

# Package ‘MEDseq’

March 28, 2022

**Type** Package

**Date** 2022-03-28

**Title** Mixtures of Exponential-Distance Models with Covariates

**Version** 1.3.3

**Description** Implements a model-based clustering method for categorical life-course sequences relying on mixtures of exponential-distance models introduced by Murphy et al. (2021) <[doi:10.1111/rssa.12712](https://doi.org/10.1111/rssa.12712)>. A range of flexible precision parameter settings corresponding to weighted generalisations of the Hamming distance metric are considered, along with the potential inclusion of a noise component. Gating covariates can be supplied in order to relate sequences to baseline characteristics. Sampling weights are also accommodated. The models are fitted using the EM algorithm and tools for visualising the results are also provided.

**Depends** R (>= 4.0.0)

**License** GPL (>= 2)

**Encoding** UTF-8

**URL** <https://cran.r-project.org/package=MEDseq>

**BugReports** <https://github.com/Keefe-Murphy/MEDseq>

**LazyData** true

**Imports** cluster, matrixStats (>= 0.53.1), nnet (>= 7.3-0), seriation, stringdist, TraMineR (>= 1.6), WeightedCluster

**Suggests** knitr, rmarkdown, viridisLite (>= 0.4.0)

**RoxygenNote** 7.1.2

**VignetteBuilder** knitr

**NeedsCompilation** no

**Author** Keefe Murphy [aut, cre] (<<https://orcid.org/0000-0002-7709-3159>>),  
Thomas Brendan Murphy [ctb] (<<https://orcid.org/0000-0002-5668-7046>>),  
Raffaella Piccarreta [ctb],  
Isobel Claire Gormley [ctb] (<<https://orcid.org/0000-0001-7713-681X>>)

**Maintainer** Keefe Murphy <[keefe.murphy@mu.ie](mailto:keefe.murphy@mu.ie)>

**Repository** CRAN

**Date/Publication** 2022-03-28 15:30:08 UTC

## R topics documented:

MEDseq-package . . . . .	2
biofam . . . . .	4
dbns . . . . .	6
get_MEDseq_results . . . . .	7
MEDseq_clustnames . . . . .	9
MEDseq_compare . . . . .	11
MEDseq_control . . . . .	15
MEDseq_entropy . . . . .	19
MEDseq_fit . . . . .	21
MEDseq_meantime . . . . .	26
MEDseq_news . . . . .	29
MEDseq_stderr . . . . .	29
mvad . . . . .	31
plot.MEDseq . . . . .	33
predict.MEDgating . . . . .	39
wKModes . . . . .	41
<b>Index</b>	<b>45</b>

---

MEDseq-package	<i>MEDseq: Mixtures of Exponential-Distance Models with Covariates</i>
----------------	--

---

### Description

Fits MEDseq models: mixtures of Exponential-Distance models with gating covariates and sampling weights. Typically used for clustering categorical/longitudinal life-course sequences.

### Usage

Fits `_MEDseq_` models introduced by Murphy et al. (2021) <doi: [10.1111/rssa.12712](https://doi.org/10.1111/rssa.12712)>, i.e. fits mixtures of exponential-distance models for clustering longitudinal life-course sequence data via the EM/CEM algorithm.

A family of parsimonious precision parameter constraints are accommodated. So too are sampling weights. Gating covariates can be supplied via formula interfaces.

The most important function in the **MEDseq** package is: `MEDseq_fit`, for fitting the models via EM/CEM.

`MEDseq_control` allows supplying additional arguments which govern, among other things, controls on the initialisation of the allocations for the EM/CEM algorithm and the various model selection options.

`MEDseq_compare` is provided for conducting model selection between different results from using different covariate combinations &/or initialisation strategies, etc.

`MEDseq_stderr` is provided for computing the standard errors of the coefficients for the covariates in the gating network.

A dedicated plotting function `plot.MEDseq` exists for visualising various aspects of the results, using new methods as well as some existing methods adapted from the **TraMineR** package.

Finally, the package also contains two data sets: `biofam` and `mvad`.

### Details

- Type: Package
- Package: MEDseq
- Version: 1.3.3
- Date: 2022-03-28 (this version), 2019-08-24 (original release)
- Licence: GPL (>=2)

### See Also

Further details and examples are given in the associated vignette document:  
`vignette("MEDseq", package = "MEDseq")`

### Author(s)

Keefe Murphy [aut, cre], Thomas Brendan Murphy [ctb], Raffaella Piccarreta [ctb], Isobel Claire Gormley [ctb]

**Maintainer:** Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

### See Also

Useful links:

- <https://cran.r-project.org/package=MEDseq>
- Report bugs at <https://github.com/Keefe-Murphy/MEDseq>

### Examples

```
# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
```

```

states      <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels      <- c("Employment", "Further Education", "Higher Education",
               "Joblessness", "School", "Training")
mvad.seq    <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a range of unweighted models without covariates
# Only consider models with a noise component
# Supply some MEDseq_control() arguments
mod1        <- MEDseq_fit(mvad.seq, G=9:10, modtype=c("CCN", "CUN", "UCN", "UUN"),
                        algo="CEM", init.z="kmodes", criterion="icl")

# Fit a model with weights and gating covariates
# Have the probability of noise-component membership be constant
mod2        <- MEDseq_fit(mvad.seq, G=11, modtype="UUN", weights=mvad$weights,
                        gating=~ gcse5eq, covars=mvad.cov, noise.gate=FALSE)

# Examine this model and its gating network
summary(mod2, network=TRUE)
plot(mod2, "clusters")

```

---

biofam

*Family life states from the Swiss Household Panel biographical survey*

---

### Description

2000 16 year-long family life sequences built from the retrospective biographical survey carried out by the Swiss Household Panel (SHP) in 2002.

### Usage

```
data(biofam)
```

### Format

A data frame with 2000 rows, 16 state variables, 1 id variable and 7 covariates and 2 weights variables.

### Details

The *biofam* data set was constructed by Müller et al. (2007) from the data of the retrospective biographical survey carried out by the Swiss Household Panel (SHP) in 2002.

The data set contains (in columns 10 to 25) sequences of family life states from age 15 to 30 (sequence length is 16) and a series of covariates. The sequences are a sample of 2000 sequences of those created from the SHP biographical survey. It includes only individuals who were at least 30 years old at the time of the survey. The *biofam* data set describes family life courses of 2000 individuals born between 1909 and 1972.

The states numbered from 0 to 7 are defined from the combination of five basic states, namely Living with parents (Parent), Left home (Left), Married (Marr), Having Children (Child), Divorced:

0 = "Parent"  
 1 = "Left"  
 2 = "Married"  
 3 = "Left+Marr"  
 4 = "Child"  
 5 = "Left+Child"  
 6 = "Left+Marr+Child"  
 7 = "Divorced"

The covariates are:

sex	
birthyr	(birth year)
nat_1_02	(first nationality)
plingu02	(language of questionnaire)
p02r01	(religion)
p02r04	(religious participation)
cspfaj	(father's social status)
cspmoj	(mother's social status)

Two additional weights variables are inserted for illustrative purpose ONLY (since biofam is a sub-sample of the original data, these weights are not adapted to the actual data):

wp00tbgp	(weights inflating to the Swiss population)
wp00tbgs	(weights respecting sample size)

## Source

Swiss Household Panel <https://forscenter.ch/projects/swiss-household-panel/>

## References

Mueller, N. S., Studer, M. and Ritschard, G. (2007). Classification de parcours de vie a l'aide de l'optimal matching. In *XIVe Rencontre de la Societe francophone de classification (SFC 2007)*, Paris, 5-7 septembre 2007, pp. 157-160.

## Examples

```

data(biofam, package="MEDseq")

biofam      <- list(covariates = biofam[2L:9L], sequences = biofam[10L:25L] + 1L)
biofam.cov  <- biofam$covariates[,colSums(is.na(biofam$covariates)) == 0]
biofam.seq  <- seqdef(biofam$sequences,
```

```

states = c("P", "L", "M", "L+M", "C", "L+C", "L+M+C", "D"),
labels = c("Parent", "Left", "Married", "Left+Marr", "Child",
           "Left+Child", "Left+Marr+Child", "Divorced")
biofam.cov$age <- 2002 - biofam.cov$birthyr

```

dbs

*Compute the Density-based Silhouette***Description**

Computes the Density-based Silhouette for a 'soft' clustering assignment matrix.

**Usage**

```

dbs(z,
    ztol = 1E-100,
    weights = NULL,
    summ = c("mean", "median"),
    clusters = NULL,
    ...)

```

**Arguments**

<code>z</code>	A numeric matrix such that rows correspond to observations, columns correspond to clusters, and rows sum to 1.
<code>ztol</code>	A small (single, numeric, non-negative) tolerance parameter governing whether small assignment probabilities are treated instead as crisp assignments. Defaults to 1E-100.
<code>weights</code>	An optional numeric vector giving observation-specific weights for computing the (weighted) mean/median DBS (see <code>summ</code> ).
<code>summ</code>	A single character string indicating whether the (possibly weighted) "mean" (the default) or "median" DBS should be computed.
<code>clusters</code>	Optional/experimental argument for giving the indicator labels of the cluster assignments. Defaults to the MAP assignment derived from <code>z</code> when not supplied. Note that actually supplying the MAP assignment here is slightly less efficient than the NULL default and <b>not</b> advised.
<code>...</code>	Catches unused arguments.

**Value**

A list with the following elements:

`silvals` A matrix where each row contains the cluster to which each observation belongs in the first column and the observation-specific DBS width in the second column.

`msw` Depending on the value of `summ`, either the mean or median DBS width.

`wmsw` Depending on the value of `summ`, either the weighted mean or weighted median DBS width.

**Note**

When calling `MEDseq_fit`, the `summ` argument can be passed via the `...` construct, in which case it governs both the `dbs` and `asw` criteria.

**Author(s)**

Keefe Murphy - <<keefe.murphy@mu.ie>>

**References**

Menardi, G. (2011). Density-based Silhouette diagnostics for clustering methods. *Statistics and Computing*, 21(3): 295-308.

**See Also**

[MEDseq\\_fit](#)

**Examples**

```
# Generate a toy z matrix
z <- abs(matrix(rnorm(50), ncol=2))
z <- z/rowSums(z)

# Return the median DBS width
dbs(z, summ="median")$msw

# For real sequence data
data(mvad)

mod <- MEDseq_fit(seqdef(mvad[,17:86]), G=11, modtype="UUN", weights=mvad$weight)

dbs(mod$z, weights=mvad$weight)
```

---

`get_MEDseq_results`      *Extract results from a MEDseq model*

---

**Description**

Utility function for extracting results of submodels from "MEDseq" objects when a range of models were run via `MEDseq_fit`.

**Usage**

```
get_MEDseq_results(x,
  what = c("z", "MAP", "DBS", "ASW"),
  rank = 1L,
  criterion = c("bic", "icl", "aic", "dbs"),
```

```

        "asw", "cv", "nec", "loglik"),
    G = NULL,
    modtype = NULL,
    noise = TRUE,
    ...)

