

Using uwIntroStats  
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## 1 Introduction

## 2 Preparing uwIntroStats

Before we can dive in and run any analyses, we first need to install the package. This is done via

```
> install.packages("uwIntroStats")
```

Regardless of the graphical user interface (GUI) that you are using, R will prompt you to select a CRAN mirror. It is essentially asking you where you want to download the package files from. Select the mirror closest to you - for us at the University of Washington it is **WA(1)** or the Fred Hutchinson Cancer Research Center (FHCRC) - and the package will download and say that it has installed. Now each time we open a new R session (whether that is at the command line, a new RGui window, or a new RStudio window) we need to load the package for use.

The **uwIntroStats** package relies on five other packages. These other packages provide key functions that the **uwIntroStats** package uses or adds functionality to. We must install these packages like we did above if we have not installed them previously, and then load **uwIntroStats**. While these packages do not need to be loaded every time (in fact, some are only used for specific functions) it is good practice to load them for the R session where you need to use **uwIntroStats**.

```
> library(Exact)
> library(geepack)
> library(plyr)
> library(sandwich)
> library(survival)
> library(uwIntroStats)
```

Don't worry about the warning message for now; that will be covered in section 3.2. Last, we load the data, **mri** that we will be using throughout this document. Information about the dataset can be found at <http://www.emersonstatistics.com/datasets/mri.pdf>. Since the data is part of the package, we can load it via

```
> data(mri)
```

The **uwIntroStats** package should be used for descriptive statistics, basic plotting (like scatterplots and boxplots), and regression analyses. The following sections will go through examples of these tasks, in addition to pointing out how our package differs from base R and other existing packages. We will assume familiarity with basic data manipulation and statistical tasks (for a refresher, see <http://www.emersonstatistics.com/GeneralMaterials/R/IntroToR.pdf>).

## 3 Descriptive Statistics

Descriptive statistics are an important part of any analysis. Often, they are just as important for the statistician or data analyst as they are for the scientific collaborators on a project. For example, descriptive statistics can help identify errors in the data, like observations that are particularly unusual (usually far out of the expected or observed range). They can also help to identify patterns of missing data, examine the study population, and identify aspects of the data that might lead to statistical issues. Descriptive statistics can also unearth relationships that had not previously been considered for

analysis, thereby generating new studies. More information on this, and an outline of an approach to analysing a dataset, was written by Scott Emerson and can be found at <http://www.emersonstatistics.com/GeneralMaterials/analysis.pdf>.

### 3.1 The basics: `descrip()`

The `uwIntroStats` package was designed with descriptive statistics in mind. The basic function for descriptive statistics is `descrip()`. This function takes in an arbitrary number of variables, and by default calculates the number of observations, the number of missing values, the mean, standard deviation, minimum value, maximum value, and the 25th, 50th, and 75th percentiles of each variable. For instance, if we wanted a quick glance at the `mri` data, we could type

```
> descrip(mri)
```

	N	Msnrg	Mean	Std Dev	Min	25%	Mdn	75%	Max
ptid:	735	0	368.0	212.3	1.000	184.5	368.0	551.5	735.0
mridate:	735	0	76423	31896	10192	66642	80992	91392	1.232e+05
age:	735	0	74.57	5.451	65.00	71.00	74.00	78.00	99.00
male:	735	0	0.4980	0.5003	0.0000	0.0000	0.0000	1.000	1.000
race:	735	0	1.318	0.6659	1.000	1.000	1.000	1.000	4.000
weight:	735	0	159.9	30.74	74.00	138.5	158.0	179.0	264.0
height:	735	0	165.8	9.710	139.0	158.0	165.9	173.2	190.5
packyrs:	735	1	19.60	27.11	0.0000	0.0000	6.500	33.75	240.0
yrsquit:	735	0	9.661	14.10	0.0000	0.0000	0.0000	18.50	56.00
alcoh:	735	0	2.109	4.852	0.0000	0.0000	0.01920	1.144	35.00
physact:	735	0	1.922	2.052	0.0000	0.5538	1.312	2.513	13.81
chf:	735	0	0.05578	0.2297	0.0000	0.0000	0.0000	0.0000	1.000
chd:	735	0	0.3347	0.6862	0.0000	0.0000	0.0000	0.0000	2.000
stroke:	735	0	0.2367	0.6207	0.0000	0.0000	0.0000	0.0000	2.000
diabetes:	735	0	0.1075	0.3099	0.0000	0.0000	0.0000	0.0000	1.000
genhlth:	735	0	2.588	0.9382	1.000	2.000	3.000	3.000	5.000
ldl:	735	10	125.8	33.60	11.00	102.0	125.0	147.0	247.0
alb:	735	2	3.994	0.2690	3.200	3.800	4.000	4.200	5.000
crt:	735	2	1.064	0.3030	0.5000	0.9000	1.000	1.200	4.000
plt:	735	7	246.0	65.80	92.00	201.8	239.0	285.0	539.0
sbp:	735	0	131.1	19.66	78.00	118.0	130.0	142.0	210.0
aai:	735	9	1.103	0.1828	0.3171	1.027	1.112	1.207	1.728
fev:	735	10	2.207	0.6875	0.4083	1.745	2.158	2.649	4.471
dsst:	735	12	41.06	12.71	0.0000	32.00	40.00	50.00	82.00
atrophy:	735	0	35.98	12.92	5.000	27.00	35.00	44.00	84.00
whgrd:	735	1	2.007	1.410	0.0000	1.000	2.000	3.000	9.000
numinf:	735	0	0.6109	0.9895	0.0000	0.0000	0.0000	1.000	5.000
volinf:	735	1	3.223	17.36	0.0000	0.0000	0.0000	0.09420	197.0

obstime:	735	0	1804	392.3	68.00	1837	1879	2044	2159
death:	735	0	0.1810	0.3852	0.0000	0.0000	0.0000	0.0000	1.000

This call gives us a quick look at the distribution of each variable in the sample, and can be easily exported to a word processing software. This is important, because displaying raw R output in a publication is not usually ideal, and taking the time to format the output is a pain.

