

# Calibration of Quinine Fluorescence Emission Vignette for the Data Set `flu` of the R package `hyperSpec`

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## Reproducing the Examples in this Vignette

The spectra files are shipped with *hyperSpec*. This allows reproduction of the whole vignette (source code and spectra files are in the package's documentation directory and its `rawdata` subdirectory). For reproducing the examples in a live session, the full file names of the spectra can be found with the command:

```
> Sys.glob (paste (.libPaths (), "hyperSpec/doc/rawdata/flu*.txt", sep = "/")).
```

 Note that loading the package, and some definitions e.g. of the color palettes are executed in `vignettes.def`.

This vignette gives an example how to

- write an import function for a spectrometer manufacturer's proprietary ASCII files,
- add further data columns to the spectra, and
- set up a linear calibration (inverse least squares).

The data set `flu` in *hyperSpec* consists of 6 fluorescence emission spectra of quinine solutions. They were acquired during an student practicum and were kindly provided by M. Kammer.

The concentrations of the solutions range from 0.05 mg/l to 0.30 mg/l. Spectra were acquired with a Perkin Elmer LS50-B fluorescence spectrometer at 350 nm excitation.

## 1 Writing an Import Function

The raw spectra are in Perkin Elmer's ASCII file format, one spectrum per file. The files are completely ASCII text, with the actual spectra starting at line 55.

The function should automatically read in all files specified by a pattern, such as `*.txt`. In order to gain some speed, the spectra matrix is preallocated after the first file is read. Also, the number of header lines are predefined instead of searching for the line after `#DATA`.

Note, that labels giving the correct units (e.g. for axis labels) are set. The label with the special name `.wavelength` corresponds to the wavelength axis, all data columns should have a label with the same name. The spectra are always in a data column called `spc`.

The function is saved in a `.R` file:

```

                                read.txt.PerkinElmer.R
read.txt.PerkinElmer <- function (files = "*.txt", skip = 54, ...) {
  ## find all files
  files <- Sys.glob (files)

  ## read the first file
  buffer <- matrix (scan (files [1], skip = skip, ...), ncol = 2, byrow = TRUE)

  ## first file gives the wavelength vector
  wavelength <- buffer [, 1]

  ## preallocate the spectra matrix: one row per file x as many columns as the
  ## first file has
  spc <- matrix (ncol = nrow (buffer), nrow = length (files))

  ## the first file's data goes into the first row
  spc [1, ] <- buffer [, 2]

  ## now read the remaining files
  for (f in seq (along = files)[-1]) {
    buffer <- matrix (scan (files [f], skip = skip, ...),
                      ncol = 2, byrow = TRUE)

    ## check whether they have the same wavelength axis
    if (! all.equal (buffer [, 1], wavelength))
      stop (paste(files [f], "has different wavelength axis."))

    spc [f, ] <- buffer[, 2]
  }

  ## finally: make the hyperSpec object
  new ("hyperSpec", wavelength = wavelength, spc = spc,
      label = list (.wavelength = expression (lambda[f1] / nm),
                    spc = "I / a.u."))
}

```

That needs to be sourced before use:

```

> source ("read.txt.PerkinElmer.R")
> flu <- read.txt.PerkinElmer ("rawdata/flu?.txt")

```

Now the spectra are in a *hyperSpec* object and can be examined e.g. by

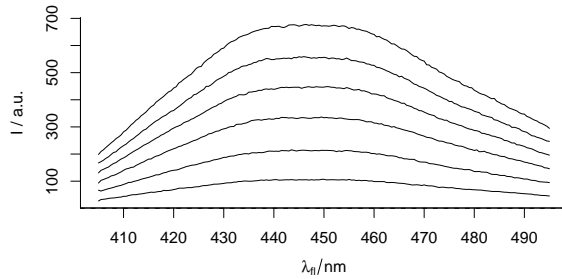
```

> flu

hyperSpec object
  6 spectra
  1 data columns
  181 data points / spectrum
wavelength: lambda[f1]/nm [numeric] 405.0 405.5 ... 495
data: (6 rows x 1 columns)
  1. spc: I / a.u. [AsIs matrix x 181] rng  27.15000 32.34467 ... 677.4947

> plot (flu)

