# **GPTwoSample Documentation**

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gptwosample.py is a tool to run two-sample tests on time series differential gene expression experiments.

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# HANDS ON TUTORIAL FOR GPTWOSAMPLE

In this tutorial we will process a full run example of gptwosample. See the full usage information in *Usage*. See format for input data .csv files in *Data format*. Make sure you cd into the extracted gptwosample folder before running this tutorial. Try printing the full help of the script using:

```
python gptwosample --help
```

If an error occurs, you probably cd one level too deep and you can cd .. up one level.

We will build up the whole running signature step by step in the following. We want to run the script verbosly and with that the script so far looks like:

```
python gptwosample -v
```

To enable plotting we provide the switch -p to the script:

```
python gptwosample -v -p
```

We want to correct for timeshifts (more on *Timeshift detection between replicates*), thus we enable the timeshift switch -t:

```
python gptwosample -v -p -t
```

Next we could additionally learn x confounding factors (see *Accounting for confounding factors* for details on confounding factors) and account for them while two-sampling:

```
python gptwosample -v -p -t -c x
```

but we do not want to account for confounders in this tutorial, leaving the data files to run on.

The output of the script shall be in the subfolder ./tutorial/, so we add the output flag -o ./tutorial/:

```
python gptwosample -v -p -t -o ./tutorial/
```

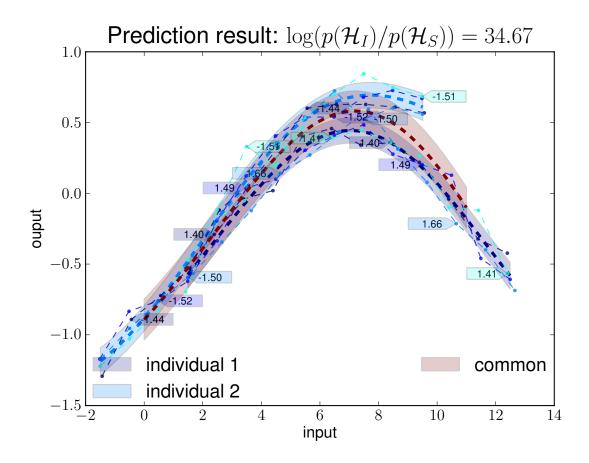
The script shall be run on the two toy condition files  $ToyCondition\{1,2\}.csv$  given in examples/ToyCondition $\{1,2\}.csv$ . These files are non optional as this package is only for comparing two timeseries experiments to each other:

```
python gptwosample -v -p -t -o ./tutorial/ examples/ToyCondition1.csv examples/ToyCondition2.csv
```

Note that the optional parameters could be collected together to give rise to a more compact call signature:

```
python gptwosample -vpto tutorial examples/ToyCondition1.csv
examples/ToyCondition2.csv
```

After hitting return the script runs gptwosample on every gene given in the ToyCondition files and plots each gene into tutorial/plots/. One example plot will look like:



The results are saved in the results.csv, which contains all predicted Bayes Factors and learnt covariance function parameters for all genes ( $Result\ structure$ ).

#### **CHAPTER**

### **TWO**

## **USAGE**

#### Calling signature:

```
gptwosample [-h] [-o DIR] [-t] [-c N] [-p] [-v] [--version] [--backend [PDF,...]] FILE FILE
```

#### where:

```
FILE treatment/control files to compare against each other
-h, --help show this help message and exit
-o DIR, --out DIR set output dir [default: ./twosample_out/]
-t, --timeshift account for timeshifts in data [default: False]
-c N, --confounder N account for N confounders in data [default: 0]
-p, --plot plot data into outdir/plots? [default: False]
-v, --verbose set verbosity level [default: 0]
--version show program's version number and exit
--backend [PDF,...] matplotlib backend - see matplotlib.use(backend)
```

6 Chapter 2. Usage

## **THREE**

## **DATA FORMAT**

The format of the two .csv files (FILE  $\,$  FILE in usage) is as follows:

arbitrary	x1	•••	xl
Gene ID 1	y1 replicate 1		yl replicate 1
Gene ID 1	y1 replicate k1		yl replicate k1
Gene ID n	y1 replicate 1		yl replicate 1
Gene ID n	y1 replicate kn		yl replicate kn

CHAPTER FOUR

# ACCOUNTING FOR CONFOUNDING FACTORS

We detect common confounding factors using probabilistic principal component analysis modeled by gaussian process latent variable models (GPLVM) citep{Lawrence2004Gaussian}. This probabilistic approach to detect low dimensional significant features can be interpreted as detecting common confounding factors in time series experiments by applying GPLVM in advance to two-sample tests of citet{Stegle2010A-robust} on the whole dataset. Two-sample tests on Gaussian Processes decide differential expression based on the bayes factor of marginal probabilities for control and treatment being modeled by one common or two separate underlying function(s). As GPLVM is based on Gaussian Processes it provides a covariance structure of confounders in the dataset. We take this covariance structure between features to build up a two-sample Gaussian Process model taking confounding factors throughout the dataset into account.