```

### Arguments

x	An object of class "MEDseq" generated by <a href="#">MEDseq_fit</a> or an object of class "MEDseqCompare" generated by <a href="#">MEDseq_compare</a> .
what	A character string indicating the desired results to extract.
rank	A number indicating what rank model results should be extracted from, where the rank is determined by <code>criterion</code> . Defaults to 1, i.e. the best model.
criterion	The <code>criterion</code> used to determine the ranking. Defaults to "bic".
G	Optional argument giving the number of components in the model for which results are desired. Can be supplied with or without also specifying <code>modtype</code> .
modtype	Optional argument giving the desired model type for which results are desired. Can be supplied with or without also specifying <code>G</code> .
noise	A logical indicating whether models with a noise component should be considered. Defaults to TRUE.
...	Catches unused arguments.

### Details

The arguments `rank` and `criterion` are invoked when one or more of the arguments `G` and `modtype` are missing. Thus, supplying `G` and `modtype` allows `rank` and `criterion` to be bypassed entirely.

### Value

The desired results extracted from the MEDseq model.

### Note

Arguments to this function can be supplied to [plot.MEDseq](#) via the `...` construct.

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### See Also

[MEDseq\\_fit](#), [plot.MEDseq](#)



## Examples

```

data(biofam)
# mod <- MEDseq_fit(seqdef(biofam[10:25] + 1L), G=9:10)

# Extract the MAP clustering of the best 9-cluster model according to the asw criterion
# get_MEDseq_results(mod, what="MAP", G=9, criterion="asw")

# Extract the DBS values of the best UUN model according to the dbs criterion
# get_MEDseq_results(mod, what="DBS", modtype="UUN", criterion="dbs")

# Plot the DBS values of this same model, by passing get_MEDseq_results arguments through plot
# plot(mod, type="dbsvals", modtype="UUN", criterion="dbs")

```

---

MEDseq_clustnames	<i>Automatic labelling of clusters using central sequences</i>
-------------------	--

---

## Description

These functions extract names for clusters according to the SPS representation of their central sequences.

## Usage

```

MEDseq_clustnames(x,
                  cluster = TRUE,
                  size = FALSE,
                  ...)

MEDseq_nameclusts(names)

```

## Arguments

x	An object of class "MEDseq" generated by <a href="#">MEDseq_fit</a> or an object of class "MEDseqCompare" generated by <a href="#">MEDseq_compare</a> .
cluster	A logical indicating whether names should be prepended with the text "Cluster g: ", where g is the cluster number. Defaults to TRUE.
size	A logical indicating whether the (typically 'soft') size of each cluster is appended to the label of each group, expressed as a percentage of the total number of observations. Defaults to FALSE.
...	Catches unused arguments.
names	The output of MEDseq_clustnames to be passed to the convenience function MEDseq_nameclusts (see Details).

## Details

Unlike the `seqclustname` function from the **WeightedCluster** package which inspired these functions, `MEDseq_clustnames` only returns the names themselves, not the factor variable indicating cluster membership with labels given by those names. Thus, `MEDseq_nameclusts` is provided as a convenience function for precisely this purpose (see Examples).

## Value

For `MEDseq_clustnames`, a character vector containing the names for each component defined by their central sequence, and optionally the cluster name (see `cluster` above) and cluster size (see `size` above). The name for the noise component, if any, will always be simply "Noise" (or "Cluster 0: Noise").

For `MEDseq_nameclusts`, a factor version of `x$MAP` with levels given by the output of `MEDseq_clustnames`.

## Note

The main `MEDseq_clustnames` function is used internally by `plot.MEDseq`, `MEDseq_meantime`, `MEDseq_stderr`, and also other print and summary methods, where its invocation can typically be controlled via a SPS logical argument. However, the optional arguments `cluster` and `size` can only be passed through `plot.MEDseq`; elsewhere `cluster=TRUE` and `size=FALSE` are always assumed.

## Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

## References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

## See Also

`seqformat`, `seqclustname`, `plot.MEDseq`, `MEDseq_meantime`, `MEDseq_stderr`

## Examples

```
# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels <- c("Employment", "Further Education", "Higher Education",
```

```

                                "Joblessness", "School", "Training")
mvad.seq      <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a model with weights and a gating covariate
# Have the probability of noise-component membership depend on the covariate
mod  <- MEDseq_fit(mvad.seq, G=5, modtype="UUN", weights=mvad$weights,
                  gating=~ gcse5eq, covars=mvad.cov, noise.gate=TRUE)

# Extract the names
names <- MEDseq_clustnames(mod, cluster=FALSE, size=TRUE)

# Get the renamed MAP cluster membership indicator vector
group <- MEDseq_nameclusts(names)

# Use the output in plots
seqplot(mvad.seq, type="d", group=group)
# same as:
# plot(mod, type="d", soft=FALSE, weighted=FALSE, cluster=FALSE, size=TRUE, border=TRUE)

# Indeed, this function is invoked by default for certain plot types
plot(mod, type="d", soft=TRUE, weighted=TRUE)
plot(mod, type="d", soft=TRUE, weighted=TRUE, SPS=FALSE)

# Invoke this function when printing the gating network coefficients
print(mod$gating, SPS=FALSE)
print(mod$gating, SPS=TRUE)

# Invoke this function in a call to MEDseq_meantime
MEDseq_meantime(mod, SPS=TRUE)

# Invoke this function in other plots
plot(mod, type="clusters", SPS=TRUE)
plot(mod, type="precision", SPS=TRUE)

```

---

MEDseq\_compare

*Choose the best MEDseq model*


---

## Description

Takes one or more sets of "MEDseq" models fitted by `MEDseq_fit` and ranks them according to a specified model selection criterion. It's possible to respect the internal ranking within each set of models, or to discard models within each set which were already deemed sub-optimal. This function can help with model selection via exhaustive or stepwise searches.

## Usage

```

MEDseq_compare(...,
               criterion = c("bic", "icl", "aic",
                           "dbs", "asw", "cv", "nec"),
               pick = 10L,

```

```

        optimal.only = FALSE)

## S3 method for class 'MEDseqCompare'
print(x,
      index = seq_len(x$pick),
      rerank = FALSE,
      digits = 3L,
      maxi = length(index),
      ...)

```

## Arguments

... One or more objects of class "MEDseq" outputted by [MEDseq\\_fit](#). All models must have been fit to the same data set. A single *named* list of such objects can also be supplied. Additionally, objects of class "MEDseqCompare" outputted by this very function can also be supplied here. This argument is only relevant for the [MEDseq\\_compare](#) function and will be ignored for the associated print function.

criterion The criterion used to determine the ranking. Defaults to "bic".

pick The (integer) number of models to be ranked and compared. Defaults to 10L. Will be constrained by the number of models within the "MEDseq" objects supplied via ... if `optimal.only` is FALSE, otherwise constrained simply by the number of "MEDseq" objects supplied. Setting `pick=Inf` is a valid way to select all models.

optimal.only Logical indicating whether to only rank models already deemed optimal within each "MEDseq" object (TRUE), or to allow models which were deemed suboptimal enter the final ranking (FALSE, the default). See Details.

x, index, rerank, digits, maxi Arguments required for the associated print function:

- x An object of class "MEDseqCompare" resulting from a call to [MEDseq\\_compare](#).
- index A logical or numeric vector giving the indices of the rows of the table of ranked models to print. This defaults to the full set of ranked models. It can be useful when the table of ranked models is large to examine a subset via this index argument, for display purposes. See `rerank`.
- rerank A logical indicating whether the ranks should be recomputed when sub-setting using `index`. Defaults to FALSE.
- digits The number of decimal places to round model selection criteria to (defaults to 3).
- maxi A number specifying the maximum number of rows/models to print. Defaults to `length(index)`.

## Details

The purpose of this function is to conduct model selection on "MEDseq" objects, fit to the same data set, with different combinations of gating network covariates or different initialisation settings.

Model selection will have already been performed in terms of choosing the optimal number of components and MEDseq model type within each supplied set of results, but [MEDseq\\_compare](#) will

respect the internal ranking of models when producing the final ranking if `optimal.only` is `FALSE`: otherwise only those models already deemed optimal within each "MEDseq" object will be ranked.

As such if two sets of results are supplied when `optimal.only` is `FALSE`, the 1st, 2nd, and 3rd best models could all belong to the first set of results, meaning a model deemed suboptimal according to one set of covariates could be superior to one deemed optimal under another set of covariates.

## Value

A list of class "MEDseqCompare", for which a dedicated print function exists, containing the following elements (each of length `pick`, and ranked according to `criterion`, where appropriate):

<code>data</code>	The name of the data set to which the models were fitted.
<code>optimal</code>	The single optimal model (an object of class "MEDseq") among those supplied, according to the chosen criterion.
<code>pick</code>	The final number of ranked models. May be different (i.e. less than) the supplied <code>pick</code> value.
<code>MEDNames</code>	The names of the supplied "MEDseq" objects.
<code>modelNames</code>	The MEDseq model names (denoting the constraints or lack thereof on the precision parameters).
<code>G</code>	The optimal numbers of components.
<code>df</code>	The numbers of estimated parameters.
<code>iters</code>	The numbers of EM/CEM iterations.
<code>bic</code>	BIC values, ranked according to <code>criterion</code> .
<code>icl</code>	ICL values, ranked according to <code>criterion</code> .
<code>aic</code>	AIC values, ranked according to <code>criterion</code> .
<code>dbS</code>	(Weighted) mean/median DBS values, ranked according to <code>criterion</code> .
<code>asw</code>	(Weighted) mean/median ASW values, ranked according to <code>criterion</code> .
<code>cv</code>	Cross-validated log-likelihood values, ranked according to <code>criterion</code> .
<code>nec</code>	NEC values, ranked according to <code>criterion</code> .
<code>loglik</code>	Maximal log-likelihood values, ranked according to <code>criterion</code> .
<code>gating</code>	The gating formulas.
<code>algo</code>	The algorithm used for fitting the model - either "EM", "CEM", "cemEM".
<code>equalPro</code>	Logical indicating whether mixing proportions were constrained to be equal across components.
<code>opti</code>	The method used for estimating the central sequence(s).
<code>weights</code>	Logical indicating whether the given model was fitted with sampling weights.
<code>noise</code>	Logical indicating the presence/absence of a noise component. Only displayed if at least one of the compared models has a noise component.
<code>noise.gate</code>	Logical indicating whether gating covariates were allowed to influence the noise component's mixing proportion. Only printed for models with a noise component, when at least one of the compared models has gating covariates.
<code>equalNoise</code>	Logical indicating whether the mixing proportion of the noise component for <code>equalPro</code> models is also equal ( <code>TRUE</code> ) or estimated ( <code>FALSE</code> ).

