

Lecture 6

Split-Apply-Combine

BIOF 339

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Goals for today

- Learn how to merge data sets
- Learn how to reshape data sets from wide to long
- Split-apply-combine
 - Split a dataset into a list of several datasets
 - Do something to each dataset
 - Put the results back together
- Use it for
 - Running tests for many variables
 - Visualizing data with p-value annotation

The data for today

This data set is taken from a breast cancer proteome database available [here](#) and modified for this exercise.

- Clinical data: [CSV](#) | [XLSX](#)
- Proteome data: [CSV](#) | [XLSX](#)

The data for today

```
# Excel
library(readxl)
clinical_data <- read_excel('lecture6_data/BreastCancer_Clinical.xlsx')

# CSV
clinical_data <- read.csv('lecture6_data/BreastCancer_Clinical.csv',
                          stringsAsFactors=F)
```

```
# Complete.TCGA.ID Gender Age.at.Initial.Pathologic.Diagnosis ER.Status
# 1 TCGA-A2-A0CM FEMALE 40 Negative
# 2 TCGA-BH-A18Q FEMALE 56 Negative
# PR.Status HER2.Final.Status Tumor Tumor..T1.Coded Node Node.Coded
# 1 Negative Negative T2 T_Other N0 Negative
# 2 Negative Negative T2 T_Other N1 Positive
# Metastasis Metastasis.Coded AJCC.Stage Converted.Stage
# 1 M0 Negative Stage IIA Stage IIA
# 2 M0 Negative Stage IIB No_Conversion
# Survival.Data.Form Vital.Status Days.to.Date.of.Last.Contact
```

The data for today

```
expression_data <- read.csv('lecture6_data/BreastCancer_Expression.csv',  
                             stringsAsFactors=F)  
head(expression_data[,1:5],2)
```

```
#      TCGA_ID NP_958782 NP_958785 NP_958786 NP_000436  
# 1 TCGA-AO-A12D  1.096131  1.111370  1.111370  1.107561  
# 2 TCGA-C8-A131  2.609943  2.650422  2.650422  2.646374
```

Merging data

Note that

1. The names of the ID columns are different
2. They are in different orders

The `merge` function takes care of both of these issues

Merging data

```
final_data = merge(clinical_data,  
                    expression_data[,1:5],  
                    by.x = 'Complete.TCGA.ID',  
                    by.y = 'TCGA_ID')  
head(final_data,2)
```

```
#      Complete.TCGA.ID Gender Age.at.Initial.Pathologic.Diagnosis ER.Status  
# 1      TCGA-A2-A0CM FEMALE                                     40 Negative  
# 2      TCGA-A2-A0D2 FEMALE                                     45 Negative  
#      PR.Status HER2.Final.Status Tumor Tumor..T1.Coded Node Node.Coded  
# 1 Negative           Negative    T2      T_Other    N0    Negative  
# 2 Negative           Negative    T2      T_Other    N0    Negative  
#      Metastasis Metastasis.Coded AJCC.Stage Converted.Stage  
# 1      M0           Negative    Stage IIA      Stage IIA  
# 2      M0           Negative    Stage IIB      Stage IIA  
#      Survival.Data.Form Vital.Status Days.to.Date.of.Last.Contact  
# 1      followup      DECEASED                                     754  
# 2      followup      LIVING                                       1027
```

Merging data

```
dim(clinical_data)
```

```
# [1] 77 30
```

```
dim(expression_data)
```

```
# [1] 83 11
```

```
dim(final_data)
```

```
# [1] 80 34
```

Note that we have extra rows, which usually means **duplication of rows/ids**. Something to check on.

Reshaping data

R usually can split data on rows.

If we want to split on variables (columns), we have to transform the data from *wide* to *long* so that each row is the data for one individual *for one variable*.

First, of course, we have to select the variables we want to include.

