

photobiologyLamps Version 0.1.10

Catalogue of Lamps

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1 Introduction

We will plot the emission spectra of the different lamps for which data is provided in the package. We plot side-by-side the lamp output as spectral energy irradiance and as spectral photon irradiance. All spectra are normalized to an area of one under the whole curve.

```
library(ggplot2)
library(photobiology)
library(photobiologyLamps)
library(photobiologygg)

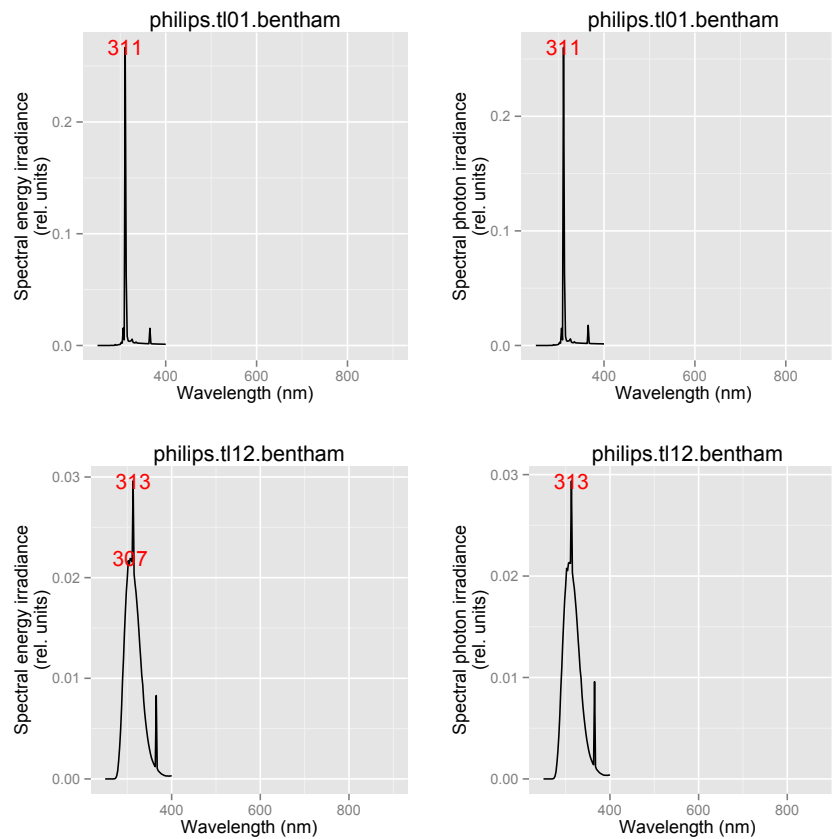
## Loading required package: proto
## Loading required package: spls2R
## Loading required package: plyr
```

We define a function to do the actual plotting so as to not repeat code, and to make changes easier in the future.

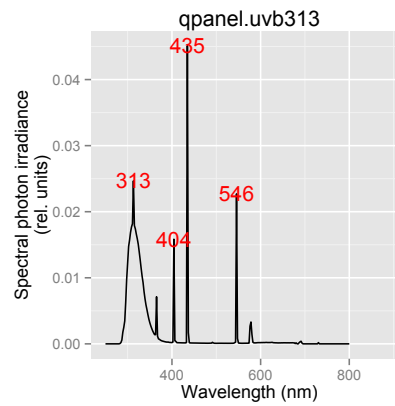
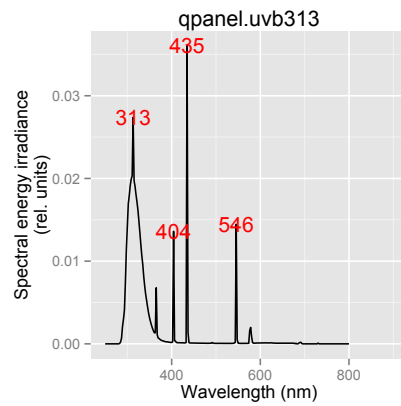
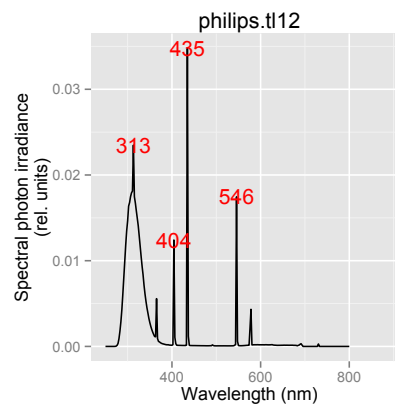
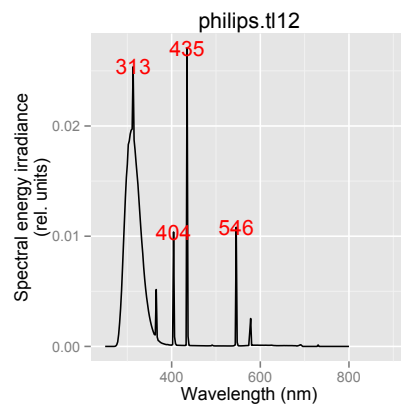
```
lamp.plotter <- function(lamp.name, w.low = 250, w.high = 900, scaled = "area") {
  w.length.out <- seq(from = w.low, to = w.high, length.out = 300)
  spectrum.data <- calc_source_output(w.length.out = w.length.out, source.name = lamp.name,
    scaled = scaled, fill = NULL)
  spectrum.data <- na.omit(spectrum.data)
  fig_energy <- ggplot(aes(x = w.length, y = s.e.irrad), data = spectrum.data) +
    xlim(w.low, w.high) + labs(x = "Wavelength (nm)", y = "Spectral energy irradiance\n(rel. units)",
    title = lamp.name) + geom_line() + stat_peaks(ignore_threshold = 0.33,
    colour = "red")
  fig_photon <- ggplot(aes(x = w.length, y = s.q.irrad), data = spectrum.data) +
    xlim(w.low, w.high) + labs(x = "Wavelength (nm)", y = "Spectral photon irradiance\n(rel. units)",
    title = lamp.name) + geom_line() + stat_peaks(ignore_threshold = 0.33,
    colour = "red")
  print(fig_energy)
  print(fig_photon)
}
```