**Note**

The criterion argument here need not comply with the criterion used for model selection within each "MEDseq" object, but be aware that a mismatch in terms of `criterion` may require the optimal model to be re-fit in order to be extracted, thereby slowing down `MEDseq_compare`.

If random starts had been used via `init.z="random"` the optimal model may not necessarily correspond to the highest-ranking model in the presence of a criterion mismatch, due to the randomness of the initialisation.

A dedicated print function exists for objects of class "MEDseqCompare" and `plot.MEDseq` can also be called on objects of class "MEDseqCompare".

**Author(s)**

Keefe Murphy - <<keefe.murphy@mu.ie>>

**References**

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

**See Also**

`MEDseq_fit`, `plot.MEDseq`

**Examples**

```
data(biofam)
seqs <- seqdef(biofam[10:25] + 1L,
              states = c("P", "L", "M", "L+M", "C",
                        "L+C", "L+M+C", "D"))
covs <- cbind(biofam[2:3], age=2002 - biofam$birthyr)

# Fit a range of models
# m1 <- MEDseq_fit(seqs, G=9:10)
# m2 <- MEDseq_fit(seqs, G=9:10, gating=~sex, covars=covs, noise.gate=FALSE)
# m3 <- MEDseq_fit(seqs, G=9:10, gating=~age, covars=covs, noise.gate=FALSE)
# m4 <- MEDseq_fit(seqs, G=9:10, gating=~sex + age, covars=covs, noise.gate=FALSE)

# Rank only the optimal models (according to the dbs criterion)
# Examine the best model in more detail
# (comp <- MEDseq_compare(m1, m2, m3, m4, criterion="dbs", optimal.only=TRUE))
# (best <- comp$optimal)
# (summ <- summary(best, parameters=TRUE))

# Examine all models visited, including those already deemed suboptimal
# Only print models with gating covariates & 10 components
# comp2 <- MEDseq_compare(comp, m1, m2, m3, m4, criterion="dbs", pick=Inf)
# print(comp2, index=comp2$gating != "None" & comp2$G == 10)
```

---

MEDseq\_control                      *Set control values for use with MEDseq\_fit*

---

### Description

Supplies a list of arguments (with defaults) for use with [MEDseq\\_fit](#).

### Usage

```
MEDseq_control(algo = c("EM", "CEM", "cemEM"),
               init.z = c("kmedoids", "kmodes", "kmodes2", "hc", "random", "list"),
               z.list = NULL,
               dist.mat = NULL,
               unique = TRUE,
               criterion = c("bic", "icl", "aic", "dbs", "asw", "cv", "nec"),
               tau0 = NULL,
               noise.gate = TRUE,
               random = TRUE,
               do.cv = FALSE,
               do.nec = FALSE,
               nfolds = 10L,
               nstarts = 1L,
               stopping = c("aitken", "relative"),
               equalPro = FALSE,
               equalNoise = FALSE,
               tol = c(1E-05, 1E-08),
               itmax = c(.Machine$integer.max, 1000L),
               opti = c("mode", "medoid", "first", "GA"),
               ordering = c("none", "decreasing", "increasing"),
               MaxNWts = 1000L,
               verbose = TRUE,
               ...)
```

### Arguments

- |        |   |
|--------|---|
| algo   | Switch controlling whether models are fit using the "EM" (the default) or "CEM" algorithm. The option "cemEM" allows running the EM algorithm starting from convergence of the CEM algorithm.   |
| init.z | The method used to initialise the cluster labels. All options respect the presence of sampling weights, if any. Defaults to "kmedoids". Other options include "kmodes", "kmodes2", Ward's hierarchical clustering ("hc", via <a href="#">hclust</a> ), "random" initialisation, and a user-supplied "list" (see z.list below). For weighted sequences, "kmedoids" is itself initialised using Ward's hierarchical clustering.<br><br>The "kmodes" and "kmodes2" options both internally call the function <a href="#">wkModes</a> , which <i>typically</i> uses random initial modes. Under "kmodes", the algorithm is instead initialised via the medoids of the clusters obtained from a call to <a href="#">hclust</a> . |

	<p>The option "kmodes2" is slightly faster, by virtue of using the <i>random</i> initial medoids. However, final results are by default still subject to randomness under both options (unless <code>set.seed</code> is invoked), as ties for modes and cluster assignments are <i>typically</i> broken at random throughout the algorithm (see the random argument below, and in <code>wkModes</code> itself).</p>
<code>z.list</code>	<p>A user supplied list of initial cluster allocation matrices, with number of rows given by the number of observations, and numbers of columns given by the range of component numbers being considered. Only relevant if <code>init.z == "z.list"</code>. These matrices are allowed correspond to both soft or hard clusterings, and will be internally normalised so that the rows sum to 1.</p>
<code>dist.mat</code>	<p>An optional distance matrix to use for initialisation when <code>init.z</code> is one of "kmedoids" or "hc". Defaults to a Hamming distance matrix. This is an experimental feature and should only be tampered with by expert users.</p>
<code>unique</code>	<p>A logical indicating whether the model is fit only to the unique observations (defaults to TRUE). When there are covariates, this means all unique combinations of covariate and sequence patterns, otherwise only the sequence patterns.</p> <p>When weights <i>are not</i> supplied to <code>MEDseq_fit</code> and <code>isTRUE(unique)</code>, weights are given by the occurrence frequency of the corresponding sequences, and the model is then fit to the unique observations only.</p> <p>When weights <i>are</i> supplied and <code>isTRUE(unique)</code>, the weights are summed for each set of duplicate observations and assigned to one retained copy of each corresponding unique sequence. Hence, observations with different weights that are otherwise duplicates are treated as duplicates and significant computational gains can be made.</p> <p>In both cases, the results will be unchanged, but setting <code>unique</code> to TRUE can often be much faster.</p>
<code>criterion</code>	<p>When either <code>G</code> or <code>modtype</code> is a vector, <code>criterion</code> governs how the 'best' model is determined when gathering output. Defaults to "bic". Note that all criteria will be returned in any case, if possible.</p>
<code>tau0</code>	<p>Prior mixing proportion for the noise component. If supplied, a noise component will be added to the model in the estimation, with <code>tau0</code> giving the prior probability of belonging to the noise component for <i>all</i> observations. Typically supplied as a scalar in the interval (0, 1), e.g. 0.1. Can be supplied as a vector when gating covariates are present and <code>noise.gate</code> is TRUE.</p>
<code>noise.gate</code>	<p>A logical indicating whether gating network covariates influence the mixing proportion for the noise component, if any. Defaults to TRUE, but leads to greater parsimony if FALSE. Only relevant in the presence of a noise component (i.e. the "CCN", "UCN", "CUN", and "UUN" models); only affects estimation in the presence of gating covariates.</p>
<code>random</code>	<p>A logical governing how ties for estimated central sequence positions are handled. When TRUE (the default), such ties are broken at random. When FALSE (the implied default prior to version 1.2.0 of this package), the first candidate state is always chosen. This argument affects all <code>opt i</code> options. If <code>verbose</code> is TRUE and there are tie-breaking operations performed, a warning message is printed once per model, regardless of the number of such operations.</p>



Note that this argument is *also* passed to `wkModes` if `init.z` is "kmodes" or "kmodes2" and that, in certain rare cases when the "CEM" algo is invoked when `equalPro` is TRUE and the precision parameter(s) are somehow constrained across clusters, this argument also governs ties for cluster assignments within `MEDseq_fit` as well.

<code>do.cv</code>	A logical indicating whether cross-validated log-likelihood scores should also be computed (see <code>nfolds</code> ). Defaults to FALSE due to significant computational burden incurred.
<code>do.nec</code>	A logical indicating whether the normalised entropy criterion (NEC) should also be computed (for models with more than one component). Defaults to FALSE. When TRUE, models with <code>G=1</code> are fitted always.
<code>nfolds</code>	The number of folds to use when <code>isTRUE{do.cv}</code> .
<code>nstarts</code>	The number of random initialisations to use when <code>init.z="random"</code> . Defaults to 1. Results will be based on the random start yielding the highest estimated log-likelihood.
<code>stopping</code>	The criterion used to assess convergence of the EM/CEM algorithm. The default ("aitken") uses Aitken's acceleration method, otherwise the "relative" change in log-likelihood is monitored (which may be less strict).
<code>equalPro</code>	Logical variable indicating whether or not the mixing proportions are to be constrained to be equal in the model. Default: <code>equalPro = FALSE</code> . Only relevant when gating covariates are <i>not</i> supplied within <code>MEDseq_fit</code> , otherwise ignored. In the presence of a noise component, only the mixing proportions for the non-noise components are constrained to be equal (by default, see <code>equalNoise</code> ), after accounting for the noise component.
<code>equalNoise</code>	Logical which is <b>only</b> invoked when <code>isTRUE(equalPro)</code> and gating covariates are not supplied. Under the default setting (FALSE), the mixing proportion for the noise component is estimated, and remaining mixing proportions are equal; when TRUE all components, including the noise component, have equal mixing proportions.
<code>tol</code>	A vector of length two giving <i>relative</i> convergence tolerances for 1) the log-likelihood of the EM/CEM algorithm, and 2) optimisation in the multinomial logistic regression in the gating network, respectively. The default is <code>c(1e-05, 1e-08)</code> . If only one number is supplied, it is used as the tolerance in both cases.
<code>itmax</code>	A vector of length two giving integer limits on the number of iterations for 1) the EM/CEM algorithm, and 2) the multinomial logistic regression in the gating network, respectively. The default is <code>c(.Machine\$integer.max, 1000)</code> . This allows termination of the EM/CEM algorithm to be completely governed by <code>tol[1]</code> . If only one number is supplied, it is used as the iteration limit for the EM/CEM algorithm only and the other element of <code>itmax</code> retains its usual default.  If, for any model with gating covariates, the multinomial logistic regression in the gating network fails to converge in <code>itmax[2]</code> iterations at any stage of the EM/CEM algorithm, an appropriate warning will be printed, prompting the user to modify this argument.
<code>opti</code>	Character string indicating how central sequence parameters should be estimated. The default "mode" is exact and thus this experimental argument should

	only be tampered with by expert users. The option "medoid" fixes the central sequence(s) to be one of the observed sequences (like k-medoids). The other options "first" and "GA" use the first-improvement and genetic algorithms, respectively, to mutate the medoid. Pre-computation of the Hamming distance matrix for the observed sequences speeds-up computation of all options other than "mode".
ordering	Experimental feature that should only be tampered with by experienced users. Allows sequences to be reordered on the basis of the column-wise entropy when opti is "first" or "GA".
MaxNWts	The maximum allowable number of weights in the call to <code>multinom</code> for the multinomial logistic regression in the gating network. There is no intrinsic limit in the code, but increasing MaxNWts will probably allow fits that are very slow and time-consuming. It may be necessary to increase MaxNWts when categorical concomitant variables with many levels are included or the number of components is high.
verbose	Logical indicating whether to print messages pertaining to progress to the screen during fitting. By default is TRUE if the session is interactive, and FALSE otherwise. If FALSE, warnings and error messages will still be printed to the screen, but everything else will be suppressed.
...	Catches unused arguments, and also allows the optional arguments <code>ztol</code> and <code>summ</code> to be passed to <code>dfs</code> ( <code>ztol</code> and <code>summ</code> ) as well as the ASW computation ( <code>summ</code> ), and the optional <code>wkModes</code> arguments <code>iter.max</code> , <code>freq.weighted</code> , and <code>fast</code> (provided <code>init.z</code> is one of "kmodes" or "kmodes2"). In such cases, the <code>wkModes</code> argument <code>random</code> is already controlled by <code>random</code> above here.