Selecting variables

```
head(final_data,2)
```

```
# Complete.TCGA.ID Gender Age.at.Initial.Pathologic.Diagnosis ER.Status
# 1 TCGA-A2-A0CM FEMALE 40 Negative
# 2 TCGA-A2-A0D2 FEMALE 45 Negative
# PR.Status HER2.Final.Status Tumor Tumor..T1.Coded Node Node.Coded
# 1 Negative Negative T2 T_Other N0 Negative
# 2 Negative Negative T2 T_Other N0 Negative
# Metastasis Metastasis.Coded AJCC.Stage Converted.Stage
# 1 M0 Negative Stage IIA Stage IIA
# 2 M0 Negative Stage IIB Stage IIA
# Survival.Data.Form Vital.Status Days.to.Date.of.Last.Contact
# 1 followup DECEASED 754
# 2 followup LIVING 1027
# Days.to.date.of.Death OS.event OS.Time PAM50.mRNA
# 1 754 1 754 Basal-like
# 2 NA 0 1027 Basal-like
# SigClust.Unsupervised.mRNA SigClust.Intrinsic.mRNA miRNA.Clusters 10/31
```

Selecting variables

```
library(dplyr)
selected_data <- select(final_data, Complete.TCGA.ID, ER.Status,
                        starts_with('NP'))
head(selected_data, 2)
```

```
#   Complete.TCGA.ID ER.Status NP_958782 NP_958785 NP_958786 NP_000436
# 1   TCGA-A2-A0CM   Negative 0.6834035 0.6944241 0.6980976 0.68707705
# 2   TCGA-A2-A0D2   Negative 0.1074909 0.1041645 0.1074909 0.09751166
```

Reshaping data

We have two goals here:

1. Keep the ER status aligned with each expression level, so we would need to repeat the column
2. Make each row have the ER status for an individual and the corresponding expression for one protein

Reshaping data

```
library(reshape2)

reshaped_data <- melt(selected_data,
                      id.vars=c( 'Complete.TCGA.ID', 'ER.Status' ))
head(reshaped_data, 5)
```