2 UV-B lamp spectra

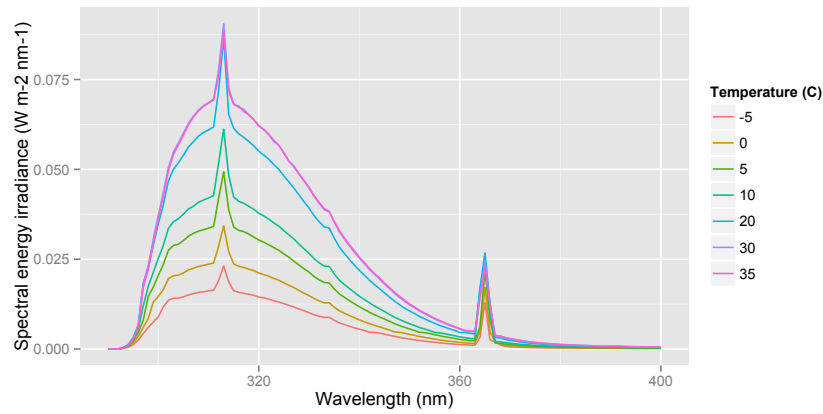
```
UVB.lamps <- c("philips.tl01.bentham", "philips.tl12.bentham")
for (lamp in UVB.lamps) {
  lamp.plotter(lamp.name = lamp)
}
```



```
UVB.M.lamps <- c("philips.tl12", "qpanel.uvb313")
for (lamp in UVB.M.lamps) {
  lamp.plotter(lamp.name = lamp)
}
```

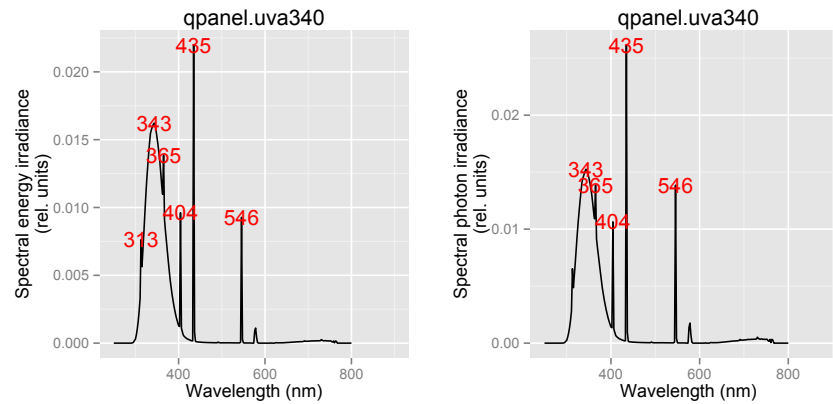


```
fig_temp <- ggplot(data = qpanel.uvb313.temperature.data, aes(x = w.length,
  y = s.e.irrad, colour = factor(temperature)))
fig_temp <- fig_temp + geom_line() + labs(x = "Wavelength (nm)", y = "Spectral energy irradiance (W m-2 nm-1)",
  colour = "Temperature (C)")
print(fig_temp)
```



