

Data manipulation with Rcell (Version 1.1-7)

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1 Introduction

Once you have your data loaded into **R**, you can filter it and plot it as shown in “Getting Started with Rcell”. To read that document type in the console

```
> vignette('Rcell')
```

But many times we want to do some manipulation or transformations on the data before plotting it. In this document you’ll see how this can be done using **Rcell**.

2 Transforming variables

If you haven’t done so, load the **Rcell** package and the example dataset with

```
> library(Rcell)
> data(ACL394filtered)
```

The easiest way to modify your dataset is to create new variables from existing ones. For example, its desirable to correct the fluorescence measure of a cell by the background fluorescence. To do this for the YFP channel we can use the `f.bg.y` variable, that contains the most common value (mode) for pixels not associated with any cell. If a cell has no fluorophores, we expect it to have a total fluorescence equivalent to `f.bg.y` times the number of pixels of the cell, `a.tot`. So the background corrected fluorescence can be calculated as `f.tot.y - f.bg.y*a.tot`. To create a new variable called `f.total.y` with the corrected value for fluorescence we can use the `transform` function. As all other **Rcell** functions, the first argument is the `cell.data` object to transform.

```
> X<-transform(X,f.total.y=f.tot.y - f.bg.y*a.tot)
```

Once created, you can use the new variable as any other variable of the dataset. You can create several variables in a single call to `transform`, as shown next for the fluorescence density variables.

```
> X<-transform(X,f.density.y=f.tot.y/a.tot,f.density.c=f.tot.c/a.tot)
```

You can keep track of the variables you’ve created with the `summary` function, that will display among other things the “transformed” variables with their definition.

```
> summary(X)
```

pos	alpha.factor
1	1.25
2	1.25
3	1.25
8	2.50
9	2.50
10	2.50
15	5.00
16	5.00
17	5.00
22	10.00
23	10.00
24	10.00
29	20.00
30	20.00
31	20.00

Table 1: example data.frame to merge

3 Merging variables

Sometimes there is no formula to specify the new variable you want to create. For example, you might want to create a variable that describes the treatment each position received. In the example dataset (`help(ACL394)`) each position received a different dose of alpha-factor pheromone, according to the Table 1.

You can create this table in Excel¹ and save it as a tab delimited text file. If you name it “mytable.txt”, then you can load it into **R** with `read.table`. The best option is to save the file in your working directory, or to change your working directory to where you saved the file (see `?setwd`).

```
> mytable<-read.table("mytable.txt",head=TRUE)
```

If the first row of your text file contains the column names (recommended), you have to set `head` to `TRUE` in `read.table`. Once loaded you can add the new data to your dataset using the `merge` function. This function looks for common variables between `X` and `mytable` and, if it finds them it merges the dataset according to those common variables. Be aware that the names of the columns of `mytable` have to match EXACTLY² to the variables of `X`³. In this case it will merge by `pos`. You can also specify the variable to merge by with the `by` argument.

```
> X<-merge(X,mytable)
```

```
merging by pos
```

```
merged vars:
```

```
alpha.factor: numeric w/values 1.25, 2.5, 5, 10, 20
```

4 Transform By

A common transformation is normalization, i.e. dividing the value of a variable by the “basal” level. For example, we might be interested in the fold increase of YFP fluorescence through time. So we need to divide the measured value at each time by the value at time zero, and we need to do this for every cell. How can we do this? The steps we should follow are the following:

¹or from R: `mytable<-data.frame(pos=with(X,unique(pos)),alpha.factor=rep(c(1.25,2.5,5,10,20),each=3))`

²**R** is case-sensitive so “pos” is different to “Pos”