Notice that there are a lot of variables measured in these data, and that some interesting phenomena are measured by multiple variables. For instance, smoking is measured by `packyrs` and `yrsquit` - if someone has zero pack-years (that is, they have never smoked) then they will also have zero years quit. Similarly, a current smoker of any amount of packs per year will have zero years quit. We can also look at the variables measured through blood samples, like LDL cholesterol (`ldl`) and get an idea of the range in our sample population. Last, in the variables like `male` which consist of only two values (sex was measured as male or female in this dataset), the mean tells us the proportion of our sample who was male (or for other binary variables, it will tell us the proportion which is coded as 1). The `descrip()` function works similarly to `summary()` in base R, but gives more information and returns it in a format that, as we described above, is easier to export from R.

Another powerful feature of `descrip()` is its ability to present stratified summaries, or summaries on a subset of the data. For instance, let's calculate descriptive statistics on the `age` variable within males and females:

```
> descrip(mri$age, strata = mri$male)
```

		N	Msng	Mean	Std Dev	Min	25%	Mdn	75%	Max
mri\$age:	All	735	0	74.57	5.451	65.00	71.00	74.00	78.00	99.00
mri\$age:	Str 0	369	0	74.41	5.258	65.00	71.00	73.00	78.00	91.00
mri\$age:	Str 1	366	0	74.73	5.642	66.00	71.00	74.00	78.00	99.00

```
> ## Call the summary function, to compare
> summary(mri$age)
```

	Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
	65.00	71.00	74.00	74.57	78.00	99.00

Here the row for `Str 0` corresponds to females (since they are coded as 0 for `male`), and males are `Str 1`. If we further wanted to restrict to only those people who were over 75 years old, we could use the `subset` argument:

```
> descrip(mri$age, strata = mri$male, subset = mri$age > 75)
```

		N	Msng	Mean	Std Dev	Min	25%	Mdn	75%	Max
mri\$age:	All	260	0	80.59	4.111	76.00	77.75	79.00	83.00	99.00
mri\$age:	Str 0	127	0	80.44	3.572	76.00	78.00	79.00	83.00	91.00
mri\$age:	Str 1	133	0	80.73	4.576	76.00	77.00	79.00	83.00	99.00

However, the `descrip()` function can only give us a fixed number of summaries (mean, count, etc). Oftentimes we need only a subset of these, or perhaps different summary measures. For this, we turn to more powerful functions.

### 3.2 Flexibility: `tableStat()` and `tabulate()`

The main draw of `tableStat()` and `tabulate()` is their ability to supply only a subset of the summary measures from `descrip()`. These functions focus on display and flexibility. The user supplies the format for the summary statistics to be displayed in, which makes exporting results from a call to

`tablestat()` or `tabulate()` even easier than it was for `descrip()`. Of the two, `tableStat()` is the base, and `tabulate()` adds some additional formatting and options. For example, let's say that we want the same summary statistics as we would get from `descrip()`, but we want to control the printout. For this, we use the `stat` argument. The options are presented in Table 1.

Name	Summary Measure	Name	Summary Measure
"count"	Number of observations on a variable	"sd"	Standard Deviation
"missing"	Number of missing observations	"variance"	Variance
"mean"	Arithmetic mean	"min"	Minimum value
"geometric mean"	Geometric mean	"max"	Maximum value
"median"	50th percentile	"quantiles"	Display quantiles
"probabilities"	Display percentiles	"mn(sd)"	Display Mean (SD) format
"range"	Maximum value - minimum value	"iqr"	Interquartile range (75th - 25th percentiles)
"all"	Display all summary measures	"row%"	Display row percentages
"col%"	Display column percentages	"tot%"	Display percentage of total

Table 1: Summary measures calculated by `tableStat()` and `tabulate()`.

The following gets us close to results we could display in a table for a publication. We want to have

*N : n; Mean (SD) : mn (sd); Range : max – min,*

which we get via:

```
> tableStat(mri$age, stat = "N: @count@; Mean (SD): @mean@ (@sd@); Range: @max@ - @min@")
Tabled descriptive statistics by strata
Call:
  tableStat(variable = mri$age, stat = "N: @count@; Mean (SD): @mean@ (@sd@); Range: @max@ - @min@")
  - NaN denotes strata with no observations
  - NA arises from missing or censored data

Format:  N: Cnt; Mean (SD): Mean (SD); Range: Max - Min

                                .ALL
N: 735.0; Mean (SD): 74.57 (5.451); Range: 99.00 - 65.00
```

