

# Package ‘ICAMSxtra’

September 22, 2020

**Title** ICAMSxtra

**Version** 0.0.1

**Author** Steve Rozen, Nanhai Jiang, Arnoud Boot, Mo Liu, Chang Jia Geng

**Maintainer** Steve Rozen <steverozen@gmail.com>

**Description** ICAMSxtra.

**biocViews**

**Imports** data.table,  
ICAMS

**Depends** R (>= 3.5)

**License** GPL-3

**Encoding** UTF-8

**LazyData** true

**Language** en-US

**Roxygen** list(markdown = TRUE)

**RoxygenNote** 7.1.1

**Suggests** BSgenome.Hsapiens.1000genomes.hs37d5,  
BSgenome.Hsapiens.UCSC.hg38,  
BSgenome.Mmusculus.UCSC.mm10,  
testthat

## R topics documented:

AnnotateIDVCFsWithTransRanges . . . . .	2
as.catalog.for.ID115 . . . . .	3
Canonicalize1Del115 . . . . .	4
CatalogRowOrder . . . . .	5
Collapse115CatalogTo83 . . . . .	5
CollapseID115CatalogsToID83s . . . . .	6
GRCh37.proportions . . . . .	6
GRCh38.proportions . . . . .	7
PlotExposure . . . . .	7
PlotExposureToPdf . . . . .	8
PlotID115AsID83ToPdf . . . . .	9
PlotID115Catalog . . . . .	10
PlotID115CatalogToPdf . . . . .	10

PlotTransBiasID115 . . . . .	11
PlotTransBiasID115ToPdf . . . . .	12
PlotTranscriptionAssociatedDamageToPdf . . . . .	13
ReadExposure . . . . .	14
ReadID115Catalog . . . . .	15
Reverse . . . . .	16
Reverse_pooled . . . . .	16
SortExposure . . . . .	16
Target . . . . .	17
Target_pooled . . . . .	17
VCFsToID115Catalogs . . . . .	18
VCFsToID115CatalogsAndPlotToPdf . . . . .	19
WriteExposure . . . . .	19
WriteID115Catalog . . . . .	20
<b>Index</b>	<b>21</b>

---

AnnotateIDVCFsWithTransRanges

*Add sequence context and transcript information to an in-memory ID VCF*

---

## Description

Add sequence context and transcript information to an in-memory ID VCF

## Usage

```
AnnotateIDVCFsWithTransRanges(
  ID.vcfs,
  ref.genome,
  trans.ranges = NULL,
  vcf.names = NULL
)
```

## Arguments

ID.vcfs	A list of in-memory ID VCF as a data.frame.
ref.genome	A ref.genome argument as described in <a href="#">ICAMS</a> .
trans.ranges	Optional. If ref.genome specifies one of the <a href="#">BSgenome</a> object <ol style="list-style-type: none"> <li><a href="#">BSgenome.Hsapiens.1000genomes.hs37d5</a></li> <li><a href="#">BSgenome.Hsapiens.UCSC.hg38</a></li> <li><a href="#">BSgenome.Mmusculus.UCSC.mm10</a></li> </ol> then the function will infer trans.ranges automatically. Otherwise, user will need to provide the necessary trans.ranges. Please refer to <a href="#">TranscriptRanges</a> for more details. If is.null(trans.ranges) do not add transcript range information.
vcf.names	list of names of the vcfs

**Value**

A list of in-memory ID VCFs each a `data.table`. These have been annotated with the sequence context (column name `seq.context`) and with transcript information in the form of a gene symbol (e.g. "TP53") and transcript strand. This information is in the columns `trans.start.pos`, `trans.end.pos`, `trans.strand`, `trans.Ensembl.gene.ID` and `trans.gene.symbol` in the output. These columns are not added if `is.null(trans.ranges)`.

**Examples**

```
file <- c(system.file("extdata/Strelka-ID-vcf",
                     "Strelka.ID.GRCh37.s1.vcf",
                     package = "ICAMSTra"))
list.of.vcfs <- ICAMS::ReadStrelkaIDVCFs(file)
if (requireNamespace("BSgenome.Hsapiens.1000genomes.hs37d5", quietly = TRUE)) {
  annotated.ID.vcfs <- AnnotateIDVCFsWithTransRanges(list.of.vcfs, ref.genome = "hg19",
                                                    trans.ranges = ICAMS::trans.ranges.GRCh37)
}
```