## Details

`MEDseq_control` is provided for assigning values and defaults within `MEDseq_fit`. While the `criterion` argument controls the choice of the optimal number of components and MEDseq model type (in terms of the constraints or lack thereof on the precision parameters), `MEDseq_compare` is provided for choosing between fits with different combinations of covariates or different initialisation settings.

## Value

A named list in which the names are the names of the arguments and the values are the values supplied to the arguments.

## Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

## References

- Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.
- Menardi, G. (2011). Density-based Silhouette diagnostics for clustering methods. *Statistics and Computing*, 21(3): 295-308.

**See Also**

[MEDseq\\_fit](#), [dbs](#), [wckMedoids](#), [pam](#), [wKModes](#), [hclust](#), [seqdist](#), [multinom](#), [MEDseq\\_compare](#)

**Examples**

```
# The CC MEDseq model is almost equivalent to k-medoids when the
# CEM algorithm is employed, mixing proportions are constrained,
# and the central sequences are restricted to the observed sequences
ctrl <- MEDseq_control(algo="CEM", equalPro=TRUE, opti="medoid", criterion="asw")

data(mvad)
# Note that ctrl must be explicitly named 'ctrl'
mod <- MEDseq_fit(seqdef(mvad[,17:86]), G=11, modtype="CC", weights=mvad$weight, ctrl=ctrl)

# Alternatively, specify the control arguments directly
mod <- MEDseq_fit(seqdef(mvad[,17:86]), G=11, modtype="CC", weights=mvad$weight,
                  algo="CEM", equalPro=TRUE, opti="medoid", criterion="asw")

# Note that supplying control arguments via a mix of the ... construct and the named argument
# 'control' or supplying MEDseq_control output without naming it 'control' can throw an error
```

---

MEDseq\_entropy

*Entropy of a fitted MEDseq model*

---

**Description**

Calculates the normalised entropy of a fitted MEDseq model.

**Usage**

```
MEDseq_entropy(x)
```

**Arguments**

x An object of class "MEDseq" generated by [MEDseq\\_fit](#) or an object of class "MEDseqCompare" generated by [MEDseq\\_compare](#).

**Details**

This function calculates the normalised entropy via

$$H = -\frac{1}{n \log(G)} \sum_{i=1}^n \sum_{g=1}^G \hat{z}_{ig} \log(\hat{z}_{ig}),$$

where  $n$  and  $G$  are the sample size and number of components, respectively, and  $\hat{z}_{ig}$  is the estimated posterior probability at convergence that observation  $i$  belongs to component  $g$ .

**Value**

A single number, given by  $1 - H$ , in the range  $[0,1]$ , such that *larger* values indicate clearer separation of the clusters.

**Note**

This function will always return a normalised entropy of 1 for models fitted using the "CEM" algorithm (see [MEDseq\\_control](#)), or models with only one component.

**Author(s)**

Keefe Murphy - <<keefe.murphy@mu.ie>>

**References**

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

**See Also**

[MEDseq\\_fit](#), [MEDseq\\_control](#)

**Examples**

```
# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels <- c("Employment", "Further Education", "Higher Education",
  "Joblessness", "School", "Training")
mvad.seq <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a model with weights and a gating covariate
# Have the probability of noise-component membership be constant
mod <- MEDseq_fit(mvad.seq, G=11, modtype="UUN", weights=mvad$weights,
  gating=~ gcse5eq, covars=mvad.cov, noise.gate=FALSE)

# Calculate the normalised entropy
MEDseq_entropy(mod)
```

MEDseq\_fit

*MEDseq: Mixtures of Exponential-Distance Models with Covariates***Description**

Fits MEDseq models: mixtures of Exponential-Distance models with gating covariates and sampling weights. Typically used for clustering categorical/longitudinal life-course sequences. Additional arguments are available via the function `MEDseq_control`.

**Usage**

```
MEDseq_fit(seqs,
           G = 1L:9L,
           modtype = c("CC", "UC", "CU", "UU",
                      "CCN", "UCN", "CUN", "UUN"),
           gating = NULL,
           weights = NULL,
           ctrl = MEDseq_control(...),
           covars = NULL,
           ...)
```

```
## S3 method for class 'MEDseq'
summary(object,
        classification = TRUE,
        parameters = FALSE,
        network = FALSE,
        SPS = FALSE,
        ...)
```

```
## S3 method for class 'MEDseq'
print(x,
      digits = 3L,
      ...)
```

**Arguments**

<code>seqs</code>	A state-sequence object of class "stslst" as created by the <code>seqdef</code> function in the <b>TraMineR</b> package. Note that the data set must have equal sequence lengths, the intervals are assumed to be evenly spaced, and missingness is not allowed.
<code>G</code>	A positive integer vector specifying the numbers of mixture components (clusters) to fit. Defaults to <code>G=1:9</code> .
<code>modtype</code>	A vector of character strings indicating the type of MEDseq models to be fitted, in terms of the constraints or lack thereof on the precision parameters. By default, all valid model types are fitted (except some only where $G > 1$ or $G > 2$ , see Note). The models are named "CC", "CU", "UC", "UU", "CCN", "CUN", "UCN",

and "UUN". The first letter denotes whether the precision parameters are constrained/unconstrained across clusters. The second letter denotes whether the precision parameters are constrained/unconstrained across sequence positions (i.e. time points). The third letter denotes whether one of the components is constrained to have zero-precision/infinite variance. Such a noise component assumes sequences in that cluster follow a uniform distribution.

gating	A <a href="#">formula</a> for determining the model matrix for the multinomial logistic regression in the gating network when fixed covariates enter the mixing proportions. Defaults to ~1, i.e. no covariates. This will be ignored where G=1. Continuous, categorical, and/or ordinal covariates are allowed. Logical covariates will be coerced to factors. Interactions, transformations, and higher order terms are permitted: the latter <b>must</b> be specified explicitly using the <code>AsIs</code> operator ( <code>I</code> ). The specification of the LHS of the formula is ignored. Intercept terms are included by default.
weights	Optional numeric vector containing observation-specific sampling weights, which are accounted for in the model fitting and other functions where applicable. weights are always internally normalised to sum to the sample size. See the unique argument to <code>MEDseq_control</code> to see how incorporating weights also yields computational benefits. Note that weights must <b>always</b> be explicitly supplied here; it is not enough to use weights when constructing the state sequence object via <code>seqdef</code> . If you <i>are</i> using a weighted "stslst" state sequence object and do not specify weights, you will be prompted to explicitly specify <code>weights=attr(seqs, "weights")</code> for a weighted model or <code>weights=NULL</code> for an unweighted model.
ctrl	A list of control parameters for the EM/CEM and other aspects of the algorithm. The defaults are set by a call to <code>MEDseq_control</code> .
covars	An optional data frame (or a matrix with named columns) in which to look for the covariates in the gating network formula, if any. If not found in covars, any supplied gating covariates are taken from the environment from which <code>MEDseq_fit</code> is called. Try to ensure the names of variables in covars do not match any of those in seqs.
...	Catches unused arguments (see <code>MEDseq_control</code> ).
x, object, digits, classification, parameters, network, SPS	Arguments required for the print and summary functions: x and object are objects of class "MEDseq" resulting from a call to <code>MEDseq_fit</code> , while digits gives the number of decimal places to round to for printing purposes (defaults to 3). classification, parameters, and network are logicals which govern whether a table of the MAP classification of observations, the mixture component parameters, and the gating network coefficients are printed, respectively. SPS governs the printing of the relevant quantities in "summaryMEDseq" objects when any of classification, parameters, &/or network are TRUE (see <code>MEDseq_clustnames</code> and <code>seqformat</code> ).

## Details

The function effectively allows 8 different MEDseq precision parameter settings for models with or without gating network covariates. By constraining the mixing proportions to be equal (see `equalPro` in `MEDseq_control`) an extra special case is facilitated in the latter case.

While model selection in terms of choosing the optimal number of components and the MEDseq model type is performed within `MEDseq_fit`, using one of the criterion options within `MEDseq_control`, choosing between multiple fits with different combinations of covariates or different initialisation settings can be done by supplying objects of class "MEDseq" to `MEDseq_compare`.

## Value

A list (of class "MEDseq") with the following named entries (of which some may be missing, depending on the criterion employed), mostly corresponding to the chosen optimal model (as determined by the criterion within `MEDseq_control`):

<code>call</code>	The matched call.
<code>data</code>	The input data, seqs.
<code>modtype</code>	A character string denoting the MEDseq model type at which the optimal criterion occurs.
<code>G</code>	The optimal number of mixture components according to criterion.
<code>params</code>	A list with the following named components: <ul style="list-style-type: none"> <li><code>theta</code> A matrix with <math>G</math> rows and <math>T</math> columns, where <math>T</math> is the number of sequence positions, giving the central sequences of each cluster. The mean of the noise component is not reported, as it does not contribute in any way to the likelihood. A dedicated <code>print</code> function is provided.</li> <li><code>lambda</code> A matrix of precision parameters. Will contain 1 row if the 1st letter of <code>modtype</code> is "C" and <math>G</math> columns otherwise. Will contain 1 column if the 2nd letter of <code>modtype</code> is "C" and <math>T</math> columns otherwise, where <math>T</math> is the number of sequence positions. Precision parameter values of zero are reported for the noise component, if any. Note that values of <code>Inf</code> are also possible, corresponding to zero-variance, which is most likely under the "UU" or "UUN" models. A dedicated <code>print</code> function is provided.</li> <li><code>tau</code> The mixing proportions: either a vector of length <math>G</math> or, if gating covariates were supplied, a matrix with an entry for each observation (rows) and component (columns).</li> </ul>
<code>gating</code>	An object of class "MEDgating" (for which dedicated <code>print</code> , <code>summary</code> , and <code>predict</code> methods exist) and either "multinom" or "glm" (only for single-component models) giving the <code>multinom</code> regression coefficients of the gating network. If gating covariates were <i>NOT</i> supplied (or the best model has just one component), this corresponds to a RHS of $\sim 1$ , otherwise the supplied gating formula. As such, a fitted gating network is always returned even in the absence of supplied covariates or clusters. If there is a noise component (and the option <code>noise.gate=TRUE</code> is invoked), its coefficients are those for the <i>last</i> component. <b>Users are cautioned against making inferences about statistical significance from summaries of the coefficients in the gating network. Users are instead advised to use the function <code>MEDseq_stderr</code>.</b>
<code>z</code>	The final responsibility matrix whose $[i, k]$ -th entry is the probability that observation $i$ belongs to the $k$ -th component. If there is a noise component, its values are found in the <i>last</i> column.