³You can see these variables with `summary(X)`

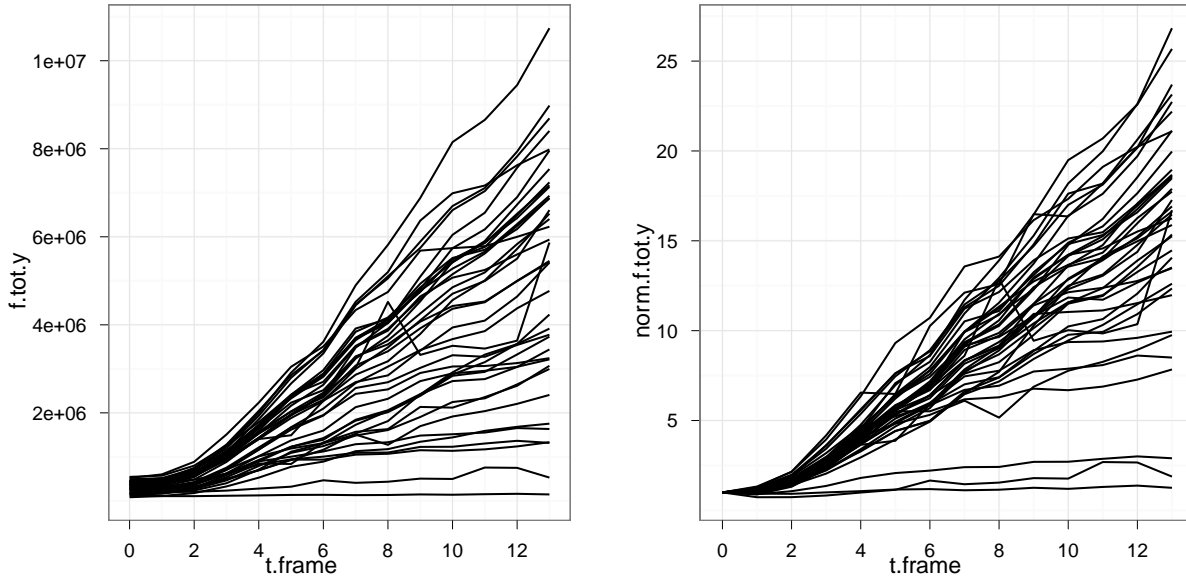


Figure 1: Left: raw single cell time course for YFP fluorescence. Right: Same data normalized to each cells value at time zero.

1. Divide the dataset by cell, creating a table for each cell.
2. Identify the value of fluorescence for time zero.
3. Create a new variable by dividing the fluorescence at each time by the value at time zero.
4. Join the cells datasets back together to retrieve the original dataset with the new variable.

All these steps are done by the function `transform.by`, but it requires information on how each step should be done. For the first step, it needs to know how to partition the dataset. This is specified by passing a quoted list of variable, whos combination of levels specify a group. For example, if you want to divide the dataset by position, the second argument of `transform.by` should be `.(pos)`. If you want to divide your dataset by cell use `.(pos,cellID)`. Note that cells in different position can have the same cellID, so the combination of `pos` and `cellID` uniquely identifies a cell. The variable `ucid` (for Unique Cell ID) is another way to uniquely identify a cell. Next we need to specify the name of the new variable to be created (`norm.f.tot.y` for example), and the definition for this variable, `f.tot.y/f.tot.y[t.frame==0]`. With the square brackets we are selecting the value of `f.tot.y` when `t.frame` is zero. Remember to use the logical operator `==` and not the assignation operator `=` within the brackets.

```
> X<-transform.by(X,.(pos,cellID),norm.f.tot.y=f.tot.y/f.tot.y[t.frame==0])
```

You can see the raw and normalized data in Figure 1. Another way to normalize the data, is dividing by the mean of the first three values.

```
> X<-transform.by(X,.(pos,cellID),norm2.f.tot.y=f.tot.y/mean(f.tot.y[t.frame<=2]))
```

5 Aggregating your data

To calculate summary statistics you can use the `aggregate` function, that returns an aggregated table. That means that the value of each cell of this aggregated table is calculated from more than one cell of the original table. For example you might be interested in getting the mean YFP fluorescence for each pheromone dose. `aggregate` accepts two notations that give equivalent results.

```
> aggregate(X,.(alpha.factor),select="f.total.y")
> aggregate(X,f.total.y~alpha.factor) #formula notation
```

	AF.nM	f.total.y
1	1.25	1071898
2	2.50	1622198
3	5.00	2234246
4	10.00	2393427
5	20.00	2377602

You can calculate other statistics using the `FUN` argument, and you can include more than one variable. Here we calculate the median for `f.tot.y`, `f.tot.c` and `a.tot`. Note the use of the wildcard in the `select` argument.

```
> aggregate(X,.(alpha.factor),select=c("f.tot.*","a.tot"),FUN=median)
```

	alpha.factor	f.tot.c	f.tot.y	a.tot
1	1.25	1047808.5	1212391	415.5
2	2.50	1055751.0	1564543	415.0
3	5.00	1037465.0	2032817	407.0
4	10.00	1001638.5	2224172	398.0
5	20.00	961167.5	2083660	380.0