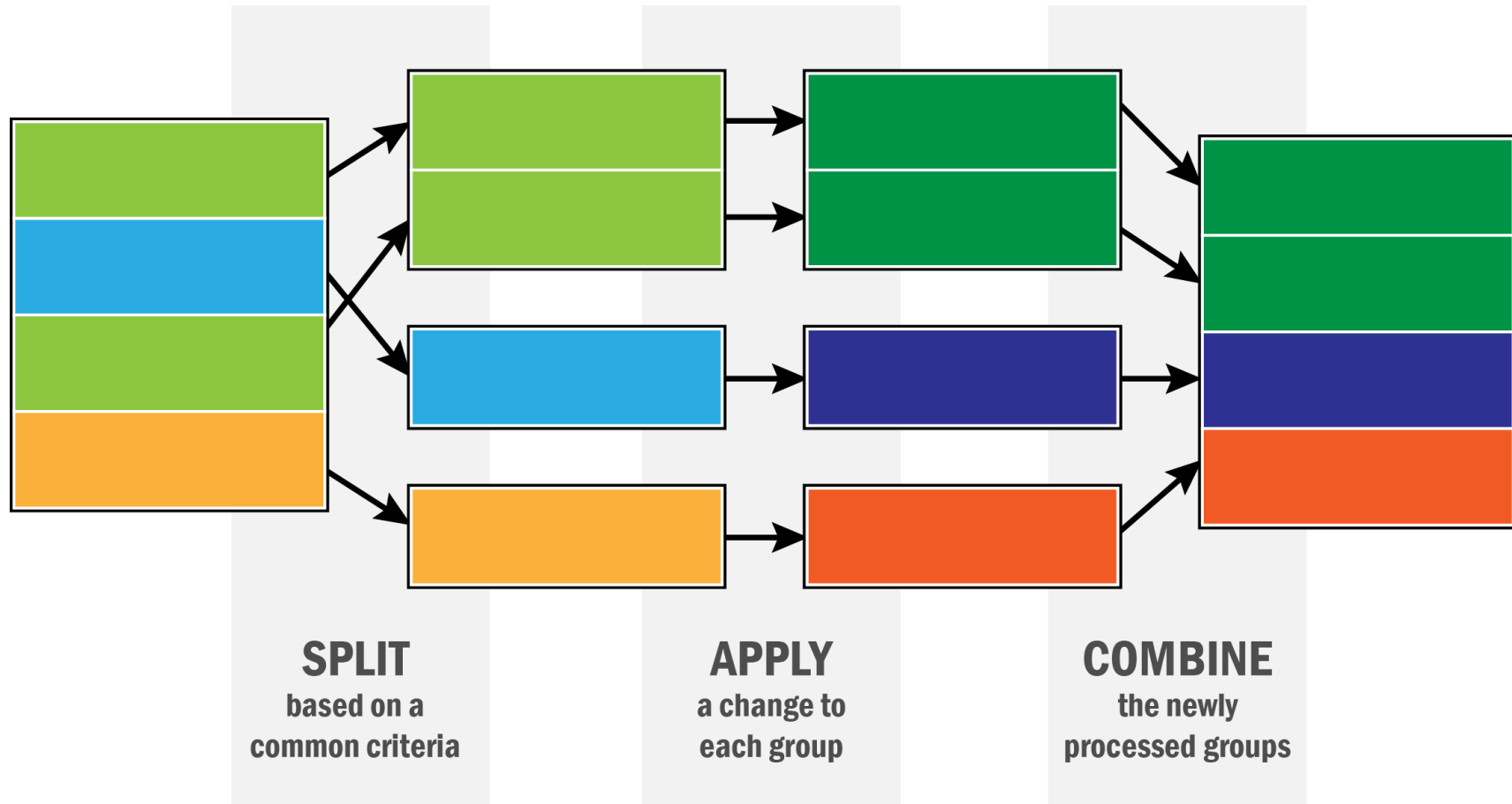
#	Complete.TCGA.ID	ER.Status	variable	value
# 1	TCGA-A2-A0CM	Negative	NP_958782	0.6834035
# 2	TCGA-A2-A0D2	Negative	NP_958782	0.1074909
# 3	TCGA-A2-A0EQ	Negative	NP_958782	-0.9126703
# 4	TCGA-A2-A0EV	Positive	NP_958782	0.4529859
# 5	TCGA-A2-A0EX	Positive	NP_958782	1.1851082

Reshaping data

```
reshaped_data <- arrange(reshaped_data, Complete.TCGA.ID) # from dplyr  
head(reshaped_data, 5)
```

#	Complete.TCGA.ID	ER.Status	variable	value
# 1	TCGA-A2-A0CM	Negative	NP_958782	0.6834035
# 2	TCGA-A2-A0CM	Negative	NP_958785	0.6944241
# 3	TCGA-A2-A0CM	Negative	NP_958786	0.6980976
# 4	TCGA-A2-A0CM	Negative	NP_000436	0.6870771
# 5	TCGA-A2-A0D2	Negative	NP_958782	0.1074909

Split-apply-combine



Splitting data by protein

```
split_data <- split(reshaped_data, reshaped_data$variable)  
class(split_data)
```

```
# [1] "list"
```

```
length(split_data)
```

```
# [1] 4
```

```
names(split_data)
```

```
# [1] "NP_958782" "NP_958785" "NP_958786" "NP_000436"
```


