygous()` function. This function selects homozygous variants that are present in all the affected individuals and occur in heterozygosity in the unaffected ones.

- **Autosomal recessive inheritance analysis: Heterozygous.**

To filter by this mode of inheritance, also known as compound heterozygous, we need two unaffected parents/ancestors and at least one affected descendant. Variants are filtered in five steps: 1. select heterozygous variants in one of the parents and homozygous in the other; 2. discard previously selected variants that are common between the two parents; 3. group variants by gene; 4. select those genes, and the variants that occur within them, which have two or more variants and there is at least one from each parent; 5. from the previously selected variants, discard those that do not occur in the affected descendants. This is implemented in the function `autosomalRecessiveHeterozygous()`.

- **Autosomal dominant inheritance analysis.**

## VariantFiltering: filtering of coding and non-coding genetic variants

The function `autosomalDominant()` identifies variants present in all the affected individual(s) discarding the ones that also occur in at least one of the unaffected subjects.

- **X-Linked inheritance analysis.**

The function `xLinked()` identifies variants that appear only in the X chromosome of the unaffected females as heterozygous, don't appear in the unaffected males analyzed and finally are present (as homozygous) in the affected male(s). This function is currently restricted to affected males, and therefore, it cannot search for X-linked segregating variants affecting daughters.

- **De Novo variants analysis**

The function `deNovo()` searches for *de novo* variants which are present in one descendant and present in both parents/ancestors. It is currently restricted to a trio of individuals.

## 6 Create a report from the filtered variants

The function `reportVariants()` allows us to easily create a report from the filtered variants into a CSV or a TSV file as follows:

```
> reportVariants(uind, type="csv", file="uind.csv")
```

The default value on the `type` argument ("shiny") starts a shiny web app which allows one to interactively filter the variants, obtaining an updated `VariantFilteringResults` object and downloading the filtered variants and the corresponding full reproducible R code, if necessary.

VarID	GENE	HGVSp	CONSEQUENCE	AAchange	AAchangeType	PolyPhen2	PROVEAN
1 rs11699009	BPIFB4	p.527F>L	nonsynonymous	F527L	Conservative	benign	Neutral
2 rs2839223	PCNT	p.704G>E	nonsynonymous	G704E	Radical	benign	Neutral
3 rs16989291	SF11	p.306W>R	nonsynonymous	W306R	Radical	probably damaging	Deleterious
4 rs79478	PARVG	p.130C>R	nonsynonymous	C130R	Radical	NA	Neutral
5 rs3764880	TLR8	p.1M>V	nonsynonymous	M1V	Conservative	possibly damaging	Neutral

**Figure 2:** Snapshot of the shiny web app run from VariantFiltering with the function `reportVariants()`. Some of the parameters has been filled for illustrative purposes.

## 7 Using the package with parallel execution

Functions in *VariantFiltering* to annotate and filter variants leverage the functionality of the Bioconductor package *BiocParallel* to perform in parallel some of the tasks and calculations and reduce the overall execution time. These functions have an argument called `BPPARAM` that allows the user to control how this parallelism is exploited. In particular the user must give as value to this argument the result from a call to the function `bpparam()`, which actually is its default behavior. Here below we modify that behavior to force a call being executed without parallelism. The interested reader should consult the help page of `bpparam()` and the vignette of the *BiocParallel* for further information.