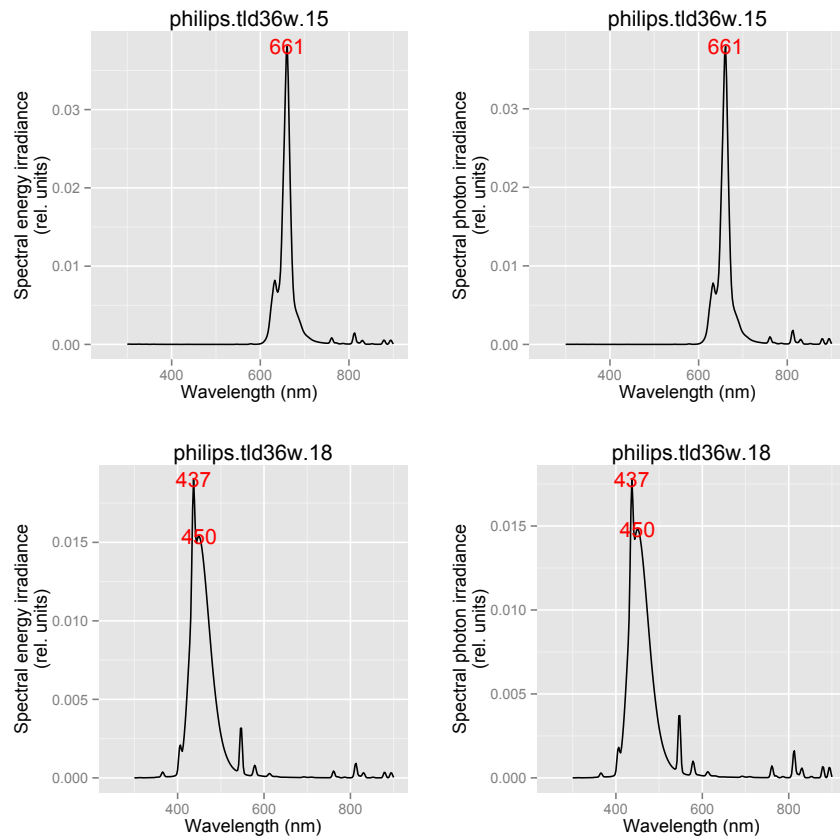
3 UV-A lamp spectra

```
UVA.lamps <- c("qpanel.uva340")
for (lamp in UVA.lamps) {
  lamp.plotter(lamp.name = lamp)
}
```



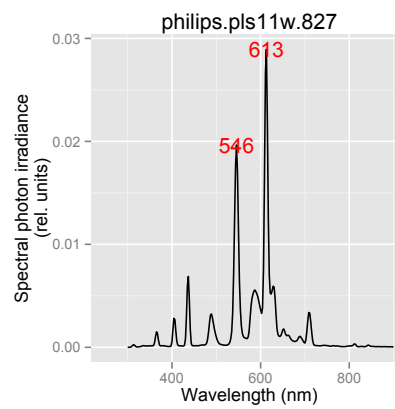
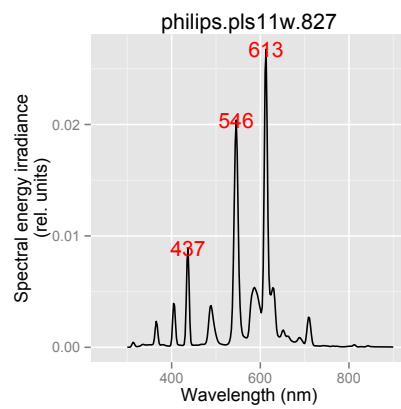
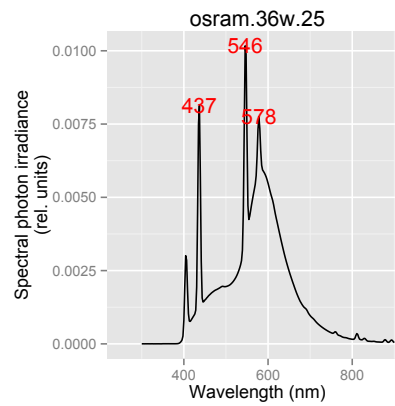
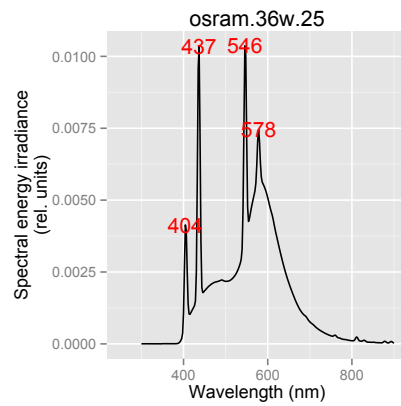
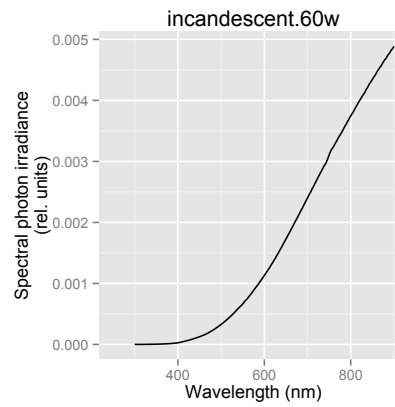
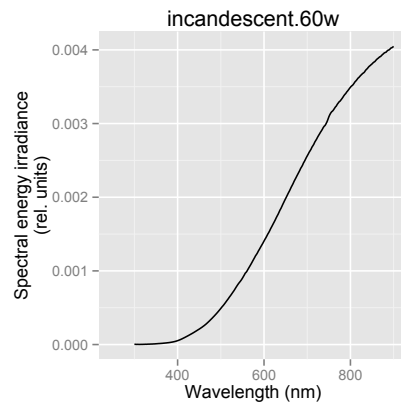
4 Narrow spectrum VIS lamps