Splitting data

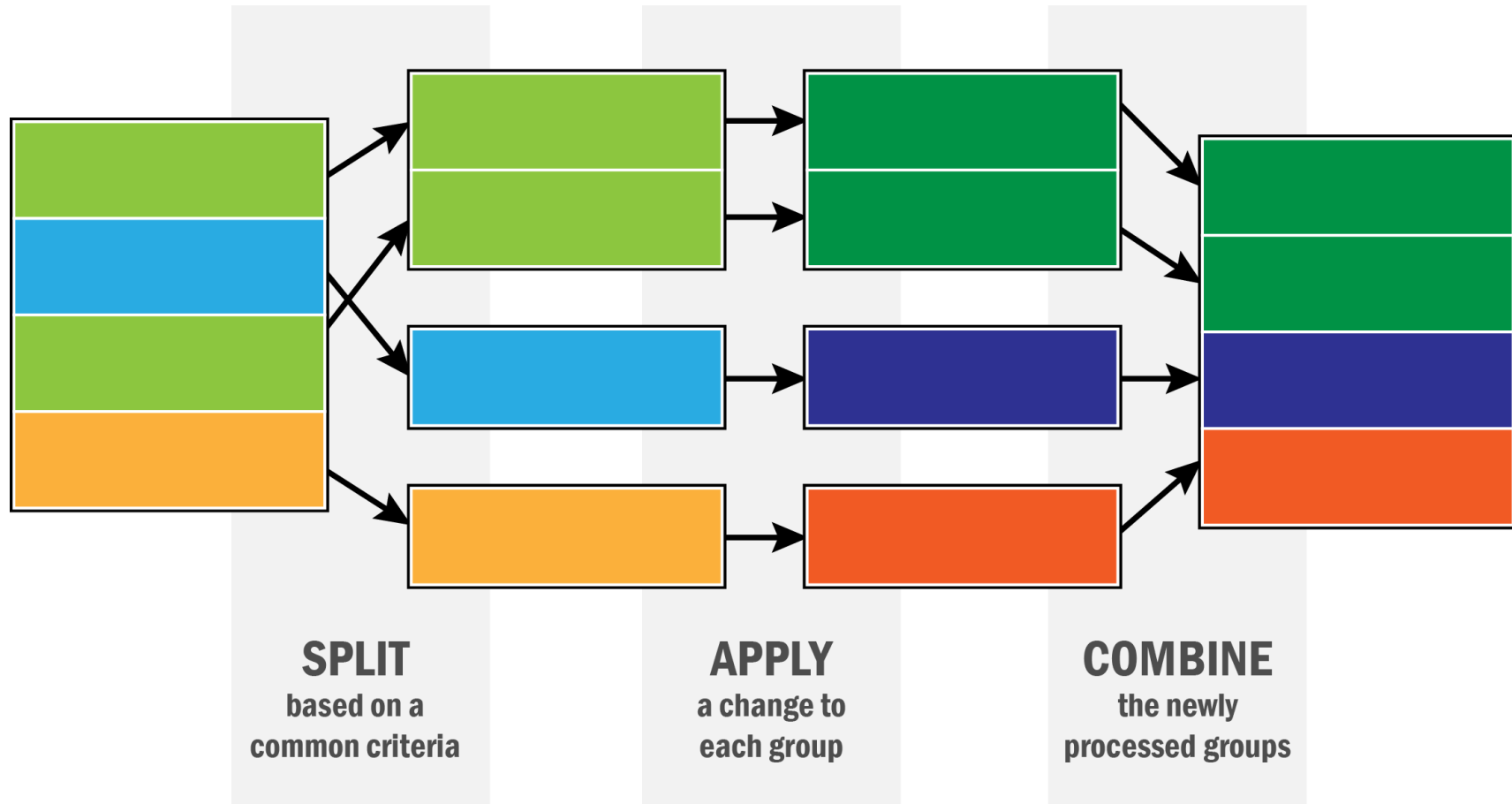
```
head(split_data[['NP_958782']],8)
```

#	Complete.TCGA.ID	ER.Status	variable	value
# 1	TCGA-A2-A0CM	Negative	NP_958782	0.6834035
# 5	TCGA-A2-A0D2	Negative	NP_958782	0.1074909
# 9	TCGA-A2-A0EQ	Negative	NP_958782	-0.9126703
# 13	TCGA-A2-A0EV	Positive	NP_958782	0.4529859
# 17	TCGA-A2-A0EX	Positive	NP_958782	1.1851082
# 21	TCGA-A2-A0EY	Positive	NP_958782	1.1748810
# 25	TCGA-A2-A0SW	Positive	NP_958782	-0.4877725
# 29	TCGA-A2-A0SX	Negative	NP_958782	-0.3985598

```
class(split_data[['NP_958782']])
```

```
# [1] "data.frame"
```