Note that all of the values enclosed in @ symbols are from Table 1 and are run by `tableStat()`, while the other values in the string are used for formatting. This function can also take stratification variables; if we want to see the above stratified by sex, we use

```
> tableStat(mri$age, strata = mri$male, stat = "N: @count@; Mean (SD): @mean@ (@sd@); Range: @min@ - @max@")
Tabled descriptive statistics by strata
Call:
```

```
tableStat(variable = mri$age, strata = mri$male, stat = "N: @count@; Mean (SD): @mean@ (@sd@); Range: @min@ - @max@")
- NaN denotes strata with no observations
- NA arises from missing or censored data
```

Format: N: Cnt; Mean (SD): Mean (SD); Range: Min - Max

```

                                strata.0                                strata.1
N: 369.0; Mean (SD): 74.41 (5.258); Range: 65.00 - 91.00 N: 366.0; Mean (SD): 74.73 (5.642); Range: 66.00 - 99.00 N: 735.0; Mean (SD): 74

```

Now if we wanted to see this in table form, in perhaps a nicer layout, we could use the `tabulate()` function. When we loaded the `uwIntroStats` package at the beginning of this document, recall that we got a warning message from R saying

The following object is masked from ‘‘package:base’’:

```
tabulate
```

This means that our function `tabulate()` overwrites the function called `tabulatte()` in the base R package, which takes a numeric vector and counts the number of times each integer occurs in it. Our function does a similar task, but on combinations of strata. The syntax is very similar to the syntax for `tableStat()`, but rather than explicitly telling `tabulate()` the strata, it creates the tables with each dimension corresponding to a variable in the order they are entered. For instance, if we want a table of `age` (in the rows) and `male` (in the columns), we would write

```
> tabulate(mri$age, mri$male)
```

Call:

```
tabulate(mri$age, mri$male)
```

	mri\$male.0	mri\$male.1	mri\$male.ALL
mri\$age.65	2	0	2
mri\$age.66	2	1	3
mri\$age.67	18	15	33
mri\$age.68	20	21	41
mri\$age.69	19	19	38
mri\$age.70	30	30	60
mri\$age.71	37	39	76
mri\$age.72	31	28	59
mri\$age.73	27	28	55
mri\$age.74	29	26	55
mri\$age.75	27	26	53
mri\$age.76	11	17	28
mri\$age.77	18	19	37
mri\$age.78	18	19	37
mri\$age.79	19	13	32
mri\$age.80	8	15	23

mri\$age.81	10	10	20
mri\$age.82	10	5	15
mri\$age.83	10	2	12
mri\$age.84	5	6	11
mri\$age.85	3	6	9
mri\$age.86	5	5	10
mri\$age.87	4	4	8
mri\$age.88	2	1	3
mri\$age.89	2	3	5
mri\$age.90	0	4	4
mri\$age.91	2	0	2
mri\$age.92	0	1	1
mri\$age.94	0	1	1
mri\$age.95	0	1	1
mri\$age.99	0	1	1
mri\$age.ALL	369	366	735

	Point Estimate	Test Statistic	df	95% CI	p-value	Warnings
Chi-squared		26.244	30		0.66262	

By default, `tabulate` gives us the count in each combination of the strata, and gives us the overall  $\chi^2$  test statistic, degrees of freedom, and p-value. We can add in other arguments as well. Let's say that we are interested in an odds ratio or a risk ratio - `tabulate()` can supply these, if we set `dispRatios = TRUE`. However, we supply both the odds ratio and the risk ratio, even though in some cases only one is appropriate. Thus it is up to the user to know which they can use. To display these ratios for `male` versus `chf` (an indicator variable of whether the patient had been diagnosed with congestive heart failure prior to MRI), we type

```
> tabulate(mri$male, mri$chf, dispRatios = TRUE)
Call:
tabulate(mri$male, mri$chf, dispRatios = TRUE)
      mri$chf.0 mri$chf.1 mri$chf.ALL
mri$male.0    354      15      369
mri$male.1    340      26      366
mri$male.ALL  694      41      735
```

	Point Estimate	Test Statistic	df	95% CI	p-value	Warnings
Chi-squared		3.2214	1		0.072679	
Odds Ratio	1.8047			[0.9396, 3.4663]		
Risk Ratio	1.2944			[1.0136, 1.6531]		

Now we are given a point estimate and a 95% confidence interval for both the odds and risk ratio, in addition to the  $\chi^2$  estimate.

Last, similar to `tableStat()`, we can identify different statistics to display. The syntax is exactly the same, and the same statistics from Table 1 are valid in `tabulate()`.



## 4 Plotting

Plots and descriptive statistics together lead to a much better understanding of the data. In our package, we have implemented two types of plots: boxplots and scatterplots. In doing so, we have attempted to address some concerns with both types of plots.

### 4.1 Boxplots

Boxplots reduce the data down to four summary measures: minimum, maximum, median, and the interquartile range (25th to 75th quantile). They also show “outliers” in the data. We put this in quotes because the definition of “outlier” depends on the function you call, and the software you are running. Even in the base R version, `boxplot()`, the boxplots can be constructed so that no outliers are shown.

Much of the criticism of boxplots lies both in this determination of outliers and in the dramatic reduction of the data. In our version, `bplot()`, we have added jittered data onto the boxplots (jittering allows us to see the data, but randomly adds noise so that we can see individual points better), and we have added the mean and standard deviation onto the plot. We can also produce stratified boxplots. For our function, the variable on the y-axis comes first, followed by the variable on the x-axis. This convention, while at odds with base R, is an attempt to make our functions match up with the regression scheme of response followed by predictors.

