# Package 'RareComb'

January 20, 2025

Title Combinatorial and Statistical Analyses of Rare Events

Version 1.1

**Description** A custom implementation of the apriori algorithm and binomial tests to identify combinations of features (genes, variants etc) significantly enriched for simultaneous mutations/events from sparse Boolean input, see Vijay Kumar Pounraja, Santhosh Girirajan (2021). Version 1.1 includes a minor adjustment to the number of combinations to be considered for multiple testing correction. This updated version is more conservative in its approach and hence more selective. <doi:10.1101/2021.10.01.462832>.

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**Encoding** UTF-8

LazyData true

**Depends** R (>= 2.10)

**Imports** magrittr, arules, dplyr, methods, pwr, stringr, tidyr, reshape2, sqldf

RoxygenNote 7.1.1

NeedsCompilation no

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analyze\_in\_out\_simultaneity

Analyze relationships between rare events among multiple input and output variables

# Description

This function takes a Boolean dataframe as input and analyzes the relationship between input and output variables for the combinations that that include at least a single output variable andmeet all the input criteria specified by the user.

#### Usage

<pre>boolean_input_mult_df</pre>		
	An input Boolean dataframe with multiple input and outcome variables	
combo_length	The length of the combinations specified by the user	
min_output_count		
	Minimum number of output variables present in the combination	
<pre>max_output_coun</pre>	t	
	Maximum number of output variables present in the combination	
<pre>min_indv_threshold</pre>		
	Minimum number of instances that support the combination	
<pre>max_freq_threshold</pre>		
	Maximum fraction of the cohort size that could support a combination (i.e., filter out highly frequent events)	
input_format	Optional   Naming convention used for input variables (Default = 'Input_')	
output_format	Optional   Naming convention used for output variables (Default = 'Output_')	
<pre>pval_filter_threshold</pre>		
	Optional   p-value cut-off to use to identify significant combinations (Default = $0.05$ )	

```
adj_pval_type Optional | Type of multiple testing corrections to use (Default = 'BH'; Alterna-
tive option = 'bonferroni')
```

# Value

A dataframe with the list of multiple-testing adjusted statistically significant combinations along with quantitative measures (frequencies, p-values etc) that support the findings.

# Author(s)

Vijay Kumar Pounraja

# Examples

<pre>boolean_input_df</pre>	Sparse Boolean dataframe with rare variant information and a single
	outcome variable

# Description

A synthetic dataset containing information about 5000 individuals (rows) and 1000 rare variants (columns).

# Usage

boolean\_input\_df

# Format

A data frame with 5000 rows and 1002 variables:

Sample\_Name Unique identifier of the samples

Input\_1 Presence and absense of rare variant 1

Input\_2 Presence and absense of rare variant 2

Input\_3 Presence and absense of rare variant 3

Input\_4 Presence and absense of rare variant 4

Input\_5 Presence and absense of rare variant 5

Input\_6 Presence and absense of rare variant 6

Input 7 Presence and absense of rare variant 7

Input\_8 Presence and absense of rare variant 8

- Input\_9 Presence and absense of rare variant 9 Input\_10 Presence and absense of rare variant 10 Input 11 Presence and absense of rare variant 11 **Input 12** Presence and absense of rare variant 12 **Input 13** Presence and absense of rare variant 13 **Input 14** Presence and absense of rare variant 14 **Input 15** Presence and absense of rare variant 15 **Input 16** Presence and absense of rare variant 16 Input 17 Presence and absense of rare variant 17 **Input 18** Presence and absense of rare variant 18 Input\_19 Presence and absense of rare variant 19 **Input 20** Presence and absense of rare variant 20 Input\_21 Presence and absense of rare variant 21 **Input 22** Presence and absense of rare variant 22 Input 23 Presence and absense of rare variant 23 Input 24 Presence and absense of rare variant 24 Input 25 Presence and absense of rare variant 25 Input\_26 Presence and absense of rare variant 26 Input 27 Presence and absense of rare variant 27 Input 28 Presence and absense of rare variant 28 Input 29 Presence and absense of rare variant 29 **Input 30** Presence and absense of rare variant 30 Input\_31 Presence and absense of rare variant 31 **Input\_32** Presence and absense of rare variant 32 Input 33 Presence and absense of rare variant 33 Input 34 Presence and absense of rare variant 34 Input 35 Presence and absense of rare variant 35 Input\_36 Presence and absense of rare variant 36 Input\_37 Presence and absense of rare variant 37 Input 38 Presence and absense of rare variant 38 Input 39 Presence and absense of rare variant 39 Input 40 Presence and absense of rare variant 40 Input 41 Presence and absense of rare variant 41 Input\_42 Presence and absense of rare variant 42 Input\_43 Presence and absense of rare variant 43 Input 44 Presence and absense of rare variant 44
- Input\_45 Presence and absense of rare variant 45