Applying a function to the split data



Applying a function to the split data

We're going to run t-tests to see if the expression for each protein differs by ER status

First we create a function that we'll run on every protein-specific data set

```
run_t_test <- function(d){  
  test <- t.test(value ~ ER.Status, data=d)  
  pvalue <- test$p.value  
  return(pvalue)  
}
```

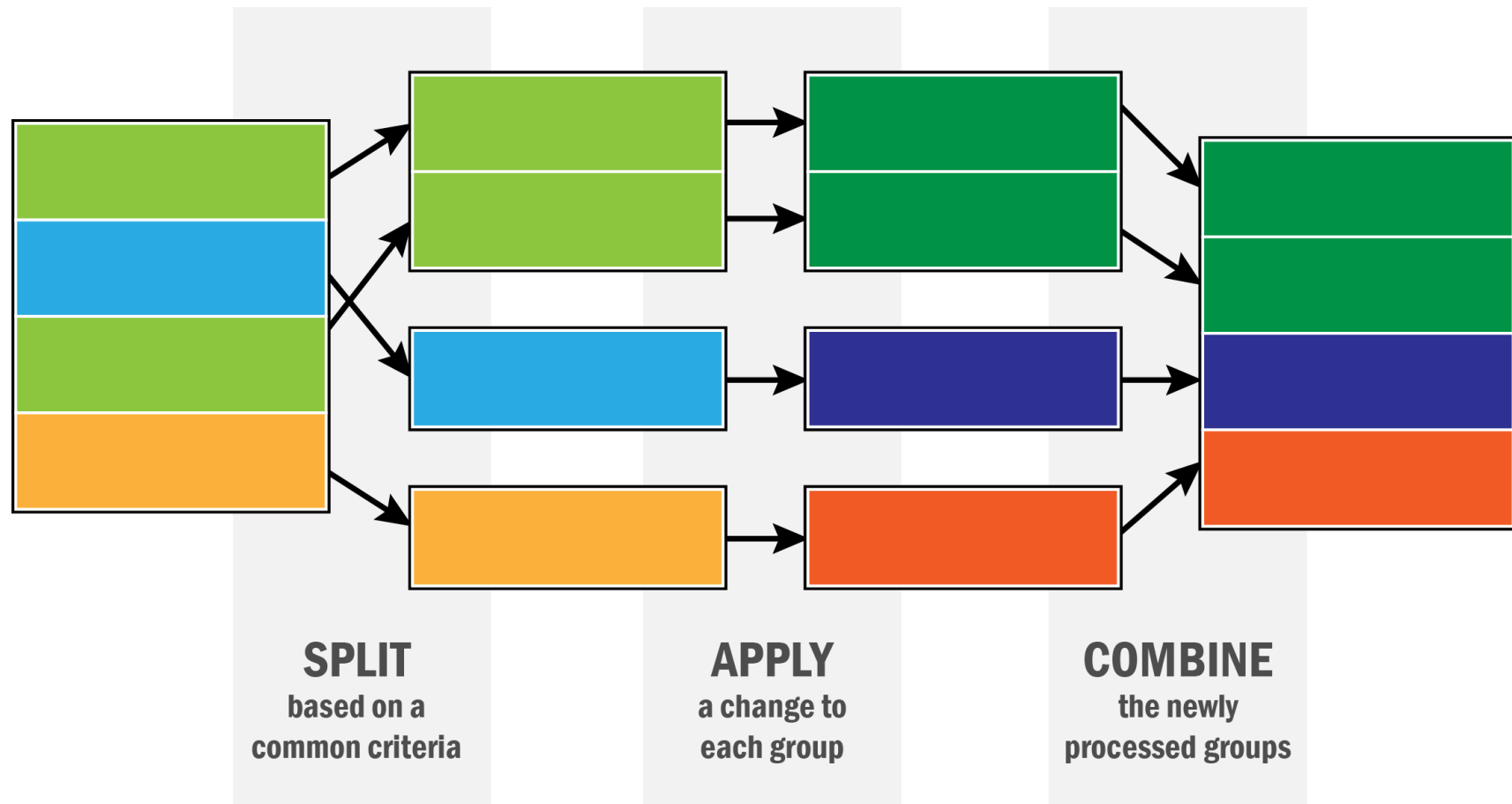
Applying a function to the split data

Now we can apply the same function to each element of the list of data frames using the `lapply` command

```
pvalues <- lapply(split_data, run_t_test)
pvalues
```

```
# $NP_958782
# [1] 0.5287476
#
# $NP_958785
# [1] 0.5243093
#
# $NP_958786
# [1] 0.519653
#
# $NP_000436
# [1] 0.5337173
```

Combining the data



Combining the data

```
pvalues_final <- unlist(pvalues)
pvalues_final
```

```
# NP_958782 NP_958785 NP_958786 NP_000436
# 0.5287476 0.5243093 0.5196530 0.5337173
```

Combining the data

```
pvalues_final <- plyr::ldply(pvalues) # Use ldply from the plyr package  
pvalues_final
```

```
#           .id      V1  
#  1 NP_958782 0.5287476  
#  2 NP_958785 0.5243093  
#  3 NP_958786 0.5196530  
#  4 NP_000436 0.5337173
```

The `ldply` function inputs a list and exports a data.frame with the elements concatenated if possible.

Plotting

The data

```
head(reshaped_data, 5)
```

#	Complete.TCGA.ID	ER.Status	variable	value
# 1	TCGA-A2-A0CM	Negative	NP_958782	0.6834035
# 2	TCGA-A2-A0CM	Negative	NP_958785	0.6944241
# 3	TCGA-A2-A0CM	Negative	NP_958786	0.6980976
# 4	TCGA-A2-A0CM	Negative	NP_000436	0.6870771
# 5	TCGA-A2-A0D2	Negative	NP_958782	0.1074909

A panel of plots