```



## 2 Adding further Data Columns

The calibration model needs the quinine concentrations for the spectra. This information can be stored together with the spectra, and also gets an appropriate label:

```
> flu$c <- seq (from = 0.05, to = 0.30, by = 0.05)
> labels (flu, "c") <- "c / (mg / l)"
> flu

hyperSpec object
  6 spectra
  2 data columns
  181 data points / spectrum
wavelength: lambda[f1]/nm [numeric] 405.0 405.5 ... 495
data: (6 rows x 2 columns)
  1. spc: I / a.u. [AsIs matrix x 181] rng 27.15000 32.34467 ... 677.4947
  2. c: c / (mg / l) [numeric] rng 0.05 0.10 ... 0.3
```

Now the *hyperSpec* object `flu` contains two data columns, holding the actual spectra and the respective concentrations. The dollar operator returns such a data column:

```
> flu$c

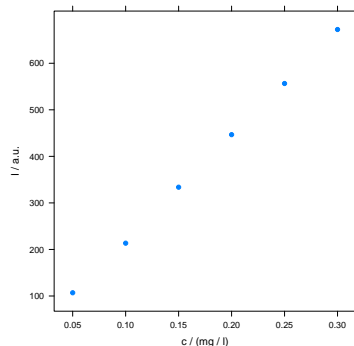
[1] 0.05 0.10 0.15 0.20 0.25 0.30
```

## 3 Linear Calibration

As R is developed for the purpose of statistical analysis, tools for a least squares calibration model are readily available.

The original spectra range from 405 to 495 nm. However, the intensities at 450 nm are perfect for a univariate calibration. Plotting them over the concentration is done by:

```
> plotc (flu[, ,450])
```



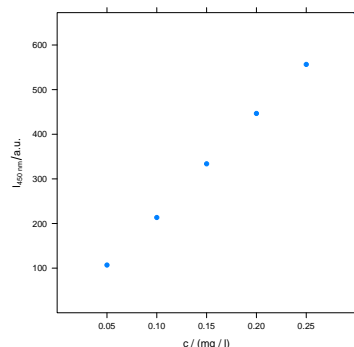
The square bracket operator extracts parts of a *hyperSpec* object. The first coordinate defines which spectra are to be used, the second which data columns, and the third gives the spectral range.

We discard all the wavelengths but 445 nm:

```
> flu <- flu [, , 450]
> labels (flu, "spc") <- expression (I ["445 nm"] / a.u.)
```

The plot could be enhanced by annotating the ordinate with the emission wavelength. Also the axes should start at the origin, so that it is easier to see whether the calibration function will go through the origin:

```
> plotc (flu, xlim = range (0, flu$c), ylim = range (0, flu$spc))
```



The actual calibration is a linear model, which can be fitted by the R function `lm`. `lm` needs a *formula* that specifies which data columns are dependent and independent variables.

The normal calibration plot gives the emission intensity as a function of the concentration, and the calibration function thus models  $I = f(c)$ , i. e.  $I = mc + b$  for a linear calibration. This is then solved for  $c$  when the calibration is used.

However, R's linear model is a quite strict in predicting: a model set up as  $I = f(c)$  will predict the intensity as a function of the concentration but not the other way round. Thus we set up an inverse calibration model<sup>1</sup>:  $c = f(I)$ . The corresponding R formula is `c ~ I`, or in our case `c ~ spc`, as the intensities are stored in the data column `$spc`:

<sup>1</sup>As we can safely assume that the error on the concentrations is far larger than the error on the instrument signal, it is actually the correct type of model from the least squares fit point of view.

In addition, `lm` (like most R model building functions) expects the data to be a *data.frame*.

There are three abbreviations that help to get the parts of the *hyperSpec* object that are frequently needed:

`flu[[]]` returns the spectra matrix. It takes the same indices as `[]`.  
`flu$.` returns the data as a *data.frame*  
`flu$..` returns a *data.frame* that has all data columns but the spectra

```
> flu[[]]

      450
[1,] 106.9497
[2,] 213.4973
[3,] 333.7757
[4,] 446.6307
[5,] 556.5237
[6,] 672.5317
```

```
> flu$.

      450      c
1 106.949666 0.05
2 213.497333 0.10
3 333.775666 0.15
4 446.630666 0.20
5 556.523666 0.25
6 672.531666 0.30
```

```
> flu$..

      c
1 0.05
2 0.10
3 0.15
4 0.20
5 0.25
6 0.30
```

Putting this together, the calibration model is calculated:

```
> calibration <- lm (c ~ spc, data = flu$.)
```

The `summary` gives a good overview of our model:

```
> summary (calibration)
```

