

# Package ‘IFAA’

September 30, 2020

**Title** IFAA: Robust association identification and Inference For Absolute Abundance in microbiome analyses

**Version** 0.0.0.9000

**Description** IFAA is a novel approach to make inference on the association of covariates with the absolute abundance (AA) of microbiome in an ecosystem.

**License** GNU General Public License version 2

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 7.1.1

**Depends** picasso (>= 1.2.0),  
glmnet (>= 2.0-16),  
expm (>= 0.999-3),  
foreach (>= 1.4.3),  
snow (>= 0.4-2),  
doSNOW (>= 1.0.15),  
rlecuyer (>= 0.3-3),  
Matrix (>= 1.2-14),  
HDCI (>= 1.0-2),  
doParallel (>= 1.0.11),  
future (>= 1.12.0)

**Suggests** knitr,  
rmarkdown

**VignetteBuilder** knitr

## R topics documented:

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IFAA	<i>Robust association identification and inference for absolute abundance in microbiome analyses</i>
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## Description

Make inference on the association of covariates of microbiome

## Usage

```
IFAA(
  MicrobData,
  CovData,
  linkIDname,
  testCov = NULL,
  ctrlCov = NULL,
  testMany = T,
  ctrlMany = F,
  nRef = 40,
  nRefMaxForEsti = 1,
  nPermu = 40,
  x1permut = T,
  refTaxa = NULL,
  reguMethod = c("mcp"),
  fwerRate = 0.25,
  paraJobs = NULL,
  bootB = 500,
  bootLassoAlpha = 0.05,
  standardize = F,
  sequentialRun = F,
  allFunc = allUserFunc(),
  refReadsThresh = 0.2,
  SDThresh = 0.05,
  SDquantilThresh = 0,
  balanceCut = 0.2,
  seed = 1
)
```

## Arguments

MicrobData	Microbiome data matrix containing microbiome abundance with each row per sample and each column per taxon/OTU/ASV. It should contain an "id" variable to correspond to the "id" variable in the covariates data: CovData.
CovData	Covariates data matrix containing covariates and confounders with each row per sample and each column per variable. It should also contain an "id" variable to correspond to the "id" variable in the microbiome data: MicrobData.
linkIDname	Variable name of the "id" variable in both MicrobData and CovData. The two data sets will be merged by this "id" variable.
testCov	Covariates that are of primary interest for testing and estimating the associations. It corresponds to $X_i$ in the equation. Default is NULL which means all covariates are testCov.
ctrlCov	Potential confounders that will be adjusted in the model. It corresponds to $W_i$ in the equation. Default is NULL which means all covariates except those in testCov are adjusted as confounders.
testMany	This takes logical value TRUE or FALSE. If TRUE, the testCov will contain all the variables in CovData provided testCov is set to be NULL. The default value is TRUE which does not do anything if testCov is not NULL.
ctrlMany	This takes logical value TRUE or FALSE. If TRUE, all variables except testCov are considered as control covariates provided ctrlCov is set to be NULL. The default value is TRUE which does not do anything if ctrlCov is not NULL.

nRef	The number of randomly picked reference taxa used in phase 1. Default number is 40.
nRefMaxForEsti	The maximum number of reference taxa used in phase 2. The default is 1.
nPermu	The number of permutation used in phase 1. Default number is 40.
refTaxa	Reference taxa will be used in phase 1. The default is NULL since the algorithm will pick those randomly. If a vector of taxa names are provided, the algorithm will use the provided taxa instead of randomly picking random reference taxa.
reguMethod	regularization approach used in phase 1 of the algorithm. Take value "mcp" or "lasso", default is "mcp".
fwerRate	The family wise error rate for identifying taxa/OTU/ASV associated with testCov in phase 1. Default is 0.25.
paraJobs	Number of parallel jobs that will be registered to run the algorithm. Default is 8. If specified as NULL, it will automatically detect the cores to decide the number of parallel jobs.
bootB	Number of bootstrap samples for obtaining confidence interval of estimates in phase 2. The default is 500.
bootLassoAlpha	The significance level in phase 2. Default is 0.05.
standardize	This takes a logical value TRUE or FALSE. If TRUE, all design matrix X in phase 1 and phase 2 will be standardized in the analyses. Default is FALSE.
sequentialRun	This takes a logical value TRUE or FALSE. Sometimes parallel jobs can not be successfully run for unknown reasons (such as hardware issues). For example, socket related errors may pop up or some slave cores return error message instead of numerical results. In those scenarios, setting sequentialRun = TRUE may help, but it will take more time to run. Default is FALSE.
refReadsThresh	The threshold of non-zero sequencing reads for choosing the reference taxon in phase 2. The default is 0.2 which means at least 20% non-zero sequencing reads.
SDThresh	The threshold of standard deviations of sequencing reads for choosing the reference taxon in phase 2. The default is 0.5 which means the standard deviation of sequencing reads should be at least 0.5.
balanceCut	The threshold of non-zero sequencing reads in each group of a binary variable for choosing the reference taxon in phase 2. The default number is 0.2 which means at least 20% sequencing reads are non-zero in each group.
seed	Random seed for reproducibility. Default is 1.

## Details

The IFAA() uses a novel approach to make inference on the association of covariates with the absolute abundance (AA) of microbiome in an ecosystem.

## Value

A list containing the estimation results.

- analysisResults\$estByCovList: A list containing estimating results for all the variables in testCov. See details.
- covariatesData: A dataset containing covariates and confounders used in the analyses.

## References

<https://arxiv.org/pdf/1909.10101.pdf>

## Examples

```
data(dataM)
dim(dataM)
dataM[1:5, 1:8]
data(dataC)
dim(dataC)
dataC[1:5, ]
results <- IFAA(MicrobData = dataM,
                CovData = dataC,
                linkIDname = "id",
                testCov = c("v1", "v2"),
                ctrlCov = c("v3"), nRef = 4,
                nPermu = 4,
                fwerRate = 0.25,
                bootB = 5)
```

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