---

as.catalog.for.ID115    *Create a catalog from a matrix, data.frame, or vector*

---

**Description**

Create a catalog from a matrix, data.frame, or vector

**Usage**

```
as.catalog.for.ID115(
  object,
  ref.genome = NULL,
  region = "unknown",
  catalog.type = "counts",
  abundance = NULL,
  infer.rownames = FALSE
)
```

**Arguments**

object	A numeric matrix, numeric data.frame, or vector. If a vector, converted to a 1-column matrix with rownames taken from the element names of the vector and with column name "Unknown". If argument <code>infer.rownames</code> is FALSE then this argument must have rownames to denote the mutation types. See <a href="#">CatalogRowOrder</a> for more details.
ref.genome	A ref.genome argument as described in <a href="#">ICAMS</a> .
region	A character string designating a region, one of genome, transcript, exome, unknown; see <a href="#">ICAMS</a> .
catalog.type	One of "counts", "density", "counts.signature", "density.signature".
abundance	If NULL, then inferred if ref.genome is one of the reference genomes known to ICAMS and region is not unknown. See <a href="#">ICAMS</a> . The argument abundance should contain the counts of different source sequences for mutations in the same format as the numeric vectors in <a href="#">all.abundance</a> .

`infer.rownames` If TRUE, and object has no rownames, then assume the rows of object are in the correct order and add the rownames implied by the number of rows in object (e.g. rownames for SBS 192 if there are 192 rows). If TRUE, **be sure the order of rows is correct.**

### Value

A catalog as described in [ICAMS](#).

---

Canonicalize1Del115      *Given a deletion and its sequence context, categorize it*

---

### Description

This function is primarily for internal use, but we export it to document the underlying logic.

### Usage

```
Canonicalize1Del115(context, del.seq, pos, trace = 0, strand)
```

### Arguments

<code>context</code>	The deleted sequence plus ample surrounding sequence on each side (at least as long as <code>del.seq</code> ).
<code>del.seq</code>	The deleted sequence in context.
<code>pos</code>	The position of <code>del.seq</code> in context.
<code>trace</code>	If > 0, then generate messages tracing how the computation is carried out.
<code>strand</code>	NULL by default. But when called by <code>PlotTransBiasInternal</code> , <code>strand</code> is either + or -. The return value will include <code>:trans</code> or <code>:nontrans</code> indicating whether the deletion occurred on the transcribed or non-transcribed strand.

### Details

See [https://github.com/steverozen/ICAMS/raw/master/data-raw/PCAWG7\\_indel\\_classification\\_2017\\_12\\_08.xlsx](https://github.com/steverozen/ICAMS/raw/master/data-raw/PCAWG7_indel_classification_2017_12_08.xlsx) for additional information on deletion mutation classification.

This function first handles deletions in homopolymers, then handles deletions in simple repeats with longer repeat units, (e.g. CACACACA, see [FindMaxRepeatDel](#)), and if the deletion is not in a simple repeat, looks for microhomology (see [FindDeIMH](#)).

See the code for unexported function [CanonicalizeID](#) and the functions it calls for handling of insertions.

### Value

A string that is the canonical representation of the given deletion type. Return NA and raise a warning if there is an un-normalized representation of the deletion of a repeat unit. See [FindDeIMH](#) for details. (This seems to be very rare.)

@keywords internal

---

CatalogRowOrder	<i>Standard order of row names in a catalog</i>
-----------------	---

---

**Description**

This data is designed for those who need to create their own catalogs from formats not supported by this package. The rownames denote the mutation types. For example, for SBS96 catalogs, the rowname AGAT represents a mutation from AGA > ATA.

**Usage**

```
catalog.row.order
```

**Format**

A list of character vectors indicating the standard orders of row names in catalogs.

**Note**

In ID (small insertion and deletion) catalogs, deletion repeat sizes range from 0 to 5+, but for plotting and end-user documentation deletion repeat sizes range from 1 to 6+. In ID83 catalogs, deletion repeat sizes range from 0 to 5.

**Examples**

```
catalog.row.order$ID115
# There are altogether 115 row names to denote the mutation types
# in ID115 catalog.
# The difference from the .ID class in \code{\link{ICAMS::catalog.row.order}} is that
# single base nonhomopolymer indels have trinucleotide context added to them in the format
# INS/DEL:C/T:1:0_P_F where P is the preceding base and F is the following base.
```

---

Collapse115CatalogTo83	<i>"Collapse" a catalog</i>
------------------------	-----------------------------

---

**Description**

Collapse115CatalogTo83 Collapse a ID 115 catalog to a ID 83 catalog.