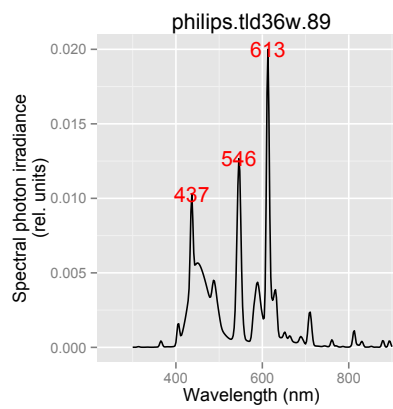
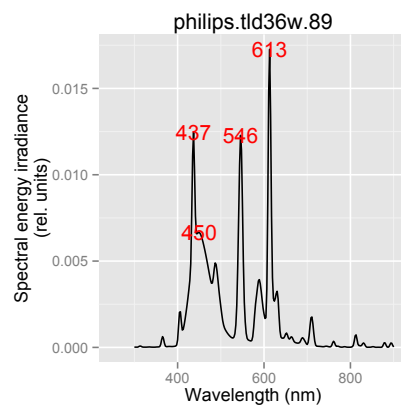
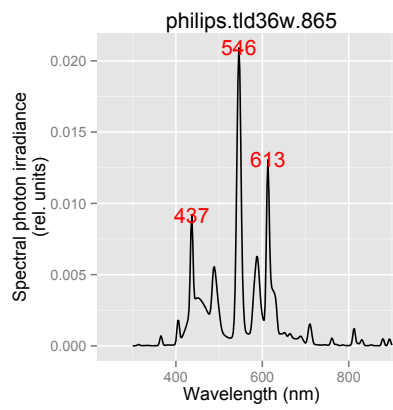
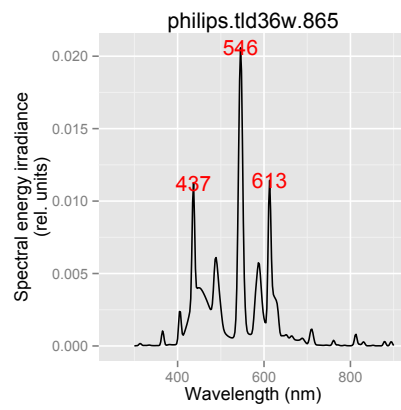
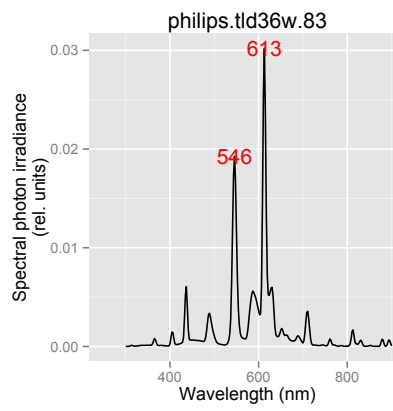
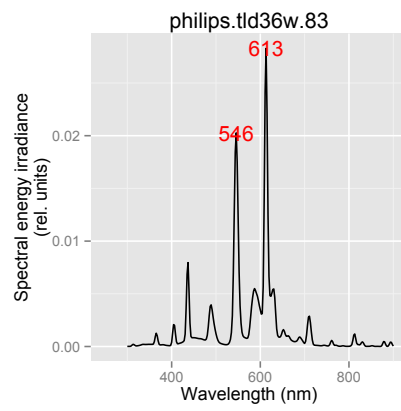
```
colour.lamps <- c("philips.tld36w.15", "philips.tld36w.18")
for (lamp in colour.lamps) {
  lamp.plotter(lamp.name = lamp)
}
```