Input\_46 Presence and absense of rare variant 46 Input\_47 Presence and absense of rare variant 47 **Input 48** Presence and absense of rare variant 48 Input 49 Presence and absense of rare variant 49 **Input 50** Presence and absense of rare variant 50 **Input 51** Presence and absense of rare variant 51 **Input 52** Presence and absense of rare variant 52 **Input 53** Presence and absense of rare variant 53 **Input 54** Presence and absense of rare variant 54 Input 55 Presence and absense of rare variant 55 Input\_56 Presence and absense of rare variant 56 Input 57 Presence and absense of rare variant 57 **Input 58** Presence and absense of rare variant 58 Input 59 Presence and absense of rare variant 59 Input 60 Presence and absense of rare variant 60 Input 61 Presence and absense of rare variant 61 **Input 62** Presence and absense of rare variant 62 Input\_63 Presence and absense of rare variant 63 Input 64 Presence and absense of rare variant 64 Input 65 Presence and absense of rare variant 65 Input 66 Presence and absense of rare variant 66 Input\_67 Presence and absense of rare variant 67 Input\_68 Presence and absense of rare variant 68 Input\_69 Presence and absense of rare variant 69 **Input 70** Presence and absense of rare variant 70 Input 71 Presence and absense of rare variant 71 **Input 72** Presence and absense of rare variant 72 Input\_73 Presence and absense of rare variant 73 Input\_74 Presence and absense of rare variant 74 Input 75 Presence and absense of rare variant 75 Input 76 Presence and absense of rare variant 76 Input 77 Presence and absense of rare variant 77 **Input 78** Presence and absense of rare variant 78 **Input 79** Presence and absense of rare variant 79 **Input 80** Presence and absense of rare variant 80 Input 81 Presence and absense of rare variant 81 Input\_82 Presence and absense of rare variant 82

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Input\_119 Presence and absense of rare variant 119

Input\_120 Presence and absense of rare variant 120 Input 121 Presence and absense of rare variant 121 **Input 122** Presence and absense of rare variant 122 **Input 123** Presence and absense of rare variant 123 **Input 124** Presence and absense of rare variant 124 **Input 125** Presence and absense of rare variant 125 **Input 126** Presence and absense of rare variant 126 **Input 127** Presence and absense of rare variant 127 Input 128 Presence and absense of rare variant 128 Input 129 Presence and absense of rare variant 129 Input\_130 Presence and absense of rare variant 130 Input 131 Presence and absense of rare variant 131 **Input 132** Presence and absense of rare variant 132 Input 133 Presence and absense of rare variant 133 Input 134 Presence and absense of rare variant 134 **Input 135** Presence and absense of rare variant 135 **Input 136** Presence and absense of rare variant 136 **Input 137** Presence and absense of rare variant 137 Input 138 Presence and absense of rare variant 138 Input 139 Presence and absense of rare variant 139 Input 140 Presence and absense of rare variant 140 Input\_141 Presence and absense of rare variant 141 Input\_142 Presence and absense of rare variant 142 **Input\_143** Presence and absense of rare variant 143 Input 144 Presence and absense of rare variant 144 Input 145 Presence and absense of rare variant 145 Input 146 Presence and absense of rare variant 146 Input\_147 Presence and absense of rare variant 147 Input\_148 Presence and absense of rare variant 148 Input 149 Presence and absense of rare variant 149 Input 150 Presence and absense of rare variant 150 Input 151 Presence and absense of rare variant 151 **Input 152** Presence and absense of rare variant 152 Input\_153 Presence and absense of rare variant 153 **Input 154** Presence and absense of rare variant 154 Input 155 Presence and absense of rare variant 155 Input\_156 Presence and absense of rare variant 156