Call:

```
lm(formula = c ~ spc, data = flu$.)
```

Residuals:

```
      1      2      3      4      5      6
-0.0009872  0.0020520 -0.0009605 -0.0007012  0.0008636 -0.0002667
```

Coefficients:

```
      Estimate Std. Error t value Pr(>|t|)
```

```

(Intercept) 3.849e-03 1.245e-03 3.091 0.0365 *
spc          4.407e-04 2.870e-06 153.596 1.08e-08 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.001362 on 4 degrees of freedom
Multiple R-squared: 0.9998, Adjusted R-squared: 0.9998
F-statistic: 2.359e+04 on 1 and 4 DF, p-value: 1.078e-08

```

In order to get predictions for new measurements, a new *data.frame* with the same independent variables (in columns with the same names) as in the calibration data are needed. Then the function `predict` can be used. It can also calculate the prediction interval. If we observe e.g. an intensity of 125 or 400 units, the corresponding concentrations and their 99 % prediction intervals are:

```

> I <- c (125, 400)
> conc <- predict (calibration, newdata = list (spc = as.matrix(I)), interval = "prediction",
+               level = .99)
> conc

      fit      lwr      upr
1 0.0589429 0.05132995 0.06655585
2 0.1801488 0.17337542 0.18692213

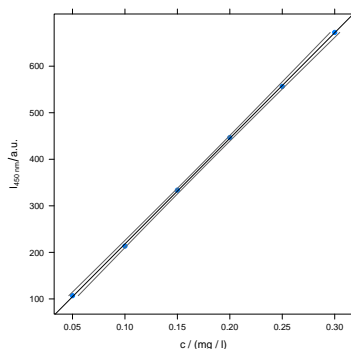
```

Finally, we can draw the calibration function and its 99 % confidence interval (also via `predict`) together with the prediction example. In order to draw the confidence interval into the calibration graph, we can either use a customized panel function:

```

> int <- list (spc = as.matrix(seq (min (flu), max(flu), length.out = 25)))
> ci <- predict (calibration, newdata = int, interval = "confidence", level = 0.99)
> panel.ci <- function (x, y, ...,
+               intensity, ci.lwr, ci.upr, ci.col = "#606060") {
+   panel.xyplot (x, y, ...)
+   panel.lmline (x, y,...)
+   panel.lines (ci.lwr, intensity, col = ci.col)
+   panel.lines (ci.upr, intensity, col = ci.col)
+ }
> plotc (flu, panel = panel.ci,
+       intensity = int$spc, ci.lwr = ci [, 2], ci.upr = ci [, 3])

```



Or, we can add the respective data to the *hyperSpec* object. The meaning of the data can be saved in a new extra data column that acts as grouping variable for the plot.

First, the spectral range of `flu` is cut to contain the fluorescence emission at 450nm only, and the new column is introduced for the original data:

```
> flu$type <- "data points"
```

Next, the calculated confidence intervals are appended:

```
> tmp <- new ("hyperSpec", spc = as.matrix(seq (min (flu), max(flu), length.out = 25)),
+           wavelength = 450)
> ci <- predict (calibration, newdata = tmp$. , interval = "confidence", level = 0.99)
> tmp <- tmp [rep (seq (tmp, index = TRUE), 3)]
> tmp$c <- as.numeric (ci)
> tmp$type <- rep (colnames (ci), each = 25)
> flu <- rbind (flu, tmp)
```

Finally, the resulting object is plotted. Our prediction example is handled by another customized panel function:

```
> panel.predict <- function (x, y, ...,
+           intensity, ci, pred.col = "red", pred.pch = 19, pred.cex = 1) {
+   panel.xyplot (x, y, ...)
+   mapply (function (i, lwr, upr, ...) {
+     panel.lines (c (lwr, upr), rep (i, 2), ...)
+   },
+           intensity, ci [, 2], ci [, 3], MoreArgs = list (col = pred.col))
+   panel.xyplot (ci [, 1], intensity, col = pred.col, pch = pred.pch, cex = pred.cex, type = "p")
+ }
> plotc (flu, groups = type, type = c("l", "p"),
+       col = c ("black", "black", "#606060", "#606060"),
+       pch = c (19, NA, NA, NA), cex = 0.5,
+       lty = c (0, 1, 1, 1),
+       panel = panel.predict,
+       intensity = I,
+       ci = conc,
+       pred.cex = 0.5)
```