```
library(ggplot2)
ggplot(reshaped_data, aes(x = ER.Status, y = value))+
  geom_boxplot()+
  facet_wrap(~variable, ncol=2)
```

Adding p-values to the plot

```
names(pvalues_final) <- c('variable', 'pvalue')
ggplot(reshaped_data, aes(x = ER.Status, y = value))+
  geom_boxplot()+
  facet_wrap(~variable, ncol=2)+
  geom_text(data=pvalues_final, aes(x=1.5, y=2, label=pvalue))
```

Formatting p-values

```
pvalues_final[, 'pvalue'] <- format.pval(pvalues_final[, 'pvalue'],  
                                         digits=3)  
ggplot(reshaped_data, aes(x = ER.Status, y = value))+  
  geom_boxplot()+  
  facet_wrap(~variable, ncol=2)+  
  geom_text(data=pvalues_final, aes(x=1.5, y=2, label=pvalue))
```

Formatting p-values

```
pvalues_final <- mutate(pvalues_final,  
                        pvalue = format.pval(pvalue, digits=3))  
ggplot(reshaped_data, aes(x = ER.Status, y = value))+  
  geom_boxplot()+  
  facet_wrap(~variable, ncol=2)+  
  geom_text(data=pvalues_final, aes(x=1.5, y=2, label=pvalue))
```

The dplyr package

Action words

`dplyr` basically gives you 6 actions to do on `data.frame` objects:

1. `mutate`: Change particular variables
2. `select`: Select (or deselect) variables
3. `arrange`: Order by some variables
4. `filter`: Select rows by some criteria
5. `group_by`: Group by some variable (so the split part of our exercise)
6. `summarise`: Summarise a variable using some function