For our first example, let’s create a boxplot showing the age distribution of the `diabetes` variable:

```
> ## Base R boxplot
> boxplot(mri$age ~ mri$diabetes)
> ## Our version of boxplot
> bplot(mri$age, mri$diabetes)
```

In comparing the two, see that our plot displays the mean line and  $\pm$  standard deviation box in blue and overlays the jittered data on top of the boxplots produced by the base R function. Now if we also wanted to stratify by race, we write

```
> bplot(mri$age, mri$diabetes, strata = mri$race)
```

This boxplot breaks down diabetes by race, again showing the mean and standard deviation bars. Note that like normal plots, we can define the axis labels and title by hand.

### 4.2 Scatterplots

Scatterplots are a very common way to view data. They plot all of the data in one window, which allows the user to get a good overall view. However, many times the data is clumped together in some spots, and it is hard to see all of the points. Thus, similar to our approach in `bplot()`, our `scatter()` function jitters the data slightly by default. To illustrate this difference, we consider the distribution of cerebral atrophy by age group. In the base R scatterplot,

```
> plot(mri$age, mri$atrophy)
```

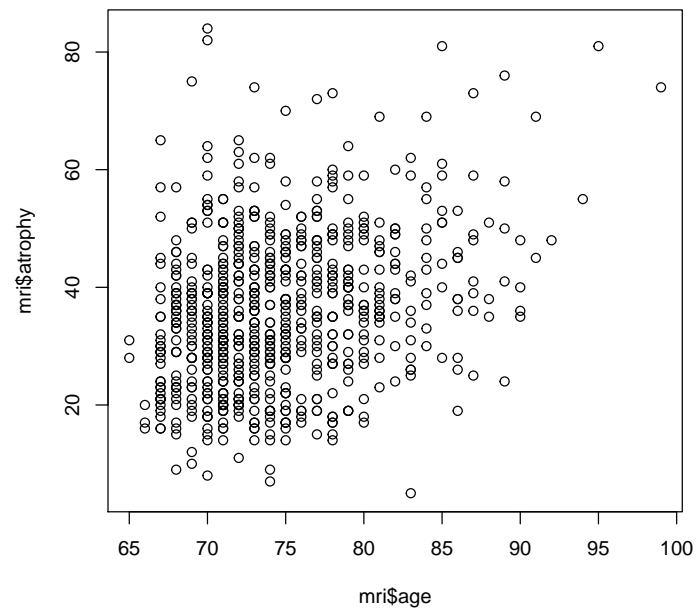


Figure 1: Base R Scatterplot

we see that a lot of the values sit on top of each other. In our version (which again uses the Y followed by X convention),

```
> scatter(mri$atrophy, mri$age)
```

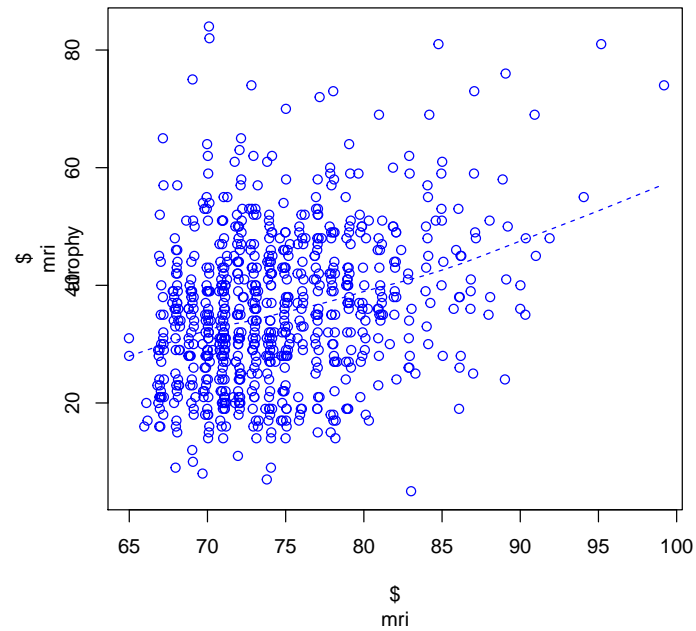


Figure 2: Scatterplot of atrophy vs age

the jittered data allows us to see where all of the points lie much better. Also notice that `scatter()` displays a loess curve by default. We can also display a line of best fit calculated by least squares by adding `plotLSfit = TRUE`. Last, we can stratify, which is particularly useful if we want to see differences across a third variable, and `scatter()` will automatically color the data appropriately:

```
> scatter(mri$atrophy, mri$age, strata = mri$race)
```

