Functional PCA in R A software primer using fdapace

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1 Overview

The basic work-flow behind the PACE approach for sparse ¹ functional data is as follows (see eg. [9, 7] for more information):

- 1. Calculate the smoothed mean $\hat{\mu}$ (using local linear smoothing) aggregating all the available readings together.
- 2. Calculate for each curve seperately its own raw covariance and then aggregate all these raw covariances to generate the sample raw covariance.
- 3. Use the off-diagonal elements of the sample raw covariance to estimate the smooth covariance.
- 4. Perform eigenanalysis on the smoothed covariance to obtain the estimated eigenfunctions $\hat{\phi}$ and eigenvalues $\hat{\lambda}$, then project that smoothed covariance on a positive semi-definite surface [5].
- 5. Use Conditional Expectation (PACE step) to estimate the corresponding scores $\hat{\xi}$.

For densely observed functional data simplified procedures are available to obtain the eigencomponents and associated functional principal components scores (see eg. [3] for more information). In particular in this case we:

- 1. Calculate the cross-sectional mean $\hat{\mu}$.
- 2. Calculate the cross-sectional covariance surface (which is guaranteed to be positive semi-definite).
- 3. Perform eigenanalysis on the covariance to estimate the eigenfunctions $\hat{\phi}$ and eigenvalues $\hat{\lambda}$.
- 4. Use numerical integration to estimate the corresponding scores $\hat{\xi}$.

In the case of sparse FPCA the most computational intensive part is the smoothing of the sample's raw covariance function. For this, we employ a local weighted bilinear smoother.

A sibling MATLAB package for fdapace can be found in http://www.stat.ucdavis.edu/PACE.

¹As a working assumption a dataset is treated as sparse if it has on average less than 20, potentially irregularly sampled, measurements per subject. A user can manually change the automatically determined dataType if that is necessary.

2 FPCA in R using fdapace

The simplest scenario is that one has two lists yList and tList where yList is a list of vectors, each containing the observed values Y_{ij} for the ith subject and tList is a list of vectors containing corresponding time points. In this case one uses:

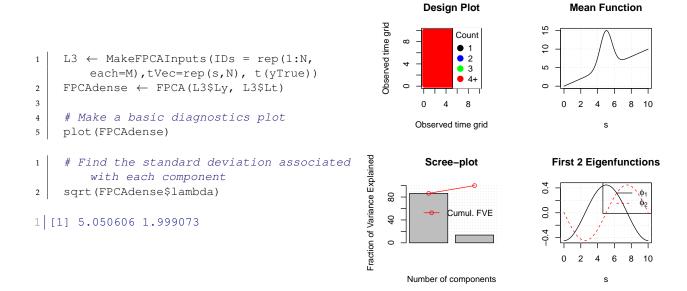
```
1 |FPCAobj ← FPCA(Ly=yList, Lt=tList)
```

The generated FPCAobj will contain all the basic information regarding the desired FPCA.