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Input_490	Presence and absense of rare variant 490
Input_491	Presence and absense of rare variant 491
Input_492	Presence and absense of rare variant 492
Input_493	Presence and absense of rare variant 493
Input_494	Presence and absense of rare variant 494
Input_495	Presence and absense of rare variant 495
Input_496	Presence and absense of rare variant 496
Input_497	Presence and absense of rare variant 497
Input_498	Presence and absense of rare variant 498
Input_499	Presence and absense of rare variant 499
Input_500	Presence and absense of rare variant 500
Output_1	Disease outcome or phenotype

#### Description

A synthetic dataset containing information about 5000 individuals (rows) and 1000 rare variants (columns) and 3 outcome variables.

#### Usage

boolean\_input\_mult\_df

#### Format

A data frame with 5000 rows and 1004 variables:

Sample\_Name Unique identifier of the samples

Input\_1 Presence and absense of rare variant 1

Input\_2 Presence and absense of rare variant 2

Input\_3 Presence and absense of rare variant 3

Input\_4 Presence and absense of rare variant 4

Input\_5 Presence and absense of rare variant 5

Input\_6 Presence and absense of rare variant 6

Input\_7 Presence and absense of rare variant 7

Input\_8 Presence and absense of rare variant 8

Input\_9 Presence and absense of rare variant 9

Input\_10 Presence and absense of rare variant 10

Input\_11 Presence and absense of rare variant 11 Input\_12 Presence and absense of rare variant 12 **Input 13** Presence and absense of rare variant 13 Input 14 Presence and absense of rare variant 14 **Input 15** Presence and absense of rare variant 15 **Input 16** Presence and absense of rare variant 16 **Input 17** Presence and absense of rare variant 17 **Input 18** Presence and absense of rare variant 18 Input 19 Presence and absense of rare variant 19 **Input 20** Presence and absense of rare variant 20 Input\_21 Presence and absense of rare variant 21 **Input 22** Presence and absense of rare variant 22 Input 23 Presence and absense of rare variant 23 Input 24 Presence and absense of rare variant 24 Input 25 Presence and absense of rare variant 25 **Input 26** Presence and absense of rare variant 26 Input 27 Presence and absense of rare variant 27 Input\_28 Presence and absense of rare variant 28 Input 29 Presence and absense of rare variant 29 Input 30 Presence and absense of rare variant 30 Input 31 Presence and absense of rare variant 31 Input\_32 Presence and absense of rare variant 32 Input\_33 Presence and absense of rare variant 33 **Input 34** Presence and absense of rare variant 34 Input 35 Presence and absense of rare variant 35 **Input 36** Presence and absense of rare variant 36 Input 37 Presence and absense of rare variant 37 Input\_38 Presence and absense of rare variant 38 Input\_39 Presence and absense of rare variant 39 Input 40 Presence and absense of rare variant 40 Input 41 Presence and absense of rare variant 41 Input 42 Presence and absense of rare variant 42 **Input 43** Presence and absense of rare variant 43 Input 44 Presence and absense of rare variant 44 Input\_45 Presence and absense of rare variant 45 Input 46 Presence and absense of rare variant 46 Input\_47 Presence and absense of rare variant 47