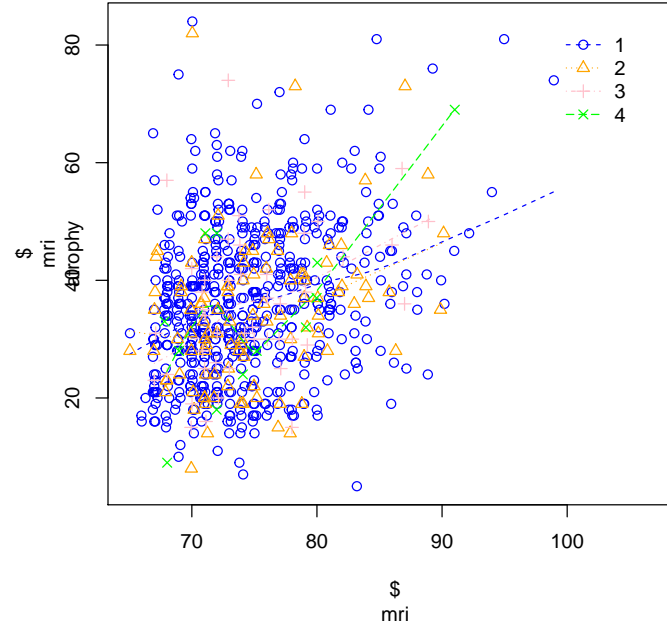


Figure 3: Scatterplot of atrophy vs age stratified by race

## 5 Transformations of a Variable

In many cases (especially in regression, which we will cover in section 7), it is more useful to model a transformation of a variable than the raw data itself. We present three functions to easily transform a variable. In each case, the function returns a list of: a matrix with the new transformed variable, the type of transformation performed, a vector describing how the variable was created, the name of the created variable, and the original data.

### 5.1 Dummy Variables

Dummy variables are a convenient way of creating indicator variables against some reference. For example, in the `mri` data, if we wanted to create a dummy variable for `race`, this would create 3 indicator variables (since there are four levels of race). By default, our `dummy()` function creates these indicators using the smallest value as the reference.

```

> dummy.race <- dummy(mri$race)
> head(dummy.race, n = 10)
      2 vs 1 3 vs 1 4 vs 1
[1,]      1      0      0
[2,]      1      0      0
[3,]      1      0      0
[4,]      0      0      0
[5,]      0      0      0
[6,]      0      0      1
[7,]      0      0      0
[8,]      1      0      0
[9,]      0      0      0
[10,]     0      0      0

```

Thus the set of dummy variables for race created above has: an indicator of whether the patient is black or white, an indicator of whether the patient is Asian or white, and an indicator of whether the patient is listed as “other” or white. These variables would allow us, in a regression, to compare each of the races against the white reference group.

## 5.2 Linear Splines

Linear splines create a piecewise linear fit between user specified “knot” points; these are the points in the variable that we allow the slope of the line to change. There are two major ways to parameterize linear splines, and we have implemented both in this package. The first is based on the absolute slope between knots - that is, if we had a variable (like `ldl` in the `mri` dataset) that took on values between 11 and 247 and we placed knots at 70 and 150, we would see three columns: the first would show the minimum of 70 and the value of `ldl`; the second would show the minimum of `ldl` - 70 and 80; and the last column would show the remaining part of `ldl`. For example, if `ldl` = 71 for one patient, we would see

```
70 1 0
```

. If `ldl` = 247, we would see

```
70 80 97
```

.

The second parameterization is denoted by us as “change”. This uses the change between knots to specify the slopes. Under this parameterization, we would see, for a value of 115,

```
115 45 0
```

, which corresponds to: 115 units between the minimum and the value, 45 units between knot 1 (at 70) and the value, and zero units between knot 2 (at 150) and the value. We do not show negative units.

These two parameterizations can be accessed via

```

> lspline.1dl <- lspline(mri$1dl, knots = c(70, 150))
> head(lspline.1dl, n = 10)
  $ mri 1dl
70 65 0
70 14 0
70 45 0
61 0 0
70 78 0
70 80 13
70 31 0
70 46 0
70 54 0
70 40 0

> lsplineD.1dl <- lsplineD(mri$1dl, knots = c(70, 150))
> head(lsplineD.1dl, n = 10)
x:min x: 70 x:150
135 65 0
84 14 0
115 45 0
61 0 0
148 78 0
163 93 13
101 31 0
116 46 0
124 54 0
110 40 0

```

### 5.3 Polynomials

Last we come to polynomials. This function creates a polynomial of the specified degree simply by multiplying the variable by itself the correct number of times. If we wanted to create a parabola in `age`, we type

```

> age.parabola <- polynomial(mri$age, degree = 2)
> head(age.parabola, n = 10)
      Linear(ctr) Square(ctr)
[1,] -2.5659864   6.5842862
[2,]  6.4340136  41.3965311
[3,] 15.4340136 238.2087760

```

```

[4,] -2.5659864  6.5842862
[5,] -4.5659864 20.8482318
[6,] -2.5659864  6.5842862
[7,]  0.4340136  0.1883678
[8,]  0.4340136  0.1883678
[9,] -7.5659864 57.2441501
[10,] -4.5659864 20.8482318

```

The “(ctr)” after each term tells us that `polynomial()` has automatically centered each term, and by default it centers at the mean of the variable.

## 6 One and Two Sample Functions

Many of our analyses boil down to one-sample or two-sample problems; What is the mean time to death? What is the median home price in Seattle? What is the difference in mean time to death between control and the treatment group? The list goes on. There are many methods of analyzing one-sample relationships, and in our package we have implemented three.