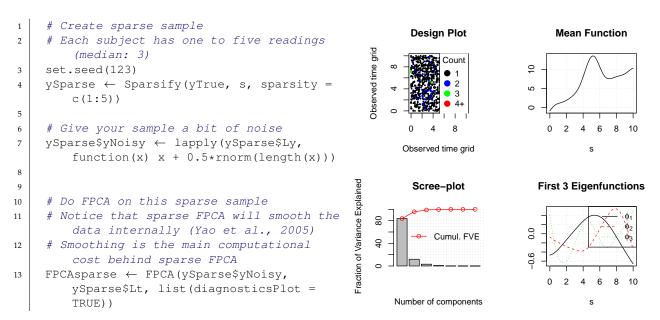
2.1 Generating a toy dense functional dataset from scratch

```
1
     library(fdapace)
2
     #devtools::load_all('.') # So we use the new interfaces
     # Set the number of subjects (N) and the
     # number of measurements per subjects (M)
    N \leftarrow 200;
6
    M \leftarrow 100;
7
8
     set.seed(123)
10
    # Define the continuum
     s \leftarrow seq(0,10,length.out = M)
11
12
     # Define the mean and 2 eigencomponents
     meanFunct \leftarrow function(s) s + 10*exp(-(s-5)^2)
14
     eigFunct1 \leftarrow function(s) +cos(2*s*pi/10) / sqrt(5)
15
     eigFunct2 \leftarrow function(s) -\sin(2*s*pi/10) / sqrt(5)
16
17
     # Create FPC scores
18
     Ksi ← matrix(rnorm(N*2), ncol=2);
19
20
     Ksi \leftarrow apply(Ksi, 2, scale)
     Ksi ← Ksi %*% diag(c(5,2))
21
22
     # Create Y_true
23
    yTrue ← Ksi %*% t(matrix(c(eigFunct1(s),eigFunct2(s)), ncol=2)) +
         t (matrix (rep (meanFunct(s), N), nrow=M))
```

2.2 Running FPCA on a dense dataset



2.3 Running FPCA on a sparse and noisy dataset



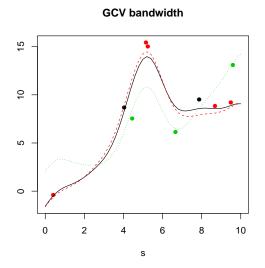
3 Further functionality

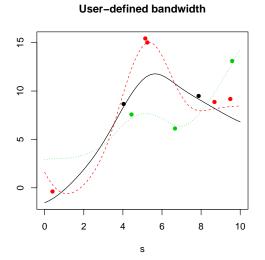
FPCA calculates the bandwidth utilized by each smoother using generalised cross-validation or k-fold cross-validation automatically. Dense data are not smoothed by default. The argument methodMuCovEst can be switched between smooth and cross-sectional if one wants to utilize different estimation techniques when work with dense data. The bandwidth used for estimating the smoothed mean and the smoothed covariance are available under ...\$bwMu and ...\$bwCov respectively. Users can nevertheless provide their own bandwidth estimates:

```
1 | FPCAsparseMuBW5 ← FPCA(ySparse$yNoisy, ySparse$Lt, optns= list(userBwMu = 5))
```

Visualising the fitted trajectories is a good way to see if the new bandwidth made any sense:

```
CreatePathPlot(FPCAsparse, subset = 1:3, main = "GCV bandwidth", pch = 16)
CreatePathPlot(FPCAsparseMuBW5, subset = 1:3, main = "User-defined bandwidth", pch = 16)
```





FPCA uses a Gaussian kernel when smoothing sparse functional data; other kernel types (eg. Epanechnikov/epan) are also available (see ?FPCA). The kernel used for smoothing the mean and covariance surface is the same. It can be found under . . . \$optns\$kernel of the returned object. For instance, one can switch the default Gaussian kernel (gauss) for a rectangular kernel (rect) as follows:

```
FPCAsparseRect ← FPCA(ySparse$yNoisy, ySparse$Lt, optns = list(kernel = 'rect')) #
Use rectangular kernel
```

FPCA returns automatically the smallest number of components required to explain 99.99% of a sample's variance. Using the function <code>selectK</code> one can determine the number of relevant components according to AIC, BIC or a different Fraction-of-Variance-Explained threshold. For example:

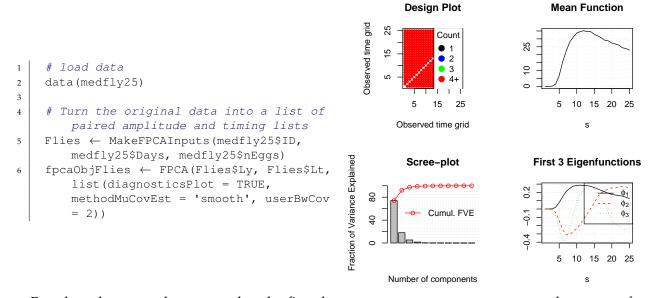
```
SelectK(FPCAsparse, criterion = 'FVE', FVEthreshold = 0.95) \# k = 2 SelectK(FPCAsparse, criterion = 'AIC') \# k = 2
```