MAP	The vector of cluster labels for the chosen model corresponding to $z$ , i.e. $\max.col(z)$ . Observations belonging to the noise component, if any, will belong to component $\emptyset$ .
BIC	A matrix of <i>all</i> BIC values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
ICL	A matrix of <i>all</i> ICL values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
AIC	A matrix of <i>all</i> AIC values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
DBS	A matrix of <i>all</i> (weighted) mean/median DBS values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note and <a href="#">dbs</a> .
DBSvals	A list of lists giving the observation-specific DBS values for <i>all</i> fitted models. The first level of the list corresponds to numbers of components, the second to the MEDseq model types.
dbs	The (weighted) mean/median DBS value corresponding to the optimal model. May not necessarily be the optimal DBS.
dbsvals	Observation-specific DBS values corresponding to the optimum model, which may not be optimal in terms of DBS.
ASW	A matrix of <i>all</i> (weighted) mean/median ASW values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
ASWvals	A list of lists giving the observation-specific ASW values for <i>all</i> fitted models. The first level of the list corresponds to numbers of components, the second to the MEDseq model types.
asw	The (weighted) mean/median ASW value corresponding to the optimal model. May not necessarily be the optimal ASW.
aswvals	Observation-specific ASW values corresponding to the optimum model, which may not be optimal in terms of ASW.
LOGLIK	A matrix of <i>all</i> maximal log-likelihood values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
DF	A matrix giving the numbers of estimated parameters (i.e. the number of 'used' degrees of freedom) for <i>all</i> visited models, with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. Subtract these numbers from the sample size to get the degrees of freedom. See Note.
ITERS	A matrix giving the total number of EM/CEM iterations for <i>all</i> visited models, with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns. See Note.
CV	A matrix of <i>all</i> cross-validated log-likelihood values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns, if available. See Note and the arguments <code>do.cv</code> and <code>nolds</code> to <a href="#">MEDseq_control</a> .
NEC	A matrix of <i>all</i> NEC values with $\text{length}\{G\}$ rows and $\text{length}(\text{modtype})$ columns, if available. See Note and the argument <code>do.nec</code> to <a href="#">MEDseq_control</a> .
bic	The BIC value corresponding to the optimal model. May not necessarily be the optimal BIC.
icl	The ICL value corresponding to the optimal model. May not necessarily be the optimal ICL.



aic	The AIC value corresponding to the optimal model. May not necessarily be the optimal AIC.
loglik	The vector of increasing log-likelihood values for every EM/CEM iteration under the optimal model. The last element of this vector is the maximum log-likelihood achieved by the parameters returned at convergence.
df	The number of estimated parameters in the optimal model (i.e. the number of 'used' degrees of freedom). Subtract this number from the sample size to get the degrees of freedom.
iters	The total number of EM/CEM iterations for the optimal model.
cv	The cross-validated log-likelihood value corresponding to the optimal model, if available. May not necessarily be the optimal one.
nec	The NEC value corresponding to the optimal model, if available. May not necessarily be the optimal NEC.
ZS	A list of lists giving the z matrices for <i>all</i> fitted models. The first level of the list corresponds to numbers of components, the second to the MEDseq model types.
uncert	The uncertainty associated with the classification.
covars	A data frame gathering the set of covariates used in the gating network, if any. Will contain zero columns in the absence of gating covariates. Supplied gating covariates will be excluded if the optimal model has only one component. May have fewer columns than covariates supplied via the covars argument also, as only the included covariates are gathered here.

Dedicated [plot](#), [print](#), and [summary](#) functions exist for objects of class "MEDseq".

### Note

Where BIC, ICL, AIC, DBS, ASW, LOGLIK, DF, ITTERS, CV, and NEC contain NA entries, this corresponds to a model which was not run; for instance a UU model is never run for single-component models as it is equivalent to CU, while a UCN model is never run for two-component models as it is equivalent to CCN. As such, one can consider the value as not really missing, but equivalent to the corresponding value. On the other hand,  $-\text{Inf}$  represents models which were terminated due to error, for which a log-likelihood could not be estimated. These objects all inherit the class "MEDCriterion" for which dedicated [print](#) and [summary](#) methods exist. For plotting, please see [plot](#).

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

### See Also

[seqdef](#), [MEDseq\\_control](#), [MEDseq\\_compare](#), [plot.MEDseq](#), [predict.MEDgating](#), [MEDseq\\_stderr](#), [I](#), [MEDseq\\_clustnames](#), [seqformat](#)

**Examples**

```

# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad      <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov  <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states    <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels    <- c("Employment", "Further Education", "Higher Education",
  "Joblessness", "School", "Training")
mvad.seq  <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a range of exponential-distance models without clustering
mod0      <- MEDseq_fit(mvad.seq, G=1)

# Fit a range of unweighted mixture models without covariates
# Only consider models with a noise component
# Supply some MEDseq_control() arguments

# mod1     <- MEDseq_fit(mvad.seq, G=9:10, modtype=c("CCN", "CUN", "UCN", "UUN"),
#                       algo="CEM", init.z="kmodes", criterion="icl")

# Fit a model with weights and a gating covariate
# Have the probability of noise-component membership be constant
mod2      <- MEDseq_fit(mvad.seq, G=11, modtype="UUN", weights=mvad$weights,
  gating=~ gcse5eq, covars=mvad.cov, noise.gate=FALSE)

# Examine this model in greater detail
summary(mod2, classification=TRUE, parameters=TRUE)
summary(mod2$gating, SPS=TRUE)
print(mod2$params$theta, SPS=TRUE)
plot(mod2, "clusters")

```

---

MEDseq\_meantime

*Compute the mean time spent in each sequence category*


---

**Description**

Computes the mean time (per cluster) spent in each sequence category (i.e. state value) for a fitted MEDseq model.

**Usage**

```
MEDseq_meantime(x,
```

```

MAP = FALSE,
weighted = TRUE,
norm = TRUE,
prop = FALSE,
map.size = FALSE,
wt.size = FALSE,
SPS = FALSE)

## S3 method for class 'MEDseqMeanTime'
print(x,
      digits = 3L,
      ...)

```

### Arguments

x	An object of class "MEDseq" generated by <a href="#">MEDseq_fit</a> or an object of class "MEDseqCompare" generated by <a href="#">MEDseq_compare</a> .
MAP	A logical indicating whether to use the MAP classification in the computation of the averages, or the 'soft' clustering assignment probabilities given by <code>x\$z</code> . Defaults to FALSE, but is always TRUE for models fitted by the CEM algorithm (see <a href="#">MEDseq_control</a> ). See <code>weighted</code> for incorporating the sampling weights (regardless of the value of MAP). See <code>map.size</code> below.
weighted	A logical indicating whether the sampling weights (if used during model fitting) are used to compute the weighted averages. These can be used alone (when MAP is TRUE) or in conjunction with the 'soft' clustering assignment probabilities (when MAP is FALSE). Defaults to TRUE. Note that, <i>by default</i> , the first column of the output is not affected by the value of <code>weighted</code> (see <code>wt.size</code> ).
norm	A logical indicating whether the mean times (outputted values after the first column) are normalised to sum to the sequence length within each cluster (defaults to TRUE). Otherwise, when FALSE, entries beyond the first column give the total (weighted) number of times a given sequence category was observed in a given cluster.
prop	A logical (defaulting to FALSE and only invoked when <code>norm</code> is also TRUE) which further normalises the output to give the <i>proportions</i> of time spent in each state on average instead of the absolute values.
map.size	A logical (defaulting to FALSE, unless the model was fitted by the CEM algorithm (see <a href="#">MEDseq_control</a> )) which overrides MAP in the <code>Size</code> column (or <code>Weighted.Size</code> column, see <code>wt.size</code> ) of the output, e.g. if <code>MAP=FALSE</code> and <code>map.size=TRUE</code> , the MAP classification is used to determine the cluster sizes but the soft cluster-membership probabilities are used to calculate quantities in remaining columns. Only relevant when <code>MAP=FALSE</code> or <code>wt.size=TRUE</code> .
wt.size	A logical (default to FALSE and only invoked when <code>weighted</code> is also TRUE) which toggles whether the weights are <i>also</i> used in the computation of the cluster sizes in the first column of the output (regardless of the values of MAP or <code>map.size</code> ).
SPS	A logical indicating whether the output should be labelled according to the state-permanence-sequence representation of the central sequences. Defaults

to FALSE. See [MEDseq\\_clustnames](#) and [seqformat](#).  
 digits Minimum number of significant digits to be printed in values.  
 ... Catches unused arguments.

### Details

Models with weights, covariates, &/or a noise component are also accounted for.

### Value

A matrix with sequence category and cluster-specific mean times, giving clusters on the rows, corresponding cluster sizes (or weighted cluster sizes) in the first column, and sequence categories in the remaining columns.

### Note

The function `plot.MEDseq` with the option `type="mt"` can be used to visualise the mean times (by cluster). However, the results displayed therein (at present) always assume `norm=TRUE`, `prop=FALSE`, and `wt.size=TRUE`, while the MAP argument is renamed to `soft`, where `MAP=!soft`.

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

### See Also

[MEDseq\\_fit](#), [MEDseq\\_control](#), [plot.MEDseq](#)

### Examples

```
data(biofam)
seqs <- seqdef(biofam[10:25] + 1L,
              states = c("P", "L", "M", "L+M", "C",
                        "L+C", "L+M+C", "D"))
mod <- MEDseq_fit(seqs, G=10, modtype="UUN")

MEDseq_meantime(mod)
MEDseq_meantime(mod, prop=TRUE)
MEDseq_meantime(mod, map.size=TRUE)
MEDseq_meantime(mod, MAP=TRUE, norm=FALSE, SPS=TRUE)
```

---

MEDseq_news	<i>Show the NEWS file</i>
-------------	---------------------------

---

**Description**

Show the NEWS file of the MEDseq package.

**Usage**

```
MEDseq_news()
```

**Value**

The MEDseq NEWS file, provided the session is interactive.

**Examples**

```
MEDseq_news()
```

---

MEDseq_stderr	<i>MEDseq gating network standard errors</i>
---------------	--

---

**Description**

Computes standard errors of the gating network coefficients in a fitted MEDseq model using either the Weighted Likelihood Bootstrap or Jackknife methods.