**CHAPTER** 

**FIVE** 

# TIMESHIFT DETECTION BETWEEN REPLICATES

A novel covariance function detecting timehifts between time series accounts for temporal mismatches between time series, (of replicates and samples) which share similar patterns, shifted in time. This allows for additional correction of confounding variation in time, as treatment might slow down reaction time of cell-cycle genes, leading to a bunch of falsely positive predicted non differential expressed genes downstream.

**CHAPTER** 

SIX

## **RESULT STRUCTURE**

The results are given in form of a results.csv. Each line corresponds to the results for one gene. The results file is structured as follows:

Gene ID | Bayes Factor | [Learnt covariance function parameters] |

The Gene ID is the ID given in the input files. The Bayes Factor is a log-score for model comparison of the individual model against the common model. The individual model assumes both samples (treatment and control) to be modelled individually by one Gaussian process each. In contrast the common model assumes both samples to be modelled by one Gaussian Process. Both likelihoods are computed and the score is created by contrasting the both likelihoods:

$$\mathcal{BF} = \ln \frac{p(\text{Individual model})}{p(\text{Common model})}$$

All plots are saved in a subfolder <outdir>/plots/

## **DEVELOPERS**

### 7.1 Package for using GPTwoSample

This module allows the user to compare two timelines with respect to diffferential expression.

It compares two timeseries against each other, depicting whether these two timeseries were more likely drawn from the same function, or from different ones. This prediction is defined by which covariance function pygp.covar you use.

Created on Jun 15, 2011

@author: Max Zwiessele, Oliver Stegle

```
 \begin{array}{c} \textbf{class} \ \texttt{gptwosample.twosample.twosample.TwoSample} (\textit{T}, \textit{Y}, \textit{covar\_common=None}, & \textit{covar\_individual\_1=None}, & \textit{covar\_individual\_2=None}) \end{array}
```

Bases: object

Run GPTwoSample on given data.

#### **Parameters:**

- T : TimePoints [n x r x t] [Samples x Replicates x Timepoints]
- Y : ExpressionMatrix [n x r x t x d] [Samples x Replicates x Timepoints x Genes]

#### Fields:

- T: Time Points [n x r x t] [Samples x Replicates x Timepoints]
- Y: Expression [n x r x t x d] [Samples x Replicates x Timepoints x Genes]
- X: Confounders [nrt x 1+q] [SamplesReplicatesTimepoints x T+q]
- lvm\_covariance: GPLVM covaraince function used for confounder learning
- n: Samples
- · r: Replicates
- t: Timepoints
- · d: Genes
- q: Confounder Components

bayes\_factors (likelihoods=None)

get list of bayes\_factors for all genes.

returns: bayes\_factor for each gene in Y

Predict all likelihoods for all genes, given in Y

#### parameters:

indices [[int]] list (or array-like) for gene indices to predict, if None all genes will be predicted

message: str printing message

**kwargs:** {...} kwargs for gptwosample.twosample.GPTwoSampleBase.predict\_model\_likelihoods

Predicts means and variances for all genes given in Y for given interpolation\_interval

#### $set_data(T, Y)$

Set data by time T and expression matrix Y:

#### Parameters:

T [real [n x r x t]] All Timepoints with shape [Samples x Replicates x Timepoints]

Y [real [n x r x t x d]] All expression values given in the form: [Samples x Replicates x Time-points x Genes]

```
 \begin{array}{c} \textbf{class} \ \texttt{gptwosample.confounder.confounder.TwoSampleConfounder} (T, \quad Y, \quad q = 4, \\ \quad lvm\_covariance = None, \\ \quad init= `random', \quad covar\_common = None, \\ \quad covar\_individual\_1 = None, \\ \quad covar\_individual\_2 = None) \\ \end{array}
```

Bases: gptwosample.twosample.TwoSample

Run GPTwoSample on given Data

#### **Parameters:**

- T : TimePoints [n x r x t] [Samples x Replicates x Timepoints]
- Y : ExpressionMatrix [n x r x t x d] [Samples x Replicates x Timepoints x Genes]
- q : Number of Confounders to use
- lvm\_covariance : optional set covariance to use in confounder learning
- init : [random, pca]

#### Fields:

- T: Time Points [n x r x t] [Samples x Replicates x Timepoints]
- Y: Expression [n x r x t x d] [Samples x Replicates x Timepoints x Genes]
- X: Confounders [nrt x 1+q] [SamplesReplicatesTimepoints x T+q]
- lvm\_covariance: GPLVM covaraince function used for confounder learning
- n: Samples
- r: Replicates
- t: Timepoints

- d: Genes
- q: Confounder Components

```
initialize_twosample_covariance (covar\_common = < function < lambda > at 0x11023d0c8 > , \\ covar\_individual\_1 = < function < lambda > at \\ 0x11023d050 > , covar\_individual\_2 = < function \\ < lambda > at 0x11023d2a8 > )
```

initialize twosample covariance with function covariance(XX), where XX is a FixedCF with the learned confounder matrix.

default is SumCF([SqexpCFARD(1), FixedCF(self.K\_conf.copy()), BiasCF()])

Learn confounder matrix with this model.