**Usage**

```
Collapse115CatalogTo83(catalog)
```

**Arguments**

catalog      A catalog as defined in [ICAMS](#).

**Value**

A catalog as defined in [ICAMS](#).

---

CollapseID115CatalogsToID83s

*"Collapse" a matrix of ID 115 catalogs to ID 83 catalog*

---

### Description

"Collapse" a matrix of ID 115 catalogs to ID 83 catalog

### Usage

CollapseID115CatalogsToID83s(catalogs)

### Arguments

catalogs            A catalog as defined in [ICAMS](#).

### Value

A catalog as defined in [ICAMS](#).

---

GRCh37.proportions

*Genic/intergenic size and proportions for H.sapiens BSgenome GRCh37*

---

### Description

This data is designed to be used in function [PlotTranscriptionAssociatedDamageToPdf](#)

### Usage

GRCh37.proportions

### Format

A list of 5 numbers with the names:

1. total.bp 2. coding.bp 3. noncoding.bp 4. prop.coding and 5. prop.noncoding

---

GRCh38.proportions	<i>Genic/intergenic size and proportions for H.sapiens BSgenome GRCh38</i>
--------------------	--

---

**Description**

This data is designed to be used in function [PlotTranscriptionAssociatedDamageToPdf](#)

**Usage**

```
GRCh38.proportions
```

**Format**

A list of 5 numbers with the names:

1. total.bp 2. coding.bp 3. noncoding.bp 4. prop.coding and 5. prop.noncoding

---

PlotExposure	<i>Plot exposures in multiple plots each with a manageable number of samples</i>
--------------	--

---

**Description**

Plot exposures in multiple plots each with a manageable number of samples

**Usage**

```
PlotExposure(
  exposure,
  samples.per.line = 30,
  plot.proportion = FALSE,
  xlim = NULL,
  ylim = NULL,
  legend.x = NULL,
  legend.y = NULL,
  cex.legend = 0.9,
  ...
)
```

**Arguments**

exposure	Exposures as a numerical matrix (or data.frame) with signatures in rows and samples in columns. Rownames are taken as the signature names and column names are taken as the sample IDs. If you want exposure sorted from largest to smallest, use <a href="#">SortExposure</a> . Do not use column names that start with multiple underscores. The exposures will often be mutation counts, but could also be e.g. mutations per megabase.
samples.per.line	Number of samples to show in each plot.

<code>plot.proportion</code>	Plot exposure proportions rather than counts.
<code>xlim, ylim</code>	Limits for the x and y axis. If NULL(default), the function tries to do something reasonable.
<code>legend.x, legend.y</code>	The x and y co-ordinates to be used to position the legend.
<code>cex.legend</code>	A numerical value giving the amount by which legend plotting text and symbols should be magnified relative to the default.
<code>...</code>	Other arguments passed to <code>barplot</code> . If <code>ylab</code> is not included, it defaults to a value depending on <code>plot.proportion</code> . If <code>col</code> is not supplied the function tries to do something reasonable.

### Value

An **invisible** list whose first element is a logic value indicating whether the plot is successful. The second element is a numeric vector giving the coordinates of all the bar midpoints drawn, useful for adding to the graph.

### Examples

```
file <- system.file("extdata",
                    "synthetic.exposure.csv",
                    package = "ICAMSxtra")
exposure <- ReadExposure(file)
PlotExposure(exposure[, 1:30])
```

---

PlotExposureToPdf	<i>Plot exposures in multiple plots each with a manageable number of samples to PDF</i>
-------------------	---

---

### Description

Plot exposures in multiple plots each with a manageable number of samples to PDF

### Usage

```
PlotExposureToPdf(
  exposure,
  file,
  mfrow = c(2, 1),
  mar = c(6, 4, 3, 2),
  oma = c(3, 2, 0, 2),
  samples.per.line = 30,
  plot.proportion = FALSE,
  xlim = NULL,
  ylim = NULL,
  legend.x = NULL,
  legend.y = NULL,
  cex.legend = 0.9,
  ...
)
```