**Usage**

```
MEDseq_stderr(mod,
               method = c("WLBS", "Jackknife"),
               N = 1000L,
               symmetric = TRUE,
               SPS = FALSE)
```

**Arguments**

mod	An object of class "MEDseq" generated by <a href="#">MEDseq_fit</a> or an object of class "MEDseqCompare" generated by <a href="#">MEDseq_compare</a> .
method	The method used to compute the standard errors (defaults to "WLBS", the Weighted Likelihood Bootstrap).
N	The (integer) number of samples to use when the "WLBS" method is employed. Defaults to 1000L. Not relevant when method="Jackknife", in which case N is always the number of observations. Must be > 1, though N being greater than or equal to the sample size is recommended under method="WLBS".

symmetric	A logical indicating whether symmetric draws from the uniform Dirichlet distribution are used for the WLBS method in the presence of existing sampling weights. Defaults to TRUE; when FALSE, the concentration parameters of the Dirichlet distribution are given by the sampling weights. Only relevant when method="WLBS" for models with existing sampling weights.
SPS	A logical indicating whether the output should be labelled according to the state-permanence-sequence representation of the central sequences. Defaults to FALSE. See <a href="#">MEDseq_clustnames</a> and <a href="#">seqformat</a> .

### Details

A progress bar is displayed as the function iterates over the N samples. The function may take a long time to run for large N. The function terminates immediately if `mod$G == 1`.

### Value

A list with the following two elements:

**Coefficients** The original matrix of estimated coefficients (`coef(mod$gating)`).

**Std. Errors** The matrix of corresponding standard error estimates.

### Note

The `symmetric` argument is an experimental feature. More generally, caution is advised in interpreting the standard error estimates under *either* the "WLBS" **or** the "Jackknife" method when there are existing sampling weights which arise from complex/stratified sampling designs.

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.

O'Hagan, A., Murphy, T. B., Scrucca, L., and Gormley, I. C. (2019). Investigation of parameter uncertainty in clustering using a Gaussian mixture model via jackknife, bootstrap and weighted likelihood bootstrap. *Computational Statistics*, 34(4): 1779-1813.

### See Also

[MEDseq\\_fit](#), [MEDseq\\_clustnames](#), [seqformat](#)

## Examples

```
# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad          <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov      <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states        <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels        <- c("Employment", "Further Education", "Higher Education",
  "Joblessness", "School", "Training")
mvad.seq      <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a model with weights and a gating covariate
# Have the probability of noise-component membership be constant
# mod          <- MEDseq_fit(mvad.seq, G=11, modtype="UUN", weights=mvad$weights,
#                             gating=~ gcse5eq, covars=mvad.cov, noise.gate=FALSE)

# Estimate standard errors using 100 WLBS samples
# (std         <- MEDseq_stderr(mod, N=100))
```

---

mvad

*MVAD: Transition from school to work*


---

## Description

The data comes from a study by McVicar and Anyadike-Danes on transition from school to work. The data consist of static background characteristics and a time series sequence of 72 monthly labour market activities for each of a cohort of 712 individuals in the Status Zero Survey. The individuals were followed up from July 1993 to June 1999. The monthly states are recorded in columns 15 (Jul.93) to 86 (Jun.99).

## Usage

```
data(mvad)
```

## Format

A data frame containing 712 rows, 72 state variables, 1 id variable and 13 covariates.

## Details

States are:

employment	(EM)
FE	further education (FE)
HE	higher education (HE)
joblessness	(JL)
school	(SC)
training	(TR)

The data set contains also ids (`id`) and sample weights (`weights`) as well as the following binary covariates:

`male`

`catholic`

Belfast, N.Eastern, Southern, S.Eastern, Western (location of school, one of five Education and Library Board areas in Northern Ireland)

Grammar (type of secondary education, 1=grammar school)

funemp (father's employment status at time of survey, 1=father unemployed)

gcse5eq (qualifications gained by the end of compulsory education, 1=5+ GCSEs at grades A-C, or equivalent)

fmpr (SOC code of father's current or most recent job at time of survey, 1=SOC1 (professional, managerial or related))

livboth (living arrangements at time of first sweep of survey (June 1995), 1=living with both parents)

### Note

The first two months of the observation period coincide with summer holidays from school. Hence, documented examples throughout this package extract only the states in columns 17 to 86; i.e. sequences of length 70 from Sep. 93 to Jun. 99.

### Source

McVicar and Anyadike-Danes (2002)

### References

McVicar, D. (2000). Status 0 four years on: young people and social exclusion in Northern Ireland. *Labour Market Bulletin*, 14, 114-119.

McVicar, D. and Anyadike-Danes, M. (2002). Predicting successful and unsuccessful transitions from school to work by using sequence methods. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 165(2): 317-334.

### Examples

```
data(mvad, package="MEDseq")

mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
```



```

mvad          <- list(covariates = mvad[c(3:4,10:14,87)],
                    sequences = mvad[,15:86],
                    weights = mvad[,2])
mvad.cov      <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states       <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels       <- c("Employment", "Further Education", "Higher Education",
                "Joblessness", "School", "Training")
mvad.seq     <- seqdef(mvad$sequences[-c(1,2)], states=states,
                    labels=labels, weights=mvad$weights)

```

---

plot.MEDseq	<i>Plot MEDseq results</i>
-------------	----------------------------

---

## Description

Produces a range of plots of the results of fitted MEDseq models.

## Usage

```

## S3 method for class 'MEDseq'
plot(x,
     type = c("clusters", "central", "precision", "gating",
             "bic", "icl", "aic", "dbs", "asw", "cv",
             "nec", "LOGLIK", "dbsvals", "aswvals",
             "uncert.bar", "uncert.profile", "loglik",
             "d", "f", "Ht", "i", "I", "ms", "mt"),
     seriated = c("observations", "both", "clusters", "none"),
     soft = NULL,
     weighted = TRUE,
     SPS = NULL,
     smeth = "TSP",
     sortv = NULL,
     subset = NULL,
     quant.scale = FALSE,
     ...)

```

## Arguments

- x** An object of class "MEDseq" generated by [MEDseq\\_fit](#) or an object of class "MEDseqCompare" generated by [MEDseq\\_compare](#).
- type** A character string giving the type of plot requested:
- "clusters" Visualise the data set with sequences grouped into their respective clusters. See `seriated`. Similar to the `type="I"` plot (see below). However, `type="clusters"` always plots the hard MAP partition and is unaffected by the `soft` argument below.

- "central" Visualise the central sequences (typically modal sequences, but this depends on the `opti` argument to `MEDseq_control` used during model-fitting). See `seriated`. The central sequence for the noise component, if any, is not shown as it doesn't contribute in any way to the likelihood. See the `type="ms"` option below for an alternative means of displaying the central sequences.
- "precision" Visualise the precision parameters in the form of a heatmap. Values of 0 and Inf are shown in "white" and "black" respectively (see `quant.scale` and `seriated`).
- "gating" Visualise the gating network, i.e. the observation index (by default) against the mixing proportions for that observation, coloured by cluster. See `seriated`. The optional argument `x.axis` can be passed via the `...` construct to change the x-axis against which mixing proportions are plotted (only advisable for models with a single gating network covariate, when `x.axis` is a quantity related to the gating network of the fitted model).
- "bic" Plots all BIC values in a fitted MEDseq object.
- "icl" Plots all ICL values in a fitted MEDseq object.
- "aic" Plots all AIC values in a fitted MEDseq object.
- "dbs" Plots all (weighted) mean/median DBS *criterion* values in a fitted MEDseq object.
- "asw" Plots all (weighted) mean/median ASW *criterion* values in a fitted MEDseq object.
- "cv" Plots all cross-validated log-likelihood values in a fitted MEDseq object.
- "nec" Plots all NEC values in a fitted MEDseq object.
- "LOGLIK" Plots all maximal log-likelihood values in a fitted MEDseq object.
- "dbsvals" Silhouette plot using observations-specific DBS values for the optimal model (coloured by cluster). See `seriated`.
- "aswvals" Silhouette plot using observations-specific ASW values for the optimal model (coloured by cluster). See `seriated`.
- "uncert.bar" Plot the observation-specific clustering uncertainties, if any, in the form of a bar plot.
- "uncert.profile" Plot the observation-specific clustering uncertainties, if any, in the form of a profile plot.
- "loglik" Plot the log-likelihood at every iteration of the EM/CEM algorithm used to fit the model.

Also available are the following options which act as wrappers to types of plots produced by the `seqplot` function in the **TraMineR** package. All are affected by the value of `seriated` and all account for the sampling weights (if any) by default (see the `weighted` argument and the related Note below).

Note also that all of the plot types below can be made to either work with the hard MAP partition, or to use the soft cluster membership probabilities, via the `soft` argument below. The soft information is used by default for all but the "i" and "I" plot types, which (by default) discard this information to instead use the MAP partition: see the `soft` argument below for modifying this default behaviour for all of the following plot types.

- "d" State distribution plots (by cluster).

- "f" Sequence frequency plots (by cluster).
  - "Ht" Transversal entropy plots (by cluster).
  - "i" Selected sequence index plots (by cluster). By default, bar widths for each observation will be proportional to their weight (if any). However, this can be overruled by specifying `weighted=FALSE`.
  - "I" Whole set index plots (by cluster). This plot effectively contains almost exactly the same information as `type="clusters"` plots, and is similarly affected by the `seriated` argument, albeit shown on a by-cluster basis rather than stacked in one plot. However, bar widths for each observation will (by default) be proportional to their weight (if any), which is not the case for `type="clusters"` plots. However, this can be overruled by specifying `weighted=FALSE`.
  - "ms" Modal state sequence plots (by cluster). This is an alternative way of displaying the central sequences beyond the `type="central"` option above. Notably, this option respects arguments passed to `get_MEDseq_results` via the `...` construct (see below), while `type="central"` does not, although still nothing is shown for the noise component. **Note:** unlike `type="central"`, this option always plots *modal* sequences, even if another `opt i` setting was invoked during model-fitting via `MEDseq_control`, in which case there will be a mismatch between the visualisation and `x$params$theta`. Similarly, there may be a mismatch if `soft` and/or `weighted` are modified from their default values of `TRUE`.
  - "mt" Mean times plots (by cluster). This is equivalent to plotting the results of `MEDseq_meantime(x,MAP=!soft,weighted=weighted,norm=TRUE,prop=FALSE,map.size=FALSE)`. Other options for `norm=FALSE`, `prop=TRUE`, `map.size=TRUE`, and `wt.size=FALSE` may be added in future versions of this package.
- `seriated` Switch indicating whether seriation should be used to improve the visualisation by re-ordering the "observations" within clusters (the default), the "clusters", "both", or "none". See `seriate` and the `smeth` and `sortv` arguments below.
- The "clusters" option (and the cluster-related part of "both") is only invoked when `type` is one of "clusters", "central", "precision", "gating", "dbsvals", "aswvals", "d", "f", "Ht", "i", "I", "ms", or "mt".
- Additionally, the "observations" option (and the observation-related part of "both") is only invoked when `type` is one of "clusters", "gating", "i" or "I", which are also the only options for which "both" is relevant.
- `soft` This argument is a single logical indicator which is only relevant for the "d", "f", "Ht", "i", "I", "ms", and "mt" plot types borrowed from **TraMineR**. When `soft=TRUE` (the default for all but the "i" and "I" type plots) the soft cluster membership probabilities are used in a manner akin to `fuzzyseqplot`. Otherwise, when `FALSE` (the default for "i" and "I" type plots), the soft information is discarded and the hard MAP partition is used instead.
- Note that soft cluster membership probabilities will not be available if `x$G=1` or the model was fitted using the `algo="CEM"` option to `MEDseq_control`. Plots may still be weighted when `soft` is `FALSE`, according to the observation-specific sampling weights, when `weighted=TRUE`. Note also that `type="Ht"` can be used in conjunction with `soft=TRUE`, unlike `fuzzyseqplot` for which `type="Ht"` is