#### **Parameters:**

x [array-like] If you provided an own lvm\_covariance you have to specify the X to use within GPLVM

lvm\_dimension\_indices [[int]] If you specified an own lvm\_covariance you have to specify the dimension indices for GPLVM

**ard\_indices** [[indices]] If you provided an own lvm\_covariance, give the ard indices of the covariance here, to be able to use the correct hyperparameters for calculating the confounder covariance matrix.

```
class gptwosample.twosample_base.TwoSampleShare(covar, *args, **kwargs)
    Bases: gptwosample.twosample.twosample_base.TwoSampleBase
```

This class provides comparison of two Timeline Groups to each other.

see gptwosample.twosample\_base.TwoSampleBase for detailed description of provided methods.

Bases: gptwosample.twosample\_base.TwoSampleBase

This class provides comparison of two Timeline Groups to one another, inlcuding timeshifts in replicates, respectively.

see gptwosample.twosample\_base.TwoSampleBase for detailed description of provided methods.

Note that this model will need one covariance function for each model, respectively!

```
\begin{tabular}{ll} {\bf class} \ {\tt gptwosample.twosample\_base.TwoSampleBase} \ ({\it learn\_hyperparameters=True}, \\ priors=None, & initial\_hyperparameters=None, \\ **kwargs) \end{tabular}
```

Bases: object

TwoSampleBase object with the given covariance function covar.

```
bayes factor(model likelihoods=None)
```

Return the Bayes Factor for the given log marginal likelihoods model\_likelihoods

#### **Parameters:**

**model\_likelihoods** [{'individual': the individual likelihoods, 'common': the common likelihoods}] The likelihoods calculated by predict\_model\_likelihoods(training\_data) for given training data training\_data.

#### get\_data (model='common', index=None)

get inputs of model model with group index index. If index is None, the whole model group will be returned.

#### get\_learned\_hyperparameters()

Returns learned hyperparameters in model structure, if already learned.

#### get\_model\_likelihoods()

Returns all calculated likelihoods in model structure. If not calculated returns None in model structure.

#### get\_predicted\_mean\_variance()

Get the predicted mean and variance as:

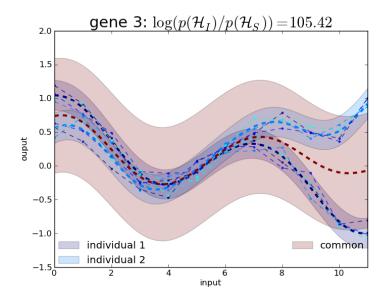
If not yet predicted it will return 'individual' and 'common' empty.

Two Instance Plots of comparing two groups to each other:

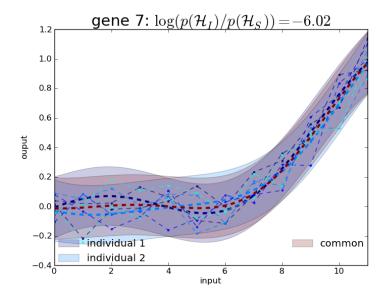
#### **Parameters:**

twosample\_object [gptwosample.twosample] GPTwoSample object, on which already 'predict'
was called.

#### **Differential Groups:**



#### **Non-Differential Groups:**



**Returns:** Proper rectangles for use in pylab.legend().

#### Parameters:

interpolation\_interval [[double]] The interval of inputs, which shall be predicted

**hyperparams** [{'covar':logtheta, ...}] Default: learned hyperparameters. Hyperparams for the covariance function's prediction.

**interval\_indices** [{'common':[boolean],'individual':[boolean]}] Indices in which to predict, for each group, respectively.

```
predict_model_likelihoods (training_data=None, interval_indices={'individual': None, 'com-
mon': None}, *args, **kwargs)
```

Predict the probabilities of the models (individual and common) to describe the data. It will optimize hyperparameters respectively.

#### **Parameters:**

**training\_data** [dict training\_data] The training data to learn from. Input are time-values and output are expression-values of e.g. a timeseries. If not given, training data must be given previously by gptwosample.twosample.basic.set\_data.