### 6.1 Correlation

While correlation is not an excellent source of inference, since it is inherently a feature of the dataset collected, sometimes it is important to know when variables are correlated. In the base R package, there are a few functions to calculate correlation: `cor()`, `var()`, `cov()`, and `cov2cor()`. Our function, `correlate()`, computes the correlation matrix between an arbitrary number of variables, and optionally also does this within a stratification variable. This is the real advantage of `correlate()`, since in base R we would have to manually subset the data into each stratum. If we want to examine the correlation between sex and diabetes, (and then stratify by race), we get

```

> ## Base R - returns only one number!
> cor(mri$male, mri$diabetes)
[1] 0.1200211
> ## uwIntroStats version
> correlate(mri$male, mri$diabetes)
Tabled correlation statistics by strata
Call:
  correlate(mri$male, mri$diabetes)
Method: Pearson
Data   : Pairwise Complete
      - NaN denotes strata with no observations
      - NA arises from missing data

##### ALL DATA
## Estimated Correlation Coefficients

```

```

      mri$male:  mri$diabetes:
mri$male:  1.0000      0.1200
mri$diabetes:  0.1200      1.0000
> ## Stratify on race
> correlate(mri$male, mri$diabetes, strata = mri$race)
Tabled correlation statistics by strata
Call:
  correlate(mri$male, mri$diabetes, strata = mri$race)
Method: Pearson
Data   : Pairwise Complete
      - NaN denotes strata with no observations
      - NA arises from missing data

##### Stratum   1
## Estimated Correlation Coefficients
      mri$male:  mri$diabetes:
mri$male:  1.0000      0.0941
mri$diabetes:  0.0941      1.0000

##### Stratum   2
## Estimated Correlation Coefficients
      mri$male:  mri$diabetes:
mri$male:  1.0000      0.2122
mri$diabetes:  0.2122      1.0000

##### Stratum   3
## Estimated Correlation Coefficients
      mri$male:  mri$diabetes:
mri$male:  1.0000      0.1155
mri$diabetes:  0.1155      1.0000

##### Stratum   4
## Estimated Correlation Coefficients
      mri$male:  mri$diabetes:
mri$male:  1.0000      0.3162
mri$diabetes:  0.3162      1.0000

##### ALL DATA
## Estimated Correlation Coefficients

```



```

          mri$male:  mri$diabetes:
mri$male:  1.0000    0.1200
mri$diabetes: 0.1200    1.0000

```

Our function takes the pain out of calculating multiple correlations within stratum variables. We also return a correlation matrix rather than a single number, which allows the user to go straight to a covariance matrix by squaring. We also allow calculation of Spearman or Pearson (default) correlation.

## 6.2 Point Estimates and Inference

As mentioned above, since correlation is a function of the data gathered, we usually need other methods for our inference. One of the most common tests is the t-test, since we are often interested in the mean and making comparisons between means. Perhaps less common are the Wilcoxon and Mann-Whitney, which use the “rank” of the variables compared.

### 6.2.1 T-tests

In the base R package, the `t.test()` function performs a t-test as you would expect. Our function `ttest()` improves on this by allowing stratification, calculation of the geometric mean, allowing for presuming unequal variances between samples, and making a cleaner output. For example, a t-test of whether the mean of the `ldl` variable is equal to 125 yields, in base R and `uwIntroStats`

```

> ## base R
> t.test(mri$ldl, mu = 125)
      One Sample t-test

data:  mri$ldl
t = 0.64326, df = 724, p-value = 0.5203
alternative hypothesis: true mean is not equal to 125
95 percent confidence interval:
 123.3527 128.2528
sample estimates:
mean of x
 125.8028
> ## uwIntroStats
> ttest(mri$ldl, null.hypoth = 125)
Call:
ttest(var1 = mri$ldl, null.hypoth = 125)

One-sample t-test :

Summary:
Variable Obs Missing Mean Std. Err. Std. Dev.      95% CI

```

```
mri$ldl 735      10  126      1.25      33.6 [123, 128]
```

```
Ho: mean = 125 ;  
Ha: mean != 125  
t = 0.6433 , df = 724  
Pr(|T| > t) = 0.520256
```

If instead we wanted a two-sample t-test of whether the difference in mean LDL between males and females were zero, we would get

```
> ## base R  
> t.test(mri$ldl[mri$male == 0], mri$ldl[mri$male == 1])  
Welch Two Sample t-test  
  
data: mri$ldl[mri$male == 0] and mri$ldl[mri$male == 1]  
t = 4.1938, df = 721.23, p-value = 3.084e-05  
alternative hypothesis: true difference in means is not equal to 0  
95 percent confidence interval:  
 5.502342 15.188221  
sample estimates:  
mean of x mean of y  
 130.9397  120.5944  
> ## uwIntroStats  
> ttest(mri$ldl, by = mri$male)  
Call:  
ttest(var1 = mri$ldl, by = mri$male)
```

Two-sample t-test allowing for unequal variances :

Summary:

	Group	Obs	Missing	Mean	Std. Err.	Std. Dev.	95% CI
mri\$male = 1	369	4	130.9	1.79	34.3	[127.4, 134.5]	
mri\$male = 0	366	6	120.6	1.69	32.1	[117.3, 123.9]	
Difference	735	10	10.3	2.47	<NA>	[5.5, 15.2]	

```
Ho: difference in means = 0 ;  
Ha: difference in means != 0  
t = 4.194 , df = 721  
Pr(|T| > t) = 3.08428e-05
```

Note that in our package, the default is to presume unequal variances between samples, which the authors believe to usually be the correct course. Also, we don't have to subset the data manually. We also run two-sided tests by default, but others can be specified.