When working with functional data (usually not very sparse) the estimation of derivatives is often of interest. Using fitted.FPCA one can directly obtain numerical derivatives by defining the appropriate order p; fdapace provides for the first two derivatives (p = 1 or 2). Because the numerically differentiated data are smoothed the user can define smoothing specific arguments

(see ?fitted.FPCA for more information); the derivation is done by using the derivative of the linear fit. Similarly using the function FPCAder, one can augment an FPCA object with functional derivatives of a sample's mean function and eigenfunctions.

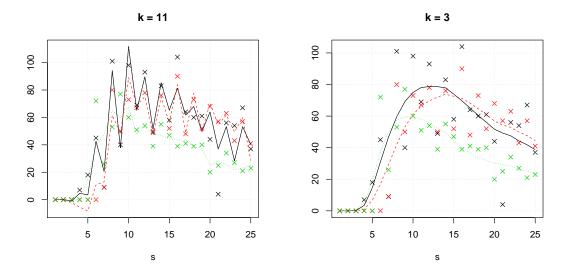
4 A real-world example

We use the medfly25 dataset that this available with fdapace to showcase FPCA and its related functionality. medfly25 is a dataset containing the eggs laid from 789 medflies (Mediterranean fruit flies, Ceratitis capitata) during the first 25 days of their lives. It is a subset of the dataset used by Carey at al. (1998) [2]; only flies having lived at least 25 days are shown. The data are rather noisy, dense and with a characteristic flat start. For that reason in contrast with above we will use a smoothing estimating procedure despite having dense data.



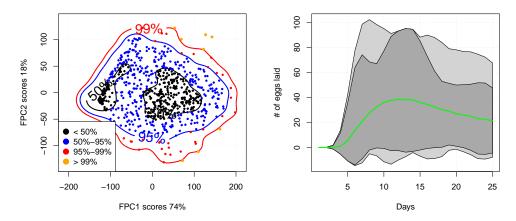
Based on the scree-plot we see that the first three components appear to encapsulate most of the relevant variation. The number of eigencomponents to reach a 99.99% FVE is 11 but just 3 eigencomponents are enough to reach a 95.0%. We can easily inspect the following visually, using the CreatePathPlot command.

```
CreatePathPlot(fpcaObjFlies, subset = c(3,5,135), main = 'k = 11', pch = 4); grid()
CreatePathPlot(fpcaObjFlies, subset = c(3,5,135), k = 3, main = 'k = 3', pch = 4); grid()
```



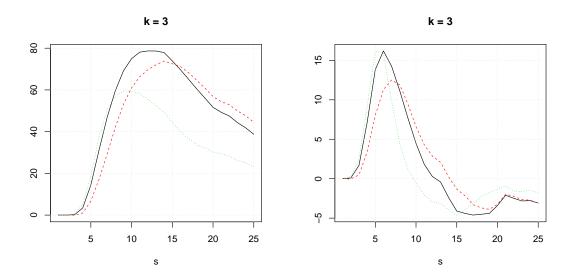
One can perform outlier detection [4] as well as visualize data using a functional box-plot. To achieve these tasks one can use the functions <code>CreateOutliersPlot</code> and <code>CreateFuncBoxPlot</code>. Different ranking methodologies (KDE, bagplot [8, 6] or point-wise) are available and can potentially identify different aspects of a sample. For example here it is notable that the kernel density estimator <code>KDE</code> variant identifies two main clusters within the main body of sample. By construction the <code>bagplot</code> method would use a single <code>bag</code> and this feature would be lost. Both functions return a (temporarily) invisible copy of a list containing the labels associated with each of sample curve. <code>CreateOutliersPlot</code> returns a (temporarily) invisible copy of a list containing the labels associated with each of sample curve.

```
CreateOutliersPlot(fpcaObjFlies, optns = list(k = 3, variant = 'KDE'))
CreateFuncBoxPlot(fpcaObjFlies, xlab = 'Days', ylab = '# of eggs laid', optns = list(k = 3, variant='bagplot'))
```