## 8 Session information

```
> toLatex(sessionInfo())
```

- R version 4.5.1 Patched (2025-08-23 r88802), x86\_64-pc-linux-gnu
- Locale: LC\_CTYPE=en\_US.UTF-8, LC\_NUMERIC=C, LC\_TIME=en\_GB, LC\_COLLATE=C, LC\_MONETARY=en\_US.UTF-8, LC\_MESSAGES=en\_US.UTF-8, LC\_PAPER=en\_US.UTF-8, LC\_NAME=C, LC\_ADDRESS=C, LC\_TELEPHONE=C, LC\_MEASUREMENT=en\_US.UTF-8, LC\_IDENTIFICATION=C
- Time zone: America/New\_York
- TZcode source: system (glibc)
- Running under: Ubuntu 24.04.3 LTS
- Matrix products: default
- BLAS: /home/biocbuild/bbs-3.22-bioc/R/lib/libRblas.so
- LAPACK: /usr/lib/x86\_64-linux-gnu/lapack/liblapack.so.3.12.0
- Base packages: base, datasets, grDevices, graphics, methods, stats, stats4, utils
- Other packages: AnnotationDbi 1.72.0, BSgenome 1.78.0, BSgenome.Hsapiens.1000genomes.hs37d5 0.99.1, Biobase 2.70.0, BiocGenerics 0.56.0, BiocIO 1.20.0, Biostrings 2.78.0, GenomicFeatures 1.62.0, GenomicRanges 1.62.0, GenomicScores 2.22.0, IRanges 2.44.0, MafDb.1Kgenomes.phase1.hs37d5 3.10.0, MatrixGenerics 1.22.0, PolyPhen.Hsapiens.dbSNP131 1.0.2, RSQLite 2.4.3, Rsamtools 2.26.0, S4Vectors 0.48.0, SIFT.Hsapiens.dbSNP137 1.0.0, SNPlocs.Hsapiens.dbSNP144.GRCh37 0.99.20, Seqinfo 1.0.0, SummarizedExperiment 1.40.0, TxDb.Hsapiens.UCSC.hg19.knownGene 3.22.1, VariantAnnotation 1.56.0, VariantFiltering 1.46.0, XVector 0.50.0, generics 0.1.4, matrixStats 1.5.0, org.Hs.eg.db 3.22.0, phastCons100way.UCSC.hg19 3.7.2, rtracklayer 1.70.0
- Loaded via a namespace (and not attached): AnnotationFilter 1.34.0, AnnotationHub 4.0.0, BiocFileCache 3.0.0, BiocManager 1.30.26, BiocParallel 1.44.0, BiocStyle 2.38.0, BiocVersion 3.22.0, DBI 1.2.3, DT 0.34.0, DelayedArray 0.36.0, Formula 1.2-5, GenomeInfoDb 1.46.0, GenomicAlignments 1.46.0, Gviz 1.54.0, HDF5Array 1.38.0, Hmisc 5.2-4, KEGGREST 1.50.0, Matrix 1.7-4, ProtGenerics 1.42.0, R6 2.6.1, RBGL 1.86.0,

## VariantFiltering: filtering of coding and non-coding genetic variants

RColorBrewer 1.1-3, RCurl 1.98-1.17, Rcpp 1.1.0, Rhdf5lib 1.32.0, S4Arrays 1.10.0, S7 0.2.0, SparseArray 1.10.0, UCSC.utils 1.6.0, XML 3.99-0.19, abind 1.4-8, backports 1.5.0, base64enc 0.1-3, biomaRt 2.66.0, biovizBase 1.58.0, bit 4.6.0, bit64 4.6.0-1, bitops 1.0-9, blob 1.2.4, cachem 1.1.0, checkmate 2.3.3, cigarillo 1.0.0, cli 3.6.5, cluster 2.1.8.1, codetools 0.2-20, colorspace 2.1-2, compiler 4.5.1, crayon 1.5.3, curl 7.0.0, data.table 1.17.8, dbplyr 2.5.1, deldir 2.0-4, dichromat 2.0-0.1, digest 0.6.37, dplyr 1.1.4, ensemblDb 2.34.0, evaluate 1.0.5, farver 2.1.2, fastmap 1.2.0, filelock 1.0.3, foreign 0.8-90, ggplot2 4.0.0, glue 1.8.0, graph 1.88.0, grid 4.5.1, gridExtra 2.3, gtable 0.3.6, h5mread 1.2.0, hms 1.1.4, htmlTable 2.4.3, htmltools 0.5.8.1, htmlwidgets 1.6.4, httpuv 1.6.16, httr 1.4.7, httr2 1.2.1, interp 1.1-6, jpeg 0.1-11, jsonlite 2.0.0, knitr 1.50, later 1.4.4, lattice 0.22-7, latticeExtra 0.6-31, lazyeval 0.2.2, lifecycle 1.0.4, magrittr 2.0.4, memoise 2.0.1, mime 0.13, nnet 7.3-20, otl 0.2.0, parallel 4.5.1, pillar 1.11.1, pkgconfig 2.0.3, png 0.1-8, prettyunits 1.2.0, progress 1.2.3, promises 1.4.0, rappdirs 0.3.3, restfulr 0.0.16, rhdf5 2.54.0, rhdf5filters 1.22.0, rjson 0.2.23, rlang 1.1.6, rmarkdown 2.30, rpart 4.1.24, rstudioapi 0.17.1, scales 1.4.0, shiny 1.11.1, shinyTree 0.3.1, shinyjs 2.1.0, shinythemes 1.2.0, stringi 1.8.7, stringr 1.5.2, tibble 3.3.0, tidyselct 1.2.1, tools 4.5.1, vctrs 0.6.5, xfun 0.53, xtable 1.8-4, yaml 2.3.10