### 6.2.2 Generic Inference

Until now, all of the methods in this section work on multiple samples. However, if we want to produce point estimates, interval estimates, and p-values for an arbitrary functional (mean, geometric mean, proportion, median, quantile, odds) of a variable, we can use the `oneSample()` function. Base R does not have a function which performs this general inference all in one spot. In the following tests, we will run one- and two-sample t-tests and an exact binomial test, all on the LDL variable. The t-tests run by `oneSample()` are an alternative to `ttest()` as described above. The output is not as verbose, but all of the information is the same:

```
> ## base R
> t.test(mri$ldl, mu = 125)

    One Sample t-test

data:  mri$ldl
t = 0.64326, df = 724, p-value = 0.5203
alternative hypothesis: true mean is not equal to 125
95 percent confidence interval:
 123.3527 128.2528
sample estimates:
mean of x
 125.8028
> ## ttest
> ttest(mri$ldl, null.hypoth = 125)
Call:
ttest(var1 = mri$ldl, null.hypoth = 125)

One-sample t-test :

Summary:
Variable Obs Missing Mean Std. Err. Std. Dev.    95% CI
mri$ldl 735      10 126      1.25      33.6 [123, 128]

Ho:  mean = 125 ;
Ha:  mean != 125
t = 0.6433 , df = 724
Pr(|T| > t) = 0.520256
> ## oneSample
> oneSample("mean", mri$ldl, null.hypothesis = 125)
Hypothesis test of two-sided alternative that Mean <> 125
Method: One sample t test
```

```

10 observations deleted due to missing values
  n      Mean   95% CIlo 95% CIhi Null Hyp   P two
725   125.8    123.4   128.3   125.0    0.5203

```

However, the true flexibility of the `oneSample()` function comes into play when we ask about different functionals. If we want inference on the geometric mean (which might make sense since LDL is a biological variable that may exhibit extreme values due to illness), in base R or with `ttest()` we have to logarithmically transform our variable and then run the t-test. With `oneSample()`, we simply run on "geometric mean":

```

> ## base R
> t.test(log(mri$ldl), mu = log(125))
      One Sample t-test

data:  log(mri$ldl)
t = -3.0054, df = 724, p-value = 0.002744
alternative hypothesis: true mean is not equal to 4.828314
95 percent confidence interval:
 4.774290 4.816983
sample estimates:
mean of x
 4.795636
> ## ttest
> ttest(log(mri$ldl), null.hypoth = log(125))
Call:
ttest(var1 = log(mri$ldl), null.hypoth = log(125))

One-sample t-test :

Summary:
  Variable Obs Missing Mean Std. Err. Std. Dev.      95% CI
log(mri$ldl) 735      10  4.8    0.0109    0.293 [4.77, 4.82]

Ho:  mean = 4.8283137373023 ;
Ha:  mean != 4.8283137373023
t = -3.005 , df = 724
Pr(|T| > t) =  0.00274401
> ## oneSample
> oneSample("geometric mean", mri$ldl, null.hypothesis = 125)
Hypothesis test of two-sided alternative that GeomMn <> 125
Method: One sample t test on log transformed data

```

```

10 observations deleted due to missing values
  n      GeomMn  95% CIlo  95% CIhi  Null Hyp  P two
725    121.0    118.4    123.6    125.0    2.744e-03

```

We can also compute an exact binomial test of the proportion of LDL values that are greater than 128:

```

> ## base R
> binom.test(sum(mri$ldl > 128, na.rm = TRUE), n = sum(!is.na(mri$ldl)))
      Exact binomial test

data:  sum(mri$ldl > 128, na.rm = TRUE) and sum(!is.na(mri$ldl))
number of successes = 336, number of trials = 725, p-value = 0.05338
alternative hypothesis: true probability of success is not equal to 0.5
95 percent confidence interval:
 0.4266728 0.5005223
sample estimates:
probability of success
 0.4634483

> ## oneSample
> oneSample("prop", mri$ldl > 128, null.hypothesis = 0.5)
Hypothesis test of two-sided alternative that Pr(Event) <> 0.5
Method: One sample inference for binomial proportions using exact distribution (LR ordering)
10 observations deleted due to missing values
  n      Pr(Event) 95% CIlo  95% CIhi  Null Hyp  P two
725    0.4634    0.4267    0.5005    0.5000  0.05338

> oneSample("prop", mri$ldl, above = 128, null.hypothesis = 0.5)
Hypothesis test of two-sided alternative that Pr>128 <> 0.5
Method: One sample inference for binomial proportions using exact distribution (LR ordering)
10 observations deleted due to missing values
  n      Pr>128  95% CIlo  95% CIhi  Null Hyp  P two
725    0.4634    0.4267    0.5005    0.5000  0.05338

```

The syntax for `oneSample()` is much more intuitive; we simply enter that we want to test a proportion, and then give the logical value to test or specify the threshold. Also, rather than having to count successes, the total, and dealing with missing values by hand, we can let `oneSample()` handle everything for us and give output that is clearer to read.

### 6.2.3 Wilcoxon and Mann-Whitney

The most similar function between our package and the base R package is the function to perform Wilcoxon Signed Rank tests and Mann-Whitney-Wilcoxon Ranked Sum tests. Our function, `wilcoxon()`, adds formatting and variances, and prints the z-score and p-value in addition to the test statistic

and p-value. Otherwise, the functions are the same, though we follow our usual syntax of Y followed by X. For example, if we want to test diseased versus healthy (in a made-up dataset), we can get

```
> ## create the data
> cf <- c(1153, 1132, 1165, 1460, 1162, 1493, 1358, 1453, 1185, 1824, 1793, 1930, 2075)
> healthy <- c(996, 1080, 1182, 1452, 1634, 1619, 1140, 1123, 1113, 1463, 1632, 1614, 1836)
> ## base R
> wilcox.test(healthy, cf, paired = TRUE)
    Wilcoxon signed rank test

data:  healthy and cf
V = 20, p-value = 0.08032
alternative hypothesis: true location shift is not equal to 0
> ## uwIntroStats
> wilcoxon(cf, healthy, paired = TRUE)
    Wilcoxon signed rank test
      obs sum ranks expected
positive  10      71    45.5
negative   3      20    45.5
zero       0       0     0.0
all       13      91    91.0

unadjusted variance  204.75
adjustment for ties    0.00
adjustment for zeroes  0.00
adjusted variance    204.75
              H0 Ha
Hypothesized Median 0 two.sided
  Test Statistic p-value
V 71              0.080322
Z 1.7821          0.037368
```