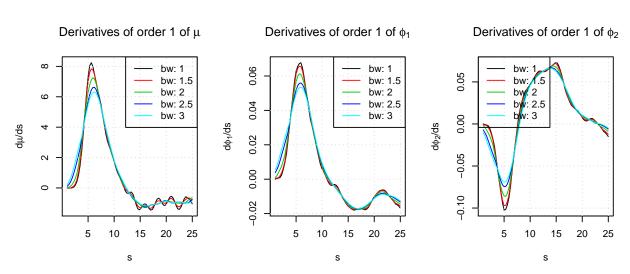


Functional data lend themselves naturally to questions about their rate of change; their derivatives. As mentioned previously using fdapace one can generate estimates of the sample's derivatives (fitted.FPCA) or the derivatives of the principal modes of variation (FPCAder). In all cases, one defines a derOptns list of options to control the derivation parameters. Getting derivatives is obtained by using a local linear smoother as above.

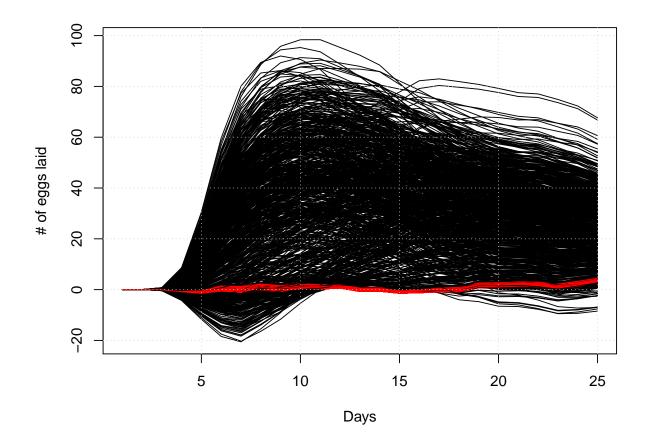
```
1    CreatePathPlot(fpcaObjFlies, subset = c(3,5,135), k = 3, main = 'k = 3', showObs =
    FALSE); grid()
2    CreatePathPlot(fpcaObjFlies, subset = c(3,5,135), k = 3, main = 'k = 3', showObs =
    FALSE, derOptns = list(p = 1, bw = 1.01, kernelType = 'epan')); grid()
```



We note that if finite support kernel types are used (eg. rect or epan), bandwidths smaller than the distance between two adjacent points over which the data are registered onto will lead to (expected) NaN estimates. In case of dense data, the grid used is (by default) equal to the grid the data were originally registered on; in the case of sparse data, the grid used (by default) spans the range of the sample's supports and uses 51 points. A user can change the number of points using the argument nRegGrid. One can investigate the effect a particular kernel type (kernelType) or bandwidth size (bw) has on the generated derivatives by using the function CreateDiagnosticsPlot but this time providing a relevant derOptns list. This will generate estimates about the mean function $\mu(t)$ as well as the first two principal modes of variation $\phi_1(t)$ and $\phi_2(t)$ for different multiples of bw.



As the medfly sample is dense we can immediately use standard multivaritte clustering functionality to identify potential subgroups within it; the function FClust is the wrapper around the clustering functionality provided by fdapace. By default FClust utilises a Gaussian Mixture Model approach based on the package Rmixmod [1], as a general rule clustering optimality is based on negative entropy criterion. In the medfly dataset clustering the data allows to immediately recognise a particular subgroup of flies that lay no or very few eggs during the period examined.



References

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