	not permissible. Finally, be advised that plotting may be time-consuming when <code>soft=TRUE</code> for "i" and "I" type plots.
weighted	This argument is a single logical indicator which is only relevant for the "d", "f", "Ht", "i", "I", "ms", and "mt" plot types borrowed from <b>TraMineR</b> . When TRUE (the default), the weights (if any) are accounted for in such plots. Note that when <code>soft</code> is TRUE, plots will still be weighted according to the soft cluster membership probabilities; thus <code>weighted=TRUE</code> and <code>soft=TRUE</code> allows both these and the observation-specific weights to be used simultaneously (the default behaviour for both arguments).
SPS	A logical indicating whether clusters should be labelled according to the state-permanence-sequence representation of their central sequence. See <a href="#">MEDseq_clustnames</a> and <a href="#">seqformat</a> . Defaults to TRUE for the plot types adapted from <b>TraMineR</b> , i.e. the "d", "f", "Ht", "i", "I", "ms", and "mt" type plots. The SPS argument is also relevant for the following type plots: "clusters", "central", and "precision", though SPS defaults to FALSE in those instances.
smeth	A character string with the name of the seriation method to be used. Defaults to "TSP". See <a href="#">seriate</a> and <code>seriation::list_seriation_methods("dist")</code> for further details and the available methods. Only relevant when <code>seriated != "none"</code> . When <code>seriated == "obs"</code> or <code>seriated == "both"</code> , the ordering of observations can be governed by <code>smeth</code> or <i>instead</i> governed by the <code>sortv</code> argument below.
sortv	A sorting method governing the ordering of observations for "clusters", "gating", "i", or "I" type plots. Potential options include "dbs" and "asw" for sorting observations by their DBS or ASW values (if available). Only relevant if <code>seriated</code> is one of "observations" or "both". Note that the <code>sortv</code> argument overrides the setting in <code>smeth</code> as it pertains to the ordering of observations if <code>sortv</code> is supplied; otherwise <code>sortv</code> is NULL and the <code>smeth</code> is invoked. Additionally, when (and only when) <code>soft=TRUE</code> and <code>type="I"</code> , the additional option <code>sortv="membership"</code> is provided in accordance with <a href="#">fuzzyseqplot</a> , on which such plots are based.
subset	An optional numeric vector giving the indices of the clusters to be plotted. For models with a noise component, values in $\emptyset : x\$G$ are admissible, where $\emptyset$ denotes the noise component, otherwise only values in $1 : x\$G$ . Only relevant for the <b>TraMineR</b> -type plots, i.e. "d", "f", "Ht", "i", "I", "ms", and "mt" type plots. Note however, that noise components are never plotted for <code>type="ms"</code> plots, so <code>subset</code> values of $\emptyset$ will be ignored in this instance.
quant.scale	Logical indicating whether precision parameter heatmaps should use quantiles to determine non-linear colour break-points when <code>type="precision"</code> . This ensures each colour represents an equal proportion of the data. The behaviour of $\emptyset$ or $\text{Inf}$ values remains unchanged; only strictly-positive finite entries are affected. Heavily imbalanced values are more likely for the "UU" and "UUN" model types, thus <code>quant.scale</code> defaults to TRUE in those instances and FALSE otherwise. Note that <code>quant.scale</code> is <i>always</i> FALSE for the "CC" and "CCN" model types.
...	Catches unused arguments, and allows arguments to <a href="#">get_MEDseq_results</a> to be passed when <code>type</code> is one of "clusters", "dbsvals", "aswvals", "uncert.bar",

"uncert.profile", "d", "f", "Ht", "i", "I", "ms", or "mt", as well as the x.axis argument when type="gating". Also allows select additional arguments to the TraMineR function `seqplot` to be used for the relevant plot types (e.g. border and/or ylab, serr where type="mt", and info where type="ms") and the size argument to `MEDseq_clustnames`, where relevant.

### Details

The type options related to model selection criteria plot values for *all* fitted models in the "MEDseq" object x. The remaining type options plot results for the optimal model, by default. However, arguments to `get_MEDseq_results` can be passed via the `...` construct to plot corresponding results for suboptimal models in x when type is one of "clusters", "d", "f", "Ht", "i", "I", "ms", or "mt". See the examples below.

### Value

The visualisation according to type of the results of a fitted MEDseq model.

### Note

Every type of plot respects the sampling weights, if any. However, those related to `seqplot` plots from **TraMineR** ("d", "f", "Ht", "i", "I", "ms", "mt") do so only when `weighted=TRUE` (the default).

For these plot types borrowed from **TraMineR**, when `weighted=TRUE`, the y-axis labels (which can be suppressed using `ylab=NA`) always display cluster sizes which correspond to the output of `MEDseq_meantime(x, MAP=!soft, weighted=weighted, map.size=FALSE, wt.size=TRUE)`, where `wt.size=TRUE` is **NOT** the default behaviour for `MEDseq_meantime`.

Finally, the plot types borrowed from **TraMineR** may be too wide to display in the preview panel. The same may also be true when type is "dbsvals" or "aswvals".

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

- Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.
- Studer, M. (2018). Divisive property-based and fuzzy clustering for sequence analysis. In G. Ritschard and M. Studer (Eds.), *Sequence Analysis and Related Approaches: Innovative Methods and Applications*, pp. 223-239. Cham: Springer International Publishing.
- Gabardin, A., Ritschard, G., Mueller, N. S., and Studer, M. (2011). Analyzing and visualizing state sequences in R with TraMineR. *Journal of Statistical Software*, 40(4): 1-37.

### See Also

`MEDseq_fit`, `seqplot`, `dbs`, `get_MEDseq_results`, `seriate`, `list_seriation_methods`, `fuzzyseqplot`, `MEDseq_meantime`, `MEDseq_clustnames`, `seqformat`

**Examples**

```

# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels <- c("Employment", "Further Education", "Higher Education",
  "Joblessness", "School", "Training")
mvad.seq <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a range of exponential-distance models without clustering
mod0 <- MEDseq_fit(mvad.seq, G=1)

# Show the central sequence and precision parameters of the optimal model
plot(mod0, type="central")
plot(mod0, type="ms")
plot(mod0, type="precision")

# Fit a range of unweighted mixture models without covariates
# Only consider models with a noise component
# mod1 <- MEDseq_fit(mvad.seq, G=9:11, modtype=c("CCN", "CUN", "UCN", "UUN"))

# Plot the DBS values for all fitted models
# plot(mod1, "dbs")

# Plot the clusters of the optimal model (according to the dbs criterion)
# plot(mod1, "clusters", criterion="dbs")

# Use seriation to order the observations and the clusters
# plot(mod1, "cluster", criterion="dbs", seriated="both")

# Use a different seriation method
# seriation::list_seriation_methods("dist")
# plot(mod1, "cluster", criterion="dbs", seriated="both", smeth="Spectral")

# Use the DBS values instead to sort the observations, and label the clusters
# plot(mod1, "cluster", criterion="dbs", seriated="both", sortv="dbs", SPS=TRUE, size=TRUE)

# Plot the observation-specific ASW values of the best CCN model (according to the asw criterion)
# plot(mod1, "aswvals", modtype="CCN", criterion="asw")

# Fit a model with weights and gating covariates
# mod2 <- MEDseq_fit(mvad.seq, G=10, modtype="UCN", weights=mvad$weights,
#   gating=~ fmpr + gcse5eq + livboth, covars=mvad.cov)

```

```

# Plot the central sequences & precision parameters of this model
# plot(mod2, "central")
# plot(mod2, "precision")

# Plot the clustering uncertainties in the form of a barplot
# plot(mod2, "uncert.bar")

# Plot the observation-specific DBS values
# plot(mod2, "dbsvals")

# Plot the transversal entropies by cluster & then the state-distributions by cluster
# Note that these plots may not display properly in the preview panel
# plot(mod2, "Ht", ylab=NA) # suppress the y-axis labels
# plot(mod2, "d", border=TRUE) # add borders

# The plots above use the soft cluster membership probabilities
# Discard this information and reproduce the per-cluster state-distributions plot
# plot(mod2, "d", soft=FALSE)

# The plots above use the observation-specific sampling weights
# Discard this information and plot the mean times per state per cluster
# plot(mod2, "mt", weighted=FALSE)

# Use type="I" and subset=0 to examine the noise component
# plot(mod2, "I", subset=0, border=TRUE, weighted=FALSE, seriated="none")

```

---

predict.MEDgating      *Predictions from MEDseq gating networks*

---

## Description

Predicts mixing proportions from MEDseq gating networks. Effectively akin to predicting from a multinomial logistic regression via `multinom`, although here the noise component (if any) is properly accounted for. So too are models with no gating covariates at all, or models with the equal mixing proportion constraint. Prior probabilities are returned by default.

## Usage

```

## S3 method for class 'MEDgating'
predict(object,
        newdata = NULL,
        type = c("probs", "class"),
        keep.noise = TRUE,
        droplevels = FALSE,
        ...)

## S3 method for class 'MEDgating'
fitted(object,
        ...)

```

```
## S3 method for class 'MEDgating'
residuals(object,
          ...)
```

### Arguments

object	An object of class "MEDgating" (typically x\$gating, where x is of class "MEDseq").
newdata	A matrix or data frame of test examples. If omitted, the fitted values are used.
type	The type of output desired. The default ("probs") returns prior probabilities, while "class" returns labels indicating the most likely group <i>a priori</i> . Note that observations classified assigned the noise component (if any) are given a label of $\emptyset$ .
keep.noise	A logical indicating whether the output should acknowledge the noise component (if any). Defaults to TRUE; when FALSE, this column is discarded and the matrix of probabilities is renormalised accordingly.
droplevels	A logical indicating whether unseen factor levels in categorical variables within newdata should be dropped (with NA predicted in their place). Defaults to FALSE.
...	Catches unused arguments or allows the type and keep.noise arguments to be passed through fitted and the keep.noise argument to be passed through residuals.

### Details

This function (unlike the predict method for `multinom` on which `predict.MEDgating` is based) accounts for sampling weights and the various ways of treating gating covariates, equal mixing proportion constraints, and noise components, although its type argument defaults to "probs" rather than "class".