Input\_48 Presence and absense of rare variant 48 Input\_49 Presence and absense of rare variant 49 **Input 50** Presence and absense of rare variant 50 Input 51 Presence and absense of rare variant 51 **Input 52** Presence and absense of rare variant 52 **Input 53** Presence and absense of rare variant 53 **Input 54** Presence and absense of rare variant 54 **Input 55** Presence and absense of rare variant 55 **Input 56** Presence and absense of rare variant 56 Input 57 Presence and absense of rare variant 57 Input\_58 Presence and absense of rare variant 58 **Input 59** Presence and absense of rare variant 59 **Input 60** Presence and absense of rare variant 60 Input 61 Presence and absense of rare variant 61 **Input 62** Presence and absense of rare variant 62 Input 63 Presence and absense of rare variant 63 Input 64 Presence and absense of rare variant 64 Input\_65 Presence and absense of rare variant 65 Input 66 Presence and absense of rare variant 66 Input 67 Presence and absense of rare variant 67 Input 68 Presence and absense of rare variant 68 Input\_69 Presence and absense of rare variant 69 Input\_70 Presence and absense of rare variant 70 Input\_71 Presence and absense of rare variant 71 **Input 72** Presence and absense of rare variant 72 Input 73 Presence and absense of rare variant 73 Input 74 Presence and absense of rare variant 74 Input\_75 Presence and absense of rare variant 75 Input\_76 Presence and absense of rare variant 76 Input 77 Presence and absense of rare variant 77 **Input 78** Presence and absense of rare variant 78 Input 79 Presence and absense of rare variant 79 **Input 80** Presence and absense of rare variant 80 Input\_81 Presence and absense of rare variant 81 **Input 82** Presence and absense of rare variant 82 Input 83 Presence and absense of rare variant 83 Input 84 Presence and absense of rare variant 84

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Input\_307 Presence and absense of rare variant 307 Input\_308 Presence and absense of rare variant 308 Input 309 Presence and absense of rare variant 309 Input 310 Presence and absense of rare variant 310 **Input 311** Presence and absense of rare variant 311 **Input 312** Presence and absense of rare variant 312 **Input 313** Presence and absense of rare variant 313 Input 314 Presence and absense of rare variant 314 Input 315 Presence and absense of rare variant 315 Input 316 Presence and absense of rare variant 316 Input\_317 Presence and absense of rare variant 317 **Input 318** Presence and absense of rare variant 318 Input 319 Presence and absense of rare variant 319 Input 320 Presence and absense of rare variant 320 Input 321 Presence and absense of rare variant 321 Input 322 Presence and absense of rare variant 322 Input 323 Presence and absense of rare variant 323 Input\_324 Presence and absense of rare variant 324 Input 325 Presence and absense of rare variant 325 Input 326 Presence and absense of rare variant 326 Input 327 Presence and absense of rare variant 327

- **Input\_328** Presence and absense of rare variant 328
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- **Input 340** Presence and absense of rare variant 340
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- Input\_487 Presence and absense of rare variant 487
- Input\_488 Presence and absense of rare variant 488
- **Input 489** Presence and absense of rare variant 489
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- Input\_491 Presence and absense of rare variant 491

Input_492 Presence and absense of rare variant 492
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Input_496 Presence and absense of rare variant 496
Input_497 Presence and absense of rare variant 497
Input_498 Presence and absense of rare variant 498
Input_499 Presence and absense of rare variant 499
Input_500 Presence and absense of rare variant 500
<b>Output_1</b> Disease outcome or phenotype 1
<b>Output_2</b> Disease outcome or phenotype 2
<b>Output_3</b> Disease outcome or phenotype 3

```
compare_enrichment
```

*Compare the enrichment in combinations of input variables between the binary outcomes (case/control)* 

# Description

This function takes a Boolean dataframe as input and quantifies the enrichment in the observed frequency of combinations that meet the criteria specified by the users compared to their corresponding expectation derived under the assumption of independence between the constituent elements of each combination. The function then reports the multiple-testing adjusted significant combinations in which enrichment is observed in cases but not in controls.