**Arguments**

exposure	Exposures as a numerical matrix (or data.frame) with signatures in rows and samples in columns. Rownames are taken as the signature names and column names are taken as the sample IDs. If you want exposure sorted from largest to smallest, use <a href="#">SortExposure</a> . Do not use column names that start with multiple underscores. The exposures will often be mutation counts, but could also be e.g. mutations per megabase.
file	The name of the PDF file to be produced.
mfrow	A vector of the form c(nr, nc). Subsequent figures will be drawn in an nr-by-nc array on the device by rows.
mar	A numerical vector of the form c(bottom, left, top, right) which gives the number of lines of margin to be specified on the four sides of the plot.
oma	A vector of the form c(bottom, left, top, right) giving the size of the outer margins in lines of text.
samples.per.line	Number of samples to show in each plot.
plot.proportion	Plot exposure proportions rather than counts.
xlim, ylim	Limits for the x and y axis. If NULL(default), the function tries to do something reasonable.
legend.x, legend.y	The x and y co-ordinates to be used to position the legend.
cex.legend	A numerical value giving the amount by which legend plotting text and symbols should be magnified relative to the default.
...	Other arguments passed to <a href="#">barplot</a> . If ylab is not included, it defaults to a value depending on plot.proportion. If col is not supplied the function tries to do something reasonable.

**Value**

An **invisible** list whose first element is a logic value indicating whether the plot is successful. The second element is a numeric vector giving the coordinates of all the bar midpoints drawn, useful for adding to the graph.

**Examples**

```
file <- system.file("extdata",
                    "synthetic.exposure.csv",
                    package = "ICAMSxtra")
exposure <- ReadExposure(file)
PlotExposureToPdf(exposure, file = file.path(tempdir(), "exposure.pdf"))
```

---

PlotID115AsID83ToPdf	<i>Plot an ID 115 signatures (default) or catalog as standard ID83 and save as pdf file</i>
----------------------	---

---

**Description**

Plot an ID 115 signatures (default) or catalog as standard ID83 and save as pdf file

**Usage**

```
PlotID115AsID83ToPdf(catalog, file, ylim = NULL)
```

**Arguments**

catalog	A catalog as defined in <a href="#">ICAMS</a> .
file	The name of the PDF file to be produced.
ylim	Has the usual meaning. Only implemented for SBS96Catalog and IndelCatalog.

**Value**

A list whose first element is a logic value indicating whether the plot is successful. For **SBS192Catalog** with "counts" catalog.type and non-null abundance and plot.SBS12 = TRUE, the list will have a second element which is a list containing the strand bias statistics.

---

PlotID115Catalog	<i>Plot <b>one</b> spectrum or signature</i>
------------------	--

---

**Description**

Plot the spectrum of **one** sample or plot **one** signature. The type of graph is based on one attribute("catalog.type") of the input catalog. You can first use [TransformCatalog](#) to get different types of catalog and then do the plotting.

**Usage**

```
PlotID115Catalog(catalog, ylim = NULL)
```

**Arguments**

catalog	A catalog as defined in <a href="#">ICAMS</a> with attributes added. See <a href="#">as.catalog</a> for more details.
ylim	Has the usual meaning. Only implemented for SBS96Catalog and IndelCatalog.

---

PlotID115CatalogToPdf	<i>Plot catalog to a PDF file</i>
-----------------------	-----------------------------------

---

**Description**

Plot catalog to a PDF file. The type of graph is based on one attribute("catalog.type") of the input catalog. You can first use [TransformCatalog](#) to get different types of catalog and then do the plotting.

**Usage**

```
PlotID115CatalogToPdf(catalog, file, ylim = NULL)
```

**Arguments**

catalog	A catalog as defined in <a href="#">ICAMS</a> with attributes added. See <a href="#">as.catalog</a> for more details.
file	The name of the PDF file to be produced.
ylim	Has the usual meaning. Only implemented for SBS96Catalog and IndelCatalog.

**Value**

A list whose first element is a logic value indicating whether the plot is successful. For **SBS192Catalog** with "counts" catalog.type and non-null abundance and plot.SBS12 = TRUE, the list will have a second element which is a list containing the strand bias statistics.

**Note**

The sizes of repeats involved in deletions range from 0 to 5+ in the mutational-spectra and signature catalog rownames, but for plotting and end-user documentation deletion repeat sizes range from 1 to 6+.