Note that in the output of `wilcoxon()`, we have displayed the data calculated by the signed rank test, and given both the test statistic (V) and the z-score (Z).

## 7 Regression

Regression is one of the most widely used and easily understood methods of statistical analysis. The base R package has many different functions to perform regression: `lm()` for linear regression, `glm()` for generalized linear regression (which allows you to perform logistic regression, for example),

`coxph()` for proportional hazards regression, and `geeglm()` as (one) method for correlated data regression. Each function has slightly different options, and remembering all of these is a chore. Part of the motivation for `regress()` is to alleviate this problem. This function provides one interface through which all types of regression can be performed. We also print the output from the regression in a much more understandable manner. Also, as we did in `ttest()`, we allow the user to calculate and use robust standard error estimates (using the Huber-White sandwich estimator), which adds flexibility. The next four subsections deal with aspects of `regress()` in more detail.

## 7.1 Basics of `regress()`

The minimum we need to enter into `regress` is a functional, a formula, and a dataset. A functional takes an object and returns a value; for instance, the mean is a functional because it takes a distribution and returns the mean. The allowed functionals are displayed in Table 2. Once we have a functional in mind (for now we choose the mean), we need to decide if we want to use robust standard error estimates. The default in `regress()` is to use these estimates, since we usually believe that the variances are not truly equal between groups compared in a regression analysis. The base R functions do presume equal variances between groups, which can lead to conservative (or anti-conservative) inference in some situations. The last change we have made to the usual regression output is displaying F-statistics rather than t-statistics for the test of each variable. We decided to use the F-distribution to make our results line up more easily with classical ANOVA tables. In most cases, the F-statistic is simply the square of the t-statistic. The inference is the same. To see `regress()` in action, consider testing the association between age and atrophy. The syntax for this computation is the same as that for any of the previous commands you might have used.

```
> ## base R
> summary(lm(atrophy ~ age, data = mri))
Call:
lm(formula = atrophy ~ age, data = mri)

Residuals:
    Min       1Q   Median       3Q      Max
-36.870  -8.589  -0.870   7.666  51.203

Coefficients:
              Estimate Std. Error t value Pr(>|t|)
(Intercept) -16.06213     6.25619  -2.567   0.0104 *
age           0.69798     0.08368   8.341 3.64e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 12.36 on 733 degrees of freedom
Multiple R-squared:  0.08669,    Adjusted R-squared:  0.08545
F-statistic: 69.58 on 1 and 733 DF,  p-value: 3.635e-16
> ## uwIntroStats
> regress("mean", atrophy ~ age, data = mri)
```

```

Call:
regress(fnctl = "mean", formula = atrophy ~ age, data = mri)

Residuals:
    Min       1Q   Median       3Q      Max
-36.870  -8.589  -0.870   7.666  51.203

Coefficients:
              Estimate Naive SE Robust SE    95%L    95%H      F stat    df Pr(>F)
[1] Intercept      -16.06    6.256    6.701   -29.22   -2.907        5.75  1    0.0168
[2] age              0.6980  0.08368  0.09002    0.5213    0.8747       60.12  1 < 0.00005

Residual standard error: 12.36 on 733 degrees of freedom
Multiple R-squared:  0.08669,    Adjusted R-squared:  0.08545
F-statistic: 60.12 on 1 and 733 DF,  p-value: 2.988e-14

```

Note that by default, our output is printed in a table, rather than having to call `summary.lm()`. The inference on the individual coefficients is different because we have used the F-distribution and are using robust standard error estimates, but we get approximately the same p-values. Also, the overall F-statistic is different in our version because we are using the robust standard error estimates. The numbers next to the coefficients tell us which to specify in any post-testing commands we run (see section 8).

## 7.2 Regression on different functionals

As mentioned above, part of the motivation for the `regress()` function was to have all types of regression in one function. Thus the other functionals we allow, and their corresponding commands in base R, are displayed in Table 2. Note that we only display the options which lead to a certain type of regression, not the rest of the syntax.

Functional	Type of Regression	Previous command (package)
"mean"	Linear Regression	<code>lm()</code> (stats - base R)
"geometric mean"	Linear Regression on logarithmically transformed Y	<code>lm()</code> ,with Y log transformed (stats - base R)
"odds"	Logistic Regression	<code>glm(family = binomial)</code> (stats - base R)
"rate"	Poisson Regression	<code>glm(family = poisson)</code> (stats - base R)
"hazard"	Proportional Hazards Regression	<code>coxph()</code> (survival)

Table 2: Allowed functionals for `regress()`.



7.3 Correlated data regression

7.4 Multiple-partial F-tests

## 8 Analysis/Post-testing

8.1 lincom

8.2 predict

## 9 Diagnostics