The partition of the dataset can be done by more than one variable, for example by dose and time. Using the function `funstofun` from the **reshape** package, you can calculate more than one statistic at once.

```
> aggregate(X,f.density.y~t.frame+alpha.factor,FUN=funstofun(median,sd),subset=t.frame%%3==0)
```

	t.frame	alpha.factor	f.density.y.median	f.density.y.sd
1	0	1.25	1037.6060	162.4781
2	3	1.25	2103.6828	420.8901
3	6	1.25	3646.6715	892.4902
4	9	1.25	3916.8426	1112.6042
5	12	1.25	4081.6656	1295.7236
6	0	2.50	1043.6030	141.9683
7	3	2.50	2283.6054	577.7679
8	6	2.50	4663.5886	1387.7653
9	9	2.50	5809.6937	1895.5383
10	12	2.50	6715.9285	2200.4825
11	0	5.00	1057.4033	163.5038
12	3	5.00	2519.7817	662.3112
13	6	5.00	5619.7677	1633.4406
14	9	5.00	8206.8911	2418.8710
15	12	5.00	9600.1285	2859.8918
16	0	10.00	1009.4154	131.5840
17	3	10.00	2686.9785	508.3823

18	6	10.00	5883.8167	1218.5057
19	9	10.00	9131.4531	2018.9390
20	12	10.00	10792.7466	2478.7091
21	0	20.00	1022.5990	120.6829
22	3	20.00	2730.6114	571.0274
23	6	20.00	6440.7103	1645.1310
24	9	20.00	9512.2292	2497.7941
25	12	20.00	11746.9381	3383.9809

6 Evaluating expressions in your dataset

Using the `with` function, you can evaluate a expression in a enviroment created from your dataset. That means that you can use the names of your variables directly, without any prefix. For example to calculate the mean of `f.tot.y` from position 1

```
> with(X,mean(f.tot.y[pos==1]))
[1] 1372297
```

If you don't use `with` you have to write the full identifier of the variable, and the code becomes longer and harder to understand. For exaple, the same result can be obtained with

```
> mean(X$data$f.tot.y[X$data$pos==1])
```

7 Exporting your data

Although you can do much of your analysis using Rcell functions, you might need to export the data to some other application or use another package within **R**. To retrieve the entire dataset in a `data.frame`, use the double square brackets notation. This returns the registers that pass the QC.filter.

```
> df<-X[[ ]]
```

This dataset is usually big, and has many varaibles or registers you are not interested in. You can subset the datset as you would a `data.frame` (but using double brackets)

```
> df<-X[[pos==1,c("cellID","f.tot.y","a.tot")]]
```

You can then save the `data.frame` to a file with `write.table`, or use it in another **R** package.

For some kinds of data analysis you need your data in a different form than the one **Rcell** uses. You can use the `reshape` function to reshape your data. For instance, a common restructuring is to display time as different columns, and individual cells as different rows. You can obtain this sort of `data.frames` with the following command.

```
> reshape(X,pos+cellID~variable+t.frame,select="f.tot.y",subset=pos<=2&cellID<=10&t.frame%%2==0)
```

	pos	cellID	f.tot.y_0	f.tot.y_2	f.tot.y_4	f.tot.y_6	f.tot.y_8	f.tot.y_10	f.tot.y_12
1	1	1	378752	748712	1350707	2028179	2155404	2072739	2214004
2	1	2	176429	300842	448582	535334	549019	562208	512430
3	1	3	384393	665472	1234888	1913377	2036718	2217148	2071306
4	1	4	245876	510412	887509	1493615	1692185	1987466	2137951
5	1	6	347597	629056	1000791	1533244	1788453	2188437	2381668
6	1	7	325715	558893	998972	1740997	2080686	2575011	2845928
7	1	8	276242	481790	842095	1310683	1464268	1735160	1826068

8	1	10	314574	559742	1050029	1832519	2166170	2765528	2981627
9	2	2	387551	620656	1049458	1327046	1317000	1409672	1742833
10	2	3	428014	655421	1239405	1616264	1942105	2268256	2583064
11	2	4	452047	718126	1381880	1808801	2260311	2624726	3127893
12	2	5	330852	228660	285463	294746	357041	354951	308372
13	2	6	166711	139176	173661	184098	201449	202782	223541
14	2	7	453124	665657	1288083	1847586	2532047	2676930	2793809
15	2	8	137043	232664	448551	652693	830410	957173	1094667
16	2	9	504990	632012	961822	1340065	1640487	1769006	1929912
17	2	10	317594	440622	763946	1221629	1711411	1788251	1834715

see `help(reshape.cell.data)` for more details.

References

- Pau, Fuchs et al. (2010). EBImage: an R package for image processing with applications to cellular phenotypes. *Bioinformatics*, 26(7):979-981.
- Colman-Lerner, Gordon et al. (2005). Regulated cell-to-cell variation in a cell-fate decision system. *Nature*, 437(7059):699-706.
- Chernomoretz, Bush et al. (2008). Using Cell-ID 1.4 with R for Microscope-Based Cytometry. *Curr Protoc Mol Biol.*, Chapter 14:Unit 14.18.