# Usage

boolean_input_df		
	An input Boolean dataframe with multiple input and a single binary outcome variable	
combo_length	The length of the combinations specified by the user	
<pre>min_indv_threshold</pre>		
	Minimum number of instances that support the combination	
max_freq_threshold		
	Maximum fraction of the cohort size that could support a combination (i.e., filter out highly frequent events)	

input_format	Optional   Naming convention used for input variables (Default = 'Input_')	
output_format	Optional   Naming convention used for output variables (Default = 'Output_')	
<pre>pval_filter_thr</pre>	eshold	
	Optional   p-value cut-off to use to identify significant combinations in cases (Default = $0.05$ )	
adj_pval_type	Optional   Type of multiple testing corrections to use (Default = 'BH'; Alternative option = 'bonferroni')	
<pre>min_power_threshold</pre>		
	Optional   Minimum statistical power (at 5% sig.threshold) required for significant combinations to be returned in the results (Default = $0.7$ )	
sample_names_ind		
	Optional   Indicator to specify if the output should includes row names that support each significant combination (Default = 'N'; Alternative option = 'Y')	

# Value

A dataframe with the list of multiple-testing adjusted statistically significant combinations along with quantitative measures (frequencies, p-values etc) that support the findings.

#### Author(s)

Vijay Kumar Pounraja

#### Examples

compare\_enrichment\_depletion

*Compare the enrichment in combinations of input variables between the binary outcomes (case/control)* 

#### Description

This function takes a Boolean dataframe as input and quantifies the enrichment in the observed frequency of combinations that meet the criteria specified by the users compared to their corresponding expectation derived under the assumption of independence between the constituent elements of each combination. The function then reports the multiple-testing adjusted significant combinations in which enrichment is observed in cases and depletion is observed in controls.

# Arguments

boolean_input_df		
	An input Boolean dataframe with multiple input and a single binary outcome variable	
combo_length	The length of the combinations specified by the user	
<pre>min_indv_thres</pre>	hold	
	Minimum number of instances that support the combination	
<pre>max_freq_thres</pre>	hold	
	Maximum fraction of the cohort size that could support a combination (i.e., filter out highly frequent events)	
input_format	Optional   Naming convention used for input variables (Default = 'Input_')	
output_format	Optional   Naming convention used for output variables (Default = 'Output_')	
pval_filter_threshold		
	Optional   p-value cut-off to use to identify significant combinations in cases (Default = 0.05)	
adj_pval_type	Optional   Type of multiple testing corrections to use (Default = 'BH'; Alterna- tive option = 'bonferroni')	
min_power_threshold		
	Optional   Minimum statistical power (at 5% sig.threshold) required for significant combinations to be returned in the results (Default = $0.7$ )	
sample_names_ind		
	Optional   Indicator to specify if the output should includes row names that support each significant combination (Default = 'N'; Alternative option = 'Y')	

#### Value

A dataframe with the list of multiple-testing adjusted statistically significant combinations along with quantitative measures (frequencies, p-values etc) that support the findings.

# Author(s)

Vijay Kumar Pounraja

#### Examples

compare\_enrichment\_modifiers

*Compare the enrichment in combinations of input variables between the binary outcomes (case/control)* 

### Description

This function takes a Boolean dataframe as input and quantifies the enrichment in the observed frequency of combinations that include at least one of the input variables supplied by the user as well as meet other user-specified criteria compared to their corresponding expectation derived under the assumption of independence between the constituent elements of each combination. The function then reports the combinations in which enrichment is observed in cases but not in controls.

# Usage

boolean_input_df		
	An input Boolean dataframe with multiple input and a single binary outcome variable	
combo_length	The length of the combinations specified by the user	
min_indv_thresh	nold	
	Minimum number of instances that support the combination	
<pre>max_freq_thresh</pre>	old	
	Maximum fraction of the cohort size that could support a combination (i.e., filter out highly frequent events)	
primary_input_e	ntities	
	List of variables that MUST be part of the combinations identified by the method	
input_format	Optional   Naming convention used for input variables (Default = 'Input_')	
output_format	Optional   Naming convention used for output variables (Default = 'Output_')	
pval_filter_threshold		
	Optional   p-value cut-off to use to identify significant combinations in cases (Default = $0.05$ )	
adj_pval_type	Optional   Type of multiple testing corrections to use (Default = 'BH'; Alternative option = 'bonferroni')	
min_power_threshold		
	Optional   Minimum statistical power (at 5% sig.threshold) required for significant combinations to be returned in the results (Default = $0.7$ )	
sample_names_ind		
	Optional   Indicator to specify if the output should includes row names that support each significant combination (Default = 'N'; Alternative option = 'Y')	

# Value

A dataframe with the list of multiple-testing adjusted statistically significant combinations along with quantitative measures (frequencies, p-values etc) that support the findings.