---

PlotTransBiasID115	<i>Plot transcription strand bias</i>
--------------------	---------------------------------------

---

**Description**

Plot transcription strand bias

**Usage**

```
PlotTransBiasID115(annotated.ID.vcf, pool, damaged.base = NULL, ymax = NULL)
```

**Arguments**

annotated.ID.vcf	An ID VCF annotated by AnnotateIDVCFsWithTransRanges. It <b>must</b> have transcript range information added.
pool	if true, 36 categories will be pooled to 4 categories by removing trinucleotide context. This can be done if the counts of individual categories are too low, to increase power.
damaged.base	One of NULL, "purine" or "pyrimidine". This function allocates approximately equal numbers of mutations from damaged.base into each of num.of.bins bin by expression level. E.g. if damaged.base is "purine", then mutations from A and G will be allocated in approximately equal numbers to each expression-level bin. The rationale for the name damaged.base is that the direction of strand bias is a result of whether the damage occurs on a purine or pyrimidine. If NULL, the function attempts to infer the damaged.base based on mutation counts.
ymax	Limit for the y axis. If not specified, it defaults to NULL and the y axis limit equals 1.5 times of the maximum mutation counts in a specific mutation type.

**Value**

A list whose first element is a logic value indicating whether the plot is successful. The second element is a named numeric vector containing the p-values printed on the plot.

**Note**

The strand bias statistics are Benjamini-Hochberg q-values based on two-sided binomial tests of the mutation counts on the transcribed and untranscribed strands relative to the actual abundances of C and T on the transcribed strand. On the plot, asterisks indicate q-values as follows \*,  $Q < 0.05$ ; \*\*,  $Q < 0.01$ ; \*\*\*,  $Q < 0.001$ .

**Examples**

```
file <- c(system.file("extdata/Strelka-ID-vcf/",
                     "Strelka.ID.GRCh37.s1.vcf",
                     package = "ICAMSxtra"))
ID.vcf <- ICAMS::ReadStrelkaIDVCFs(file)
if (requireNamespace("BSgenome.Hsapiens.1000genomes.hs37d5", quietly = TRUE)) {
  annotated.ID.vcf <- AnnotateIDVCFsWithTransRanges(ID.vcf, ref.genome = "hg19",
                                                    trans.ranges = ICAMS::trans.ranges.GRCh37,
                                                    vcf.names = "Strelka.ID.GRCh37.s1.vcf")
  #' alternatively run below code to skip call to AnnotateIDVCFsWithTransRanges
  #' load(c(system.file("extdata/annotated.ID.vcf.rda", package = "ICAMSxtra")))
  PlotTransBiasID115(annotated.ID.vcf = annotated.ID.vcf[[1]],
                    pool = TRUE)
}
```

---

PlotTransBiasID115ToPdf

*Plot transcription strand bias to a PDF file*

---

**Description**

Plot transcription strand bias to a PDF file

**Usage**

```
PlotTransBiasID115ToPdf(annotated.ID.vcfs, file, pool, damaged.base = NULL)
```

**Arguments**

annotated.ID.vcfs	ID vcfs which have been annotated with AnnotateIDVCFsWithTransRanges.
file	The name of output file.
pool	if true, 36 categories will be pooled to 4 categories by removing trinucleotide context. This can be done if the counts of individual categories are too low, to increase power.
damaged.base	One of NULL, "purine" or "pyrimidine". This function allocates approximately equal numbers of mutations from damaged.base into each of num.of.bins bin by expression level. E.g. if damaged.base is "purine", then mutations from

A and G will be allocated in approximately equal numbers to each expression-level bin. The rationale for the name `damaged.base` is that the direction of strand bias is a result of whether the damage occurs on a purine or pyrimidine. If NULL, the function attempts to infer the `damaged.base` based on mutation counts.

### Value

A list whose first element is a logic value indicating whether the plot is successful. The second element is a named numeric vector containing the p-values printed on the plot.

### Note

The strand bias statistics are Benjamini-Hochberg q-values based on two-sided binomial tests of the mutation counts on the transcribed and untranscribed strands relative to the actual abundances of C and T on the transcribed strand. On the plot, asterisks indicate q-values as follows \*,  $Q < 0.05$ ; \*\*,  $Q < 0.01$ ; \*\*\*,  $Q < 0.001$ .