### Value

The return value depends on whether newdata is supplied or not and whether the model includes gating covariates to begin with. When newdata is not supplied, the fitted values are returned (as matrix if the model contained gating covariates, otherwise as a vector as per x\$params\$tau). If newdata is supplied, the output is always a matrix with the same number of rows as the newdata.

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Murphy, K., Murphy, T. B., Piccarreta, R., and Gormley, I. C. (2021). Clustering longitudinal life-course sequences using mixtures of exponential-distance models. *Journal of the Royal Statistical Society: Series A (Statistics in Society)*, 184(4): 1414-1451. <doi:10.1111/rssa.12712>.



**See Also**[multinom](#)**Examples**

```

# Load the MVAD data
data(mvad)
mvad$Location <- factor(apply(mvad[,5:9], 1L, function(x)
  which(x == "yes")), labels = colnames(mvad[,5:9]))
mvad <- list(covariates = mvad[c(3:4,10:14,87)],
  sequences = mvad[,15:86],
  weights = mvad[,2])
mvad.cov <- mvad$covariates

# Create a state sequence object with the first two (summer) time points removed
states <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels <- c("Employment", "Further Education", "Higher Education",
  "Joblessness", "School", "Training")
mvad.seq <- seqdef(mvad$sequences[-c(1,2)], states=states, labels=labels)

# Fit a model with weights and a gating covariate
# Have the probability of noise-component membership be constant
mod <- MEDseq_fit(mvad.seq, G=11, modtype="UUN", weights=mvad$weights,
  gating=~ gcse5eq, covars=mvad.cov, noise.gate=FALSE)
(preds <- predict(mod$gating, newdata=mvad.cov[1:5,]))

# Note that the predictions are not the same as the multinom predict method
# in this instance, owing to the invocation of noise.gate=FALSE above
mod2 <- mod
class(mod2$gating) <- c("multinom", "nnet")
predict(mod2$gating, newdata=mvad.cov[1:5,], type="probs")

# We can make this function behave in the same way by invoking keep.noise=FALSE
predict(mod$gating, keep.noise=FALSE, newdata=mvad.cov[1:5,])

```

wKModes

*Weighted K-Modes Clustering with Tie-Breaking***Description**

Perform k-modes clustering on categorical data with observation-specific sampling weights and tie-breaking adjustments.

**Usage**

```

wKModes(data,
  modes,
  weights = NULL,

```

```

iter.max = .Machine$integer.max,
freq.weighted = FALSE,
fast = TRUE,
random = TRUE,
...)
```

### Arguments

<code>data</code>	A matrix or data frame of categorical data. Objects have to be in rows, variables in columns.
<code>modes</code>	Either the number of modes or a set of initial (distinct) cluster modes (where each mode is a row and <code>modes</code> has the same number of columns as <code>data</code> ). If a number, a random set of (distinct) rows in <code>data</code> is chosen as the initial modes.
<code>weights</code>	Optional numeric vector containing non-negative observation-specific case weights.
<code>iter.max</code>	The maximum number of iterations allowed. Defaults to <code>.Machine\$integer.max</code> . The algorithm terminates when <code>iter.max</code> is reached or when the partition ceases to change between iterations.
<code>freq.weighted</code>	A logical indicating whether the usual simple-matching (Hamming) distance between objects is used, or a frequency weighted version of this distance. Default to <code>FALSE</code> ; when <code>TRUE</code> , the frequency weights are computed within the algorithm and are <i>not</i> user-specified. Distinct from the observation-level <code>weights</code> above, the frequency weights are assigned on a per-feature basis and derived from the categories represented in each column of data.
<code>fast</code>	A logical indicating whether a fast version of the algorithm should be applied. Defaults to <code>TRUE</code> .
<code>random</code>	A logical indicating whether ties for the modal values &/or assignments are broken at random. Defaults to <code>TRUE</code> (the implied default had been <code>FALSE</code> prior to version 1.3.2 of this package, as per <code>k1aR::kmodes</code> ). Regarding the modes, ties are broken at random when <code>TRUE</code> and the first candidate state is always chosen for the mode when <code>FALSE</code> . Regarding assignments, tie-breaking is always first biased in favour of the observation's most recent cluster: regarding ties thereafter, these are broken at random when <code>TRUE</code> or the first other candidate cluster is always chosen when <code>FALSE</code> .
<code>...</code>	Catches unused arguments.

### Details

The k-modes algorithm (Huang, 1997) is an extension of the k-means algorithm by MacQueen (1967).

The data given by `data` is clustered by the k-modes method (Huang, 1997) which aims to partition the objects into `k` groups such that the distance from objects to the assigned cluster modes is minimised.

By default, the simple-matching (Hamming) distance is used to determine the dissimilarity of two objects. It is computed by counting the number of mismatches in all variables. Alternatively, this distance can be weighted by the frequencies of the categories in data, using the `freq.weighted` argument (see Huang, 1997, for details).

If an initial matrix of modes is supplied, it is possible that no object will be closest to one or more modes. In this case, fewer clusters than the number of supplied modes will be returned and a warning will be printed.

If called using `fast = TRUE`, the reassignment of the data to clusters is done for the entire data set before recomputation of the modes is done. For computational reasons, this option should be chosen for all but the most moderate of data sizes.

### Value

An object of class "wKModes" which is a list with the following components:

`cluster` A vector of integers indicating the cluster to which each object is allocated.

`size` The number of objects in each cluster.

`modes` A matrix of cluster modes.

`withindiff` The within-cluster (weighted) simple-matching distance for each cluster.

`iterations` The number of iterations the algorithm reached.

`weighted` A logical indicating whether observation-level weights were used or not throughout the algorithm.

`freq.weighted` A logical indicating whether feature-level `freq.weights` were used or not in the computation of the distances.

`random` A logical indicating whether ties were broken at random or not throughout the algorithm.

### Note

This code is adapted from the `kmodes` function in the **klaR** package. Specifically, modifications were made to allow for random tie-breaking for the modes and assignments (see `random` above) and the incorporation of observation-specific sampling weights, with a view to using this function as a means to initialise the allocations for MEDseq models (see the `MEDseq_control` argument `init.z` and the related options "kmodes" and "kmodes2").

Notably, the `wKModes` function, when invoked inside `MEDseq_fit`, is used regardless of whether the weights are true sampling weights, or the weights are merely aggregation weights, or there are no weights at all. Furthermore, the `MEDseq_control` argument `random` is *also* passed to `wKModes` when it is invoked inside `MEDseq_fit`.

### Author(s)

Keefe Murphy - <<keefe.murphy@mu.ie>>

### References

Huang, Z. (1997). A fast clustering algorithm to cluster very large categorical data sets in data mining. In H. Lu, H. Motoda, and H. Luu (Eds.), *KDD: Techniques and Applications*, pp. 21-34. Singapore: World Scientific.

MacQueen, J. (1967). Some methods for classification and analysis of multivariate observations. In L. M. L. Cam and J. Neyman (Eds.), *Proceedings of the Fifth Berkeley Symposium on Mathematical Statistics and Probability*, Volume 1, pp. 281-297. Berkeley, CA, USA: University of California Press.

**See Also**

[MEDseq\\_control](#), [MEDseq\\_fit](#)

**Examples**

```
# Load the MVAD data & aggregate the state sequences
data(mvad)
aggMVAD <- wcAggregateCases(mvad[,17:86], weights=mvad$weight)

# Create a state sequence object without the first two (summer) time points
states <- c("EM", "FE", "HE", "JL", "SC", "TR")
labels <- c("Employment", "Further Education", "Higher Education",
           "Joblessness", "School", "Training")
mvad.seq <- seqdef(mvad[aggMVAD$aggIndex, 17:86],
                 states=states, labels=labels,
                 weights=aggMVAD$aggWeights)

# Run k-modes without the weights
resX <- wKModes(mvad.seq, 2)

# Run k-modes with the weights
resW <- wKModes(mvad.seq, 2, weights=aggMVAD$aggWeights)

# Examine the modal sequences of both solutions
seqformat(seqdef(resX$modes), from="STS", to="SPS", compress=TRUE)
seqformat(seqdef(resW$modes), from="STS", to="SPS", compress=TRUE)
```

# Index

- \* **clustering**
  - MEDseq\_compare, 11
  - MEDseq\_fit, 21
  - MEDseq\_stderr, 29
- \* **control**
  - MEDseq\_control, 15
- \* **datasets**
  - biofam, 4
  - mvad, 31
- \* **main**
  - MEDseq\_compare, 11
  - MEDseq\_fit, 21
  - MEDseq\_stderr, 29
  - plot.MEDseq, 33
- \* **package**
  - MEDseq-package, 2
- \* **plotting**
  - plot.MEDseq, 33
- \* **prediction**
  - predict.MEDgating, 39
- \* **utility**
  - dbs, 6
  - get\_MEDseq\_results, 7
  - MEDseq\_clustnames, 9
  - MEDseq\_entropy, 19
  - MEDseq\_meantime, 26
  - MEDseq\_news, 29
  - predict.MEDgating, 39
  - wKModes, 41
- biofam, 3, 4
- dbs, 6, 18, 19, 24, 37
- fitted.MEDgating (predict.MEDgating), 39
- formula, 22
- fuzzyseqplot, 35–37
- get\_MEDseq\_results, 7, 35–37
- hclust, 15, 19
- I, 22, 25
- list\_seriation\_methods, 37
- MEDseq (MEDseq-package), 2
- MEDseq-package, 2
- MEDseq\_clustnames, 9, 22, 25, 28, 30, 36, 37
- MEDseq\_compare, 2, 8, 9, 11, 12, 14, 18, 19, 23, 25, 27, 29, 33
- MEDseq\_control, 2, 15, 18, 20–25, 27, 28, 34, 35, 43, 44
- MEDseq\_entropy, 19
- MEDseq\_fit, 2, 7–9, 11, 12, 14–20, 21, 22, 23, 27–30, 33, 37, 43, 44
- MEDseq\_meantime, 10, 26, 35, 37
- MEDseq\_nameclusts (MEDseq\_clustnames), 9
- MEDseq\_news, 29
- MEDseq\_stderr, 2, 10, 23, 25, 29
- multinom, 18, 19, 23, 39–41
- mvad, 3, 31
- pam, 19
- plot, 25
- plot.MEDseq, 3, 8, 10, 14, 25, 28, 33
- predict, 23
- predict.MEDgating, 25, 39
- print.MEDseq (MEDseq\_fit), 21
- print.MEDseqCompare (MEDseq\_compare), 11
- print.MEDseqMeanTime (MEDseq\_meantime), 26
- residuals.MEDgating
  - (predict.MEDgating), 39
- seqclustname, 10
- seqdef, 21, 22, 25
- seqdist, 19
- seqformat, 10, 22, 25, 28, 30, 36, 37
- seqplot, 34, 37
- seriate, 35–37
- set.seed, 16

summary.MEDseq (MEDseq\_fit), [21](#)

wcKMedoids, [19](#)

wkModes, [15–19](#), [41](#)