## Author(s)

Vijay Kumar Pounraja

#### Examples

compare\_expected\_vs\_observed

Compare the observed frequencies of combinations with their expected frequencies under the assumption of independence within a single group

# Description

This function takes a Boolean dataframe as input and compares the observed frequency of combinations that meet the criteria specified by the users with their corresponding expectation derived under the assumption of independence between the constituent elements of each combination

## Usage

boolean_input_df		
	An input Boolean dataframe with multiple input variables	
combo_length	The length of the combinations specified by the user	
<pre>min_indv_threshold</pre>		
	Minimum number of instances that support the combination	
<pre>max_freq_threshold</pre>		
	Maximum fraction of the cohort size that could support a combination (i.e., filter out highly frequent events)	
input_format	Optional   Naming convention used for input variables (Default = 'Input_')	

pval_filter_threshold	
	Optional   p-value cut-off to use for multiple testing adjustment (Default = $0.05$ )
adj_pval_type	Optional   Type of multiple testing corrections to use (Default = 'BH'; Alterna- tive option = 'bonferroni')

#### Value

A dataframe with the list of multiple-testing adjusted statistically significant combinations along with quantitative measures (frequencies, p-values etc) that support the findings.

# Author(s)

Vijay Kumar Pounraja

# Examples

custom_left_join	Perform successive left joins to fetch information about the constituent
	elements of the combinations

# Description

Fetching the frequency of multiple individual elements that make up the combinations of varying length and hence varying variable names or to join two similar data frames using identical variable names necessitates this function that supplements and joins data based on the length of the combinations.

#### Usage

```
custom_left_join(
   left_df,
   right_df,
   combo_length = combo_length,
   diff_colnames = diff_colnames
)
```

left_df	The data frame with information about the combinations
right_df	The data frame with information either about the combinations or their con- stituent elements
combo_length	The length of the combinations specified by the user used to determine the num- ber of successive joins to attempt

# input\_list

# Value

An output dataframe with the results of the join operation

#### Author(s)

Vijay Kumar Pounraja

input\_list

A list of 50 random input variables

#### Description

A list of 50 random input variables

# Usage

input\_list

#### Format

A list of 50 random input variables:

run\_apriori\_freqitems Generate frequent items using the apriori algorithm

#### Description

This function takes in a factorized Boolean matrix and generate frequent itemsets that meet all the user provided criteria provided by the calling function.

### Usage

```
run_apriori_freqitems(
    apriori_input_df,
    combo_length,
    support_threshold,
    input_colname_list,
    confidence_threshold = confidence_threshold,
    include_output_ind = include_output_ind,
    output_colname_list = output_colname_list
)
```

# Arguments

apriori_input_df		
	An input factorized Boolean dataframe with multiple input and outcome variables	
combo_length	The length of the combinations specified by the user	
<pre>support_threshold</pre>		
	Minimum support value calculated based on the minimum absolute observed frequency threshold specified by the user	
input_colname_list		
	A list of column names that identify the input variables	
confidence_threshold		
	Minimum confidence threshold specified by the user	
include_output_ind		
	Specifies if the outcome variables must also be made part of the analysis using the algorithm	
output_colname_list		
	A list of column names that identify the outcome variables	

# Details

This is a function leveraged by few of the four main methods available to the users.

# Value

A list of frequent item sets that meet all the constraints supplied to the apriori algorithm

#### Author(s)

Vijay Kumar Pounraja

run\_apriori\_rules Generate rules using the apriori algorithm

# Description

This function takes in a factorized Boolean matrix and generate rules that meet all the user provided criteria while restricting the RHS of the rule based on the list of variables allowed in RHS provided by the calling function.

```
run_apriori_rules(
    apriori_input_df,
    combo_length,
    support_threshold,
    input_colname_list,
    confidence_threshold = confidence_threshold,
    output_colname_list = output_colname_list
)
```

#### Arguments

apriori_input_d	f
	An input factorized Boolean dataframe with multiple input and outcome variables
combo_length	The length of the combinations specified by the user
support_thresho	ld
	Minimum support value calculated based on the minimum absolute observed frequency threshold specified by the user
<pre>input_colname_l</pre>	ist
	A list of column names that identify the input variables
confidence_thre	shold
	Minimum confidence threshold specified by the user
<pre>output_colname_</pre>	list
	Optional   A list of column names that identify the outcome variables

# Details

This is a function leveraged by few of the four main methods available to the users.