### Examples

```
library(ICAMS)
file <- c(system.file("extdata/Strelka-ID-vcf/",
                     "Strelka.ID.GRCh37.s1.vcf",
                     package = "ICAMSxtra"))
ID.vcf <- ICAMS::ReadStrelkaIDVCFs(file)
if (requireNamespace("BSgenome.Hsapiens.1000genomes.hs37d5", quietly = TRUE)) {
  annotated.ID.vcf <- AnnotateIDVCFsWithTransRanges(ID.vcf, ref.genome = "hg19",
                                                    trans.ranges = ICAMS::trans.ranges.GRCh37,
                                                    vcf.names = "Strelka.ID.GRCh37.s1")
  PlotTransBiasID115ToPdf(annotated.ID.vcfs = annotated.ID.vcf,
                          file = file.path(tempdir(), "test.pdf"),
                          pool = TRUE)
}
```

---

PlotTranscriptionAssociatedDamageToPdf

*Plot indel counts on transcribed and nontranscribed strands to pdf*

---

### Description

Plot indel counts on transcribed and nontranscribed strands to pdf

### Usage

```
PlotTranscriptionAssociatedDamageToPdf(
  list.of.vcfs,
  ref.genome,
  names.of.vcfs,
  proportions,
  file
)
```

**Arguments**

<code>list.of.vcfs</code>	List of in-memory ID VCFs. The list names will be the sample ids in the output catalog.
<code>ref.genome</code>	A <code>ref.genome</code> argument as described in <a href="#">ICAMS</a> .
<code>names.of.vcfs</code>	list of names of vcfs
<code>proportions</code>	The gene proportions for the genome e.g. <code>GRCh37.proportions</code> or <code>GRCh38.proportions</code>
<code>file</code>	The name of the PDF file to be produced.

**Value**

a list of tables of p-values for each vcf

**Note**

The strand bias statistics are Benjamini-Hochberg q-values based on two-sided binomial tests of the mutation counts on the transcribed and untranscribed strands relative to the actual abundances of C and T on the transcribed strand. On the plot, asterisks indicate q-values as follows \*,  $Q < 0.05$ ; \*\*,  $Q < 0.01$ ; \*\*\*,  $Q < 0.001$ .

**Examples**

```
dirs <- c(system.file("extdata/Strelka-ID-vcf", "Strelka.ID.GRCh37.s1.vcf", package="ICAMSxtra"),
          system.file("extdata/Strelka-ID-vcf", "Strelka.ID.GRCh37.s2.vcf", package="ICAMSxtra"))

list.of.vcfs <- ICAMS::ReadStrelkaIDVCFs(dirs, names.of.VCFs = c("s1", "s2"))
PlotTranscriptionAssociatedDamageToPdf(list.of.vcfs = list.of.vcfs,
                                       ref.genome = "hg19",
                                       names.of.vcfs = c("s1", "s2"),
                                       proportions = GRCh37.proportions,
                                       file = file.path(tempdir(), "test.pdf"))
```

---

ReadExposure

---

*Read an exposure matrix from a file*


---

**Description**

Read an exposure matrix from a file

**Usage**

```
ReadExposure(file, check.names = FALSE)
```

**Arguments**

<code>file</code>	CSV file containing an exposure matrix.
<code>check.names</code>	Passed to <a href="#">read.csv</a> . <b>IMPORTANT:</b> If TRUE this will replace the double colon in identifiers of the form <code>&lt;tumor_type&gt;::&lt;sample_id&gt;</code> with two periods (i.e. <code>&lt;tumor_type&gt;.&lt;sample_id&gt;</code> ). If <code>check.names</code> is true, generate a warning if double colons were present.

**Value**

Matrix of exposures.

**Examples**

```
file <- system.file("extdata",
                    "synthetic.exposure.csv",
                    package = "ICAMSxtra")
exposure <- ReadExposure(file)
```

---

ReadID115Catalog	<i>Read catalog</i>
------------------	---------------------

---

**Description**

Read a catalog in standardized format from path.

**Usage**

```
ReadID115Catalog(
  file,
  ref.genome = NULL,
  region = "unknown",
  catalog.type = "counts"
)
```

**Arguments**

file	Path to a catalog on disk in the standardized format.
ref.genome	A ref.genome argument as described in <a href="#">ICAMS</a> .
region	region A character string designating a genomic region; see <a href="#">as.catalog</a> and <a href="#">ICAMS</a> .
catalog.type	One of "counts", "density", "counts.signature", "density.signature".

**Details**

See also [WriteCatalog](#)

**Value**

A catalog as an S3 object; see [as.catalog](#).

**Comments**

To add or change attributes of the catalog, you can use function [attr](#).  
For example, `attr(catalog, "abundance") <- custom.abundance`.

**Note**

In ID (small insertion and deletion) catalogs, deletion repeat sizes range from 0 to 5+, but for plotting and end-user documentation deletion repeat sizes range from 1 to 6+.

---

Reverse	<i>Transcription bias of indels classified into 115 categories (purine)</i>
---------	---

---

**Description**

This data is designed to be used as an example in function [PlotTransBiasID115](#) and [PlotTransBiasID115ToPdf](#).