**interval\_indices:** gptwosample.data.data\_base.get\_model\_structure() interval indices, which assign data to individual or common model, respectively.

```
args [[..]] see pygp.gpr.gp_base.GP
kwargs [{..}] see pygp.gpr.gp_base.GP
set data(training data)
```

#### Parameters:

Set the data of prediction.

**training\_data** [dict training\_data] The training data to learn from. Input are time-values and output are expression-values of e.g. a timeseries.

Training data training\_data has following structure:

```
{'input' : {'group 1':[double] ... 'group n':[double]},
'output' : {'group 1':[double] ... 'group n':[double]}}
```

## 7.2 GPTwoSample plot

The easiest way to plot your results in an easy and convenient way.

#### 7.2.1 Plot GPTwoSample predictions

Module for easy plotting of GPTwoSample results.

gptwosample.plot.plot\_basic.plot\_results plots training data, as well as sausage\_plots for a GPTwoSample experiment. You can give interval indices for plotting, if u chose

Created on Feb 10, 2011

@author: Max Zwiessele, Oliver Stegle

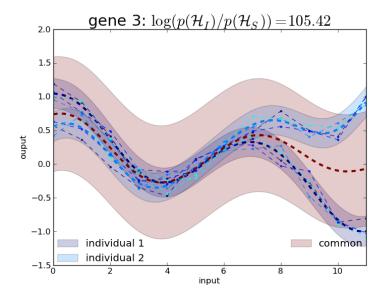
Plot the results given by last prediction.

Two Instance Plots of comparing two groups to each other:

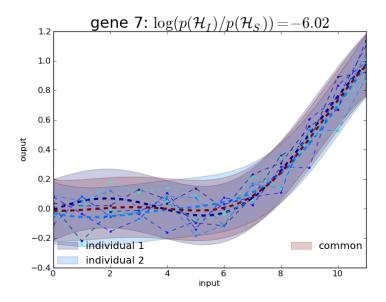
#### **Parameters:**

twosample\_object [gptwosample.twosample] GPTwoSample object, on which already 'predict' was called.

#### **Differential Groups:**



#### **Non-Differential Groups:**



**Returns:** Proper rectangles for use in pylab.legend().

# .. automodule:: gptwosample.plot.interval # :members:

### 7.3 Package for data handling

Use this Package for easiest way to handle the data for GPTwoSample.

#### 7.3.1 Data Structure Module

This Module is for easy access to data structures gptwosample works with.

Created on Mar 18, 2011

@author: Max Zwiessele

exception gptwosample.data.data\_base.DataStructureError(\*args, \*\*kwargs)

Bases: exceptions. TypeError

Thrown, if DataStructure given does not fit. Training data training data has following structure:

```
{input_id : {'group 1':[double] ... 'group n':[double]},
output_id : {'group 1':[double] ... 'group n':[double]}}
```

gptwosample.data.data\_base.get\_model\_structure(individual=None, common=None)

Returns the valid structure for model dictionaries, used in gptwosample. Make sure to use this method if you want to use the model structure in this package!

```
qptwosample.data.data_base.get_training_data_structure(x1, x2, y1, y2)
```

Get the structure for training data, given two inputs x1 and x2 with corresponding outputs y1 and y2. Make sure, that replicates have to be tiled one after the other for proper resampling of data!

```
gptwosample.data.data_base.has_model_structure(structure)
```

Returns the valid structure for model dictionaries, used in gptwosample. Make sure to use this method if you want to use the model structure in this package!

#### 7.3.2 Data IO tool

For convienent usage this module provides IO operations for data

Created on Jun 9, 2011

@author: Max Zwiessele, Oliver Stegle

```
gptwosample.data.dataIO.get_data_from_csv(path_to_file, delimiter=', ', count=-1, ver-
bose=True, message='Reading File')
```

Return data from csv file with delimiter delimiter in form of a dictionary. Missing Values are all values x which cannot be converted float(x)

The file format has to fullfill following formation:

arbitrary	x1	•••	xl
Gene Name 1	y1 replicate 1		yl replicate 1
Gene Name 1	y1 replicate k1		yl replicate k1
Gene Name n	y1 replicate 1		yl replicate 1
Gene Name n	y1 replicate kn		yl replicate kn

Returns: {"input":[x1,...,x1], "Gene Name 1":[[y1 replicate 1, ... y1 replicate 1], ... ,[y1 replicate k, ..., y1 replikate k]]}

```
gptwosample.data.dataIO.write_data_to_csv (data, path_to_file, header='GPTwoSample', de-
limiter='.')
```

Write given data in training\_data\_structure (see <code>gptwosample.data.data\_base</code> for details) into file for path\_to\_file.

#### **Parameters:**

data [dict] data to write in training data structure

path to file [String] The path to the file to write to

header [String] Name of the tabledelimiter [character] delimiter for the csv file

## **PYTHON MODULE INDEX**

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