## Value

A list of rules that meet all the constraints supplied to the apriori algorithm

#### Author(s)

Vijay Kumar Pounraja

run\_apriori\_rules\_inout\_simult

Generate rules using the apriori algorithm

# Description

This function takes in a factorized Boolean matrix and generate rules that meet all the user provided criteria while allowing the outcome variables to be part of either LHS or RHS of the rules but restricting the input variables to the LHS of the rules.

```
run_apriori_rules_inout_simult(
    apriori_input_df,
    combo_length,
    support_threshold,
    input_colname_list,
    output_colname_list = output_colname_list
)
```

# Arguments

apriori_input_df		
	n input factorized Boolean dataframe with multiple input and outcome vari- les	
combo_length Th	e length of the combinations specified by the user	
<pre>support_threshold</pre>		
	inimum support value calculated based on the minimum absolute observed equency threshold specified by the user	
input_colname_list		
A	list of column names that identify the input variables	
output_colname_list		
Op	ptional   A list of column names that identify the outcome variables	

# Details

This is a function leveraged by few of the four main methods available to the users.

#### Value

A list of rules that meet all the constraints supplied to the apriori algorithm

# Author(s)

Vijay Kumar Pounraja

run\_apriori\_rules\_modifiers

Generate rules using the apriori algorithm

# Description

This function takes in a factorized Boolean matrix and generate rules that meet all the user provided criteria while restricting the RHS of the rule based on the list of variables allowed in RHS provided by the calling function.

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```
run_apriori_rules_modifiers(
    apriori_input_df,
    combo_length,
    support_threshold,
    input_colname_list,
    output_colname_list = output_colname_list
)
```

# Arguments

apriori_input_df		
	An input factorized Boolean dataframe with multiple input and outcome variables	
combo_length	The length of the combinations specified by the user	
<pre>support_threshold</pre>		
	Minimum support value calculated based on the minimum absolute observed frequency threshold specified by the user	
input_colname_list		
	A list of column names that identify the input variables	
output_colname_list		
	Optional   A list of column names that identify the outcome variables	

# Details

This is a function leveraged by few of the four main methods available to the users.

#### Value

A list of rules that meet all the constraints supplied to the apriori algorithm

# Author(s)

Vijay Kumar Pounraja

run\_apriori\_w\_sample\_names

Generate frequent items along with the names of supporting observations using the apriori algorithm

# Description

This function takes in a factorized Boolean matrix and generate frequent item sets that meet all the user provided criteria provided by the calling function. This function includes in it's output the identifiers of observations that support each significant combination.

```
run_apriori_w_sample_names(
    apriori_input_df,
    combo_length,
    support_threshold,
    input_colname_list,
    input_sample_list,
    confidence_threshold = confidence_threshold,
    include_output_ind = include_output_ind,
    output_colname_list = output_colname_list
)
```

#### Arguments

apriori\_input\_df An input factorized Boolean dataframe with multiple input and outcome variables combo\_length The length of the combinations specified by the user support\_threshold Minimum support value calculated based on the minimum absolute observed frequency threshold specified by the user input\_colname\_list A list of column names that identify the input variables input\_sample\_list A list of row names that identify the samples/observations confidence\_threshold Minimum confidence threshold specified by the user include\_output\_ind Specifies if the outcome variables must also be made part of the analysis using the algorithm output\_colname\_list A list of column names that identify the outcome variables

# Details

This is a function leveraged by few of the four main methods available to the users.

#### Value

A list of frequent item sets that meet all the constraints supplied to the apriori algorithm

# Author(s)

Vijay Kumar Pounraja

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