**Usage**

```
reverse
```

**Format**

A vector which contains the 115 categories of indel events, but in purine format  
An object of class character of length 36.

---

Reverse_pooled	<i>Transcription bias of indels classified into 13 categories (purine)</i>
----------------	--

---

**Description**

This data is designed to be used as an example in function [PlotTransBiasID115](#) and [PlotTransBiasID115](#) when pool = TRUE.

**Usage**

```
reverse_pooled
```

**Format**

A vector which contains the 13 categories of indel events, standardised to purine format.  
An object of class character of length 4.

---

SortExposure	<i>Sort columns of an exposure matrix from largest to smallest (or vice versa)</i>
--------------	--

---

**Description**

Sort columns of an exposure matrix from largest to smallest (or vice versa)

**Usage**

```
SortExposure(exposure, decreasing = TRUE)
```



**Arguments**

- `exposure` Exposures as a numerical matrix (or `data.frame`) with signatures in rows and samples in columns. Rownames are taken as the signature names and column names are taken as the sample IDs.
- `decreasing` If TRUE, sort from largest to smallest.

**Value**

The original exposure with columns sorted.

**Examples**

```
file <- system.file("extdata",
                    "synthetic.exposure.csv",
                    package = "ICAMSxtra")
exposure <- ReadExposure(file)
exposure.sorted <- SortExposure(exposure)
```

---

Target	<i>Transcription bias of indels classified into 115 categories (pyrimidine)</i>
--------	---

---

**Description**

This data is designed to be used as an example in function [PlotTransBiasID115](#) and [PlotTransBiasID115ToPdf](#).

**Usage**

`target`

**Format**

A vector which contains the 115 categories of indel events, standardised to pyrimidine format.  
An object of class character of length 36.

---

Target_pooled	<i>Transcription bias of indels classified into 13 categories (pyrimidine)</i>
---------------	--

---

**Description**

This data is designed to be used as an example in function [PlotTransBiasID115](#) and [PlotTransBiasID115](#)

**Usage**

`target_pooled`

**Format**

A vector which contains the 13 categories of indel events, standardised to pyrimidine format.  
An object of class character of length 4.

---

VCFsToID115Catalogs      *Create ID (small insertion and deletion) catalog from ID VCFs*


---

## Description

Create ID (small insertion and deletion) catalog from ID VCFs

## Usage

```
VCFsToID115Catalogs(
  list.of.vcfs,
  ref.genome,
  region = "unknown",
  flag.mismatches = 0
)
```

## Arguments

<code>list.of.vcfs</code>	List of in-memory ID VCFs. The list names will be the sample ids in the output catalog.
<code>ref.genome</code>	A <code>ref.genome</code> argument as described in <a href="#">ICAMS</a> .
<code>region</code>	A character string acting as a region identifier, one of "genome", "exome".
<code>flag.mismatches</code>	Optional. If > 0, then if there are mismatches to references in the ID (insertion/deletion) VCF, generate messages showing the mismatched rows and continue. Otherwise stop if there are mismatched rows. See <a href="#">AnnotateIDVCF</a> for more details.

## Value

A list of two elements. 1st element `catalog` is the ID (small insertion and deletion) catalog with attributes added. See [as.catalog](#) for more details. 2nd element `annotated.vcfs` is a list of data frames which contain the original VCF with three additional columns `seq.context.width`, `seq.context` and `ID.class` added. The category assignment of each ID mutation in VCF can be obtained from `ID.class` column.

## Note

In ID (small insertion and deletion) catalogs, deletion repeat sizes range from 0 to 5+, but for plotting and end-user documentation deletion repeat sizes range from 1 to 6+.

---

VCFsToID115CatalogsAndPlotToPdf

*Read a list of vcfs and plot ID115 catalogs as pdf*


---

### Description

Read a list of vcfs and plot ID115 catalogs as pdf

### Usage

```
VCFsToID115CatalogsAndPlotToPdf(
  list.of.vcfs,
  ref.genome,
  region = "unknown",
  flag.mismatches = 0,
  file,
  ylim = NULL
)
```

### Arguments

<code>list.of.vcfs</code>	List of in-memory ID VCFs. The list names will be the sample ids in the output catalog.
<code>ref.genome</code>	A <code>ref.genome</code> argument as described in <a href="#">ICAMS</a> .
<code>region</code>	A character string acting as a region identifier, one of "genome", "exome".
<code>flag.mismatches</code>	Optional. If > 0, then if there are mismatches to references in the ID (insertion/deletion) VCF, generate messages showing the mismatched rows and continue. Otherwise stop if there are mismatched rows. See <a href="#">AnnotateIDVCF</a> for more details.
<code>file</code>	The name of the PDF file to be produced.
<code>ylim</code>	Has the usual meaning. Only implemented for SBS96Catalog and IndelCatalog.