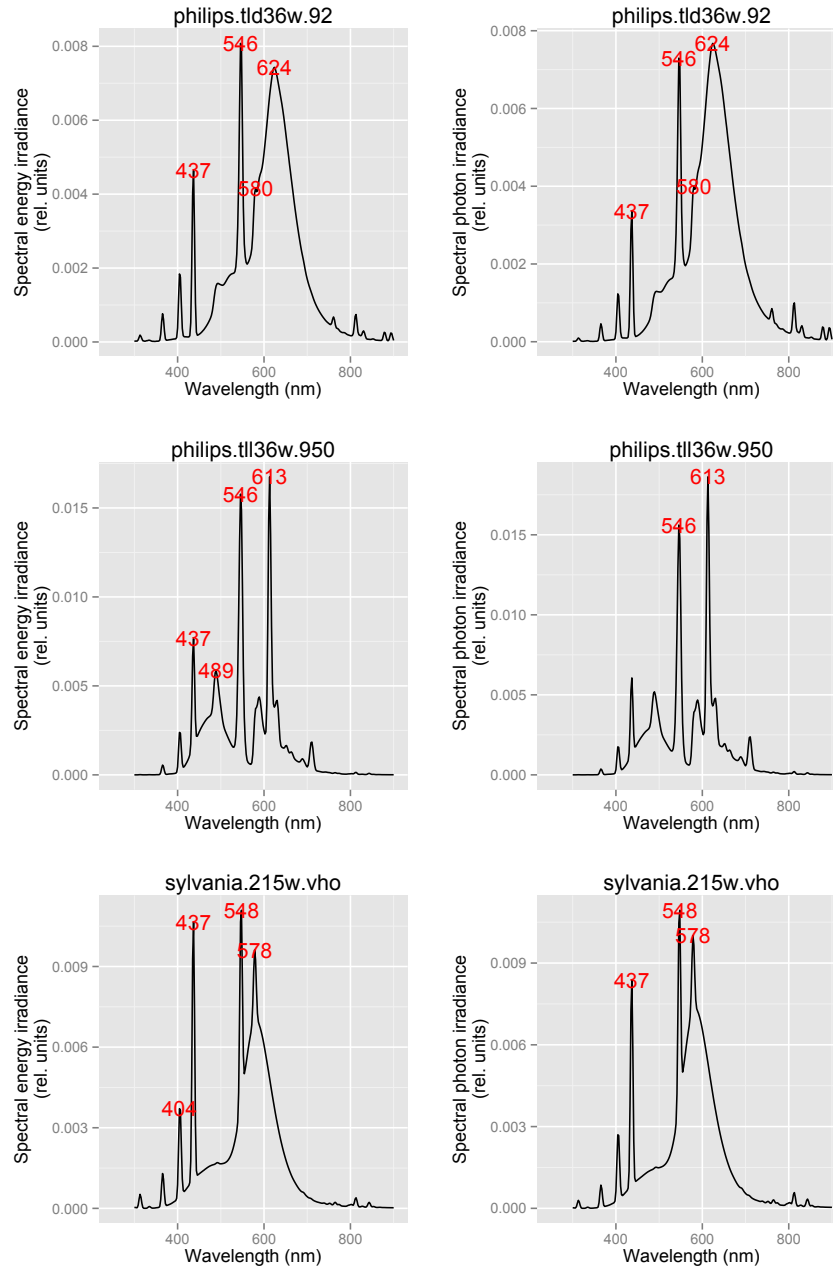


5 Broad VIS lamps

```
white.lamps <- c("incandescent.60w", "osram.36w.25", "philips.pls11w.827", "philips.tld36w.83",
  "philips.tld36w.865", "philips.tld36w.89", "philips.tld36w.92", "philips.tl136w.950",
  "sylvania.215w.vho")
for (lamp in white.lamps) {
  lamp.plotter(lamp.name = lamp)
}
```







6 Calibration lamps


```

FEL.data <- FEL_spectrum(250:900)
D2.data <- D2_spectrum(250:900)
calibration.lamps <- c("FEL", "D2")
for (lamp in calibration.lamps) {
  lamp.plotter(lamp.name = lamp, scaled = NULL)
}

```