---

WriteExposure

*Write an exposure matrix to a file*


---

### Description

Write an exposure matrix to a file

### Usage

```
WriteExposure(exposure, file)
```

### Arguments

<code>exposure</code>	Exposures as a numerical matrix (or <code>data.frame</code> ) with signatures in rows and samples in columns. Rownames are taken as the signature names and column names are taken as the sample IDs.
<code>file</code>	File to which to write the exposure matrix (as a CSV file).

**Examples**

```
file <- system.file("extdata",  
                    "synthetic.exposure.csv",  
                    package = "ICAMExtra")  
exposure <- ReadExposure(file)  
WriteExposure(exposure, file = file.path(tempdir(), "synthetic.exposure.csv"))
```

---

WriteID115Catalog	<i>Write a catalog to a file.</i>
-------------------	-----------------------------------

---

**Description**

Write a catalog to a file.

**Usage**

```
WriteID115Catalog(catalog, file, strict = TRUE)
```

**Arguments**

catalog	A catalog as defined in <a href="#">ICAMS</a> with attributes added. See <a href="#">as.catalog</a> for more details.
file	The path of the file to be written.
strict	If TRUE, then stop if additional checks on the input fail.

# Index

## \* datasets

- CatalogRowOrder, [5](#)
  - GRCh37.proportions, [6](#)
  - GRCh38.proportions, [7](#)
  - Reverse, [16](#)
  - Reverse\_pooled, [16](#)
  - Target, [17](#)
  - Target\_pooled, [17](#)
- all.abundance, [3](#)
- AnnotateIDVCF, [18](#), [19](#)
- AnnotateIDVCFsWithTransRanges, [2](#)
- as.catalog, [10](#), [11](#), [15](#), [18](#), [20](#)
- as.catalog.for.ID115, [3](#)
- attr, [15](#)
- barplot, [8](#), [9](#)
- BSgenome, [2](#)
- BSgenome.Hsapiens.1000genomes.hs37d5, [2](#)
- BSgenome.Hsapiens.UCSC.hg38, [2](#)
- BSgenome.Mmusculus.UCSC.mm10, [2](#)
- Canonicalize1Del115, [4](#)
- CanonicalizeID, [4](#)
- catalog.row.order (CatalogRowOrder), [5](#)
- CatalogRowOrder, [3](#), [5](#)
- Collapse115CatalogTo83, [5](#)
- CollapseID115CatalogsToID83s, [6](#)
- FindDelMH, [4](#)
- FindMaxRepeatDel, [4](#)
- GRCh37.proportions, [6](#)
- GRCh38.proportions, [7](#)
- ICAMS, [2–6](#), [10](#), [11](#), [14](#), [15](#), [18–20](#)
- PlotExposure, [7](#)
- PlotExposureToPdf, [8](#)
- PlotID115AsID83ToPdf, [9](#)
- PlotID115Catalog, [10](#)
- PlotID115CatalogToPdf, [10](#)
- PlotTransBiasID115, [11](#), [16](#), [17](#)
- PlotTransBiasID115ToPdf, [12](#), [16](#), [17](#)
- PlotTranscriptionAssociatedDamageToPdf, [6](#), [7](#), [13](#)
- read.csv, [14](#)
- ReadExposure, [14](#)
- ReadID115Catalog, [15](#)
- Reverse, [16](#)
- reverse (Reverse), [16](#)
- Reverse\_pooled, [16](#)
- reverse\_pooled (Reverse\_pooled), [16](#)
- SortExposure, [7](#), [9](#), [16](#)
- Target, [17](#)
- target (Target), [17](#)
- Target\_pooled, [17](#)
- target\_pooled (Target\_pooled), [17](#)
- TranscriptRanges, [2](#)
- TransformCatalog, [10](#)
- VCFsToID115Catalogs, [18](#)
- VCFsToID115CatalogsAndPlotToPdf, [19](#)
- WriteCatalog, [15](#)
- WriteExposure, [19](#)
- WriteID115Catalog, [20](#)