

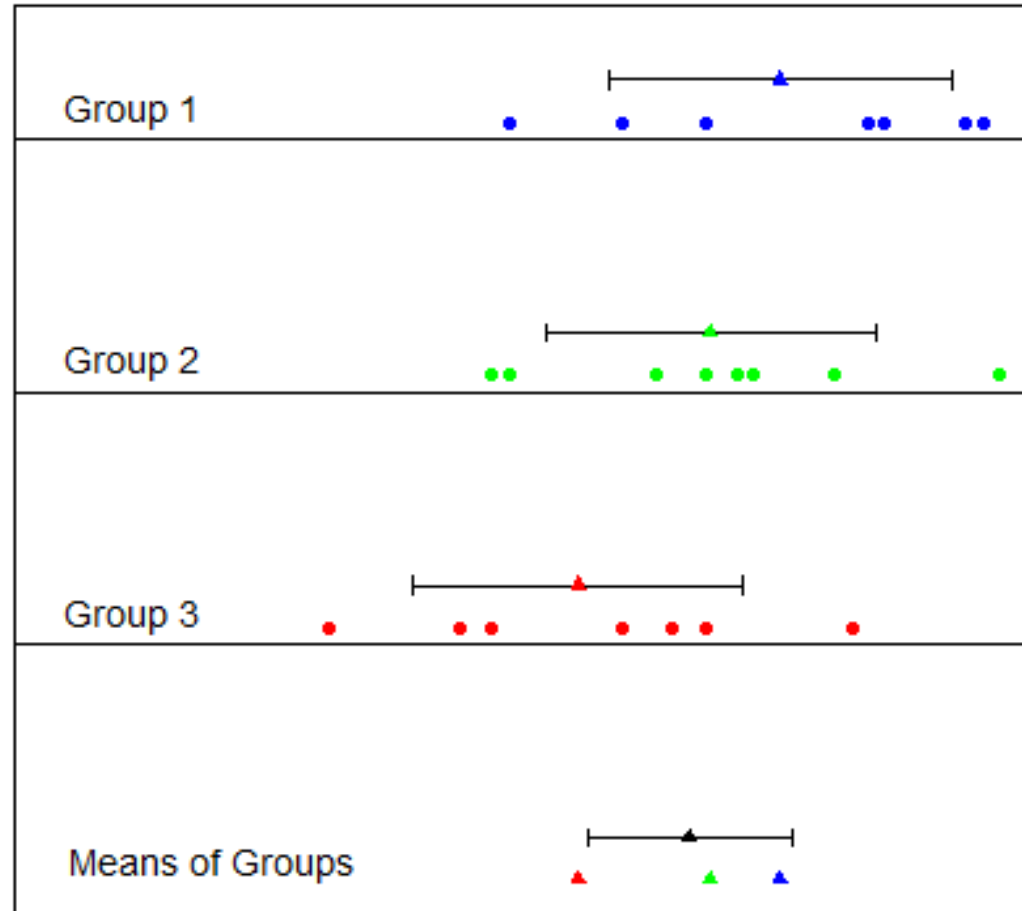
ANOVA – Categorical Data as Predictors

One-Way ANOVA

- ANOVA is used to test for differences among several means without increasing the Type I error rate, only doing one test
- The ANOVA uses data from all groups to estimate standard errors, which can increase the power of the analysis
- Basic Idea
 - Calculate the mean of the observations within each group
 - Compare the variance of these means to the average variance within each group
 - As the means become more different, the variance among the means increases

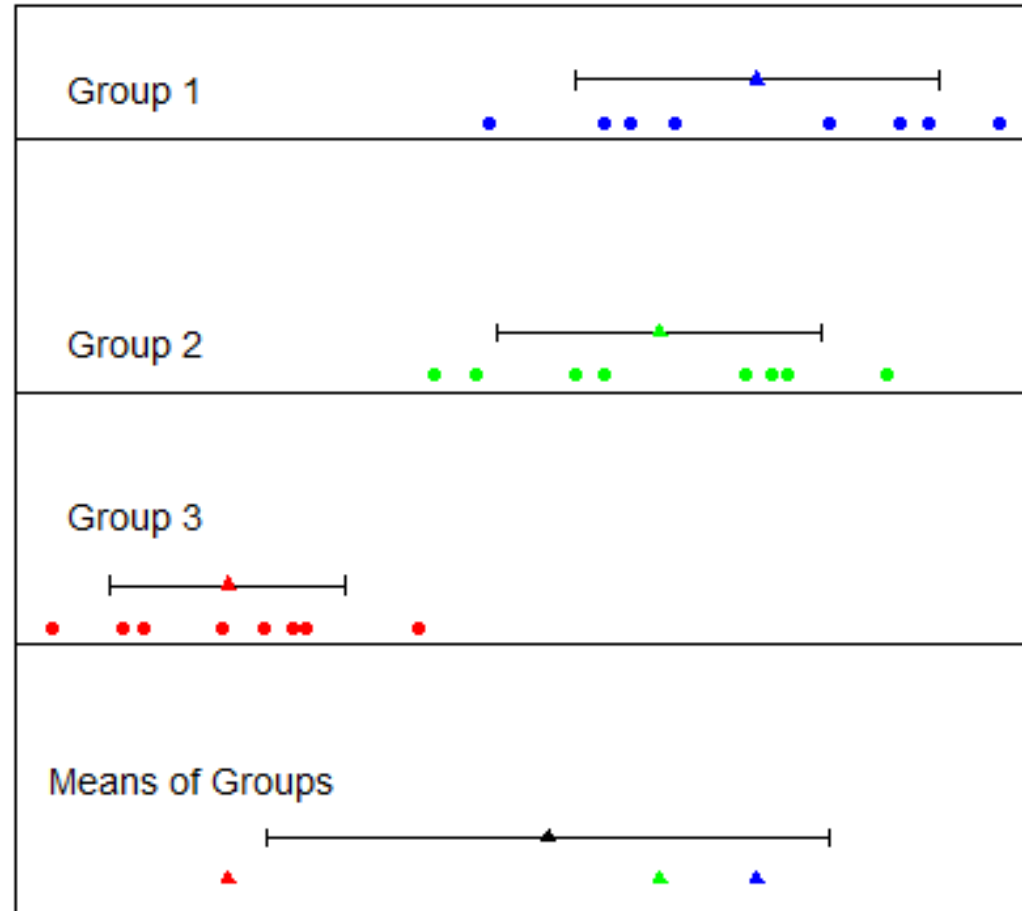
Why Look at Variance When Interested in Means?

- The variability within each sample is approximately the same
- The variability in the mean values of the samples is consistent with the variability within the individual samples



Why Look at Variance When Interested in Means?

- The variability in the sample means is much larger than would be expected given the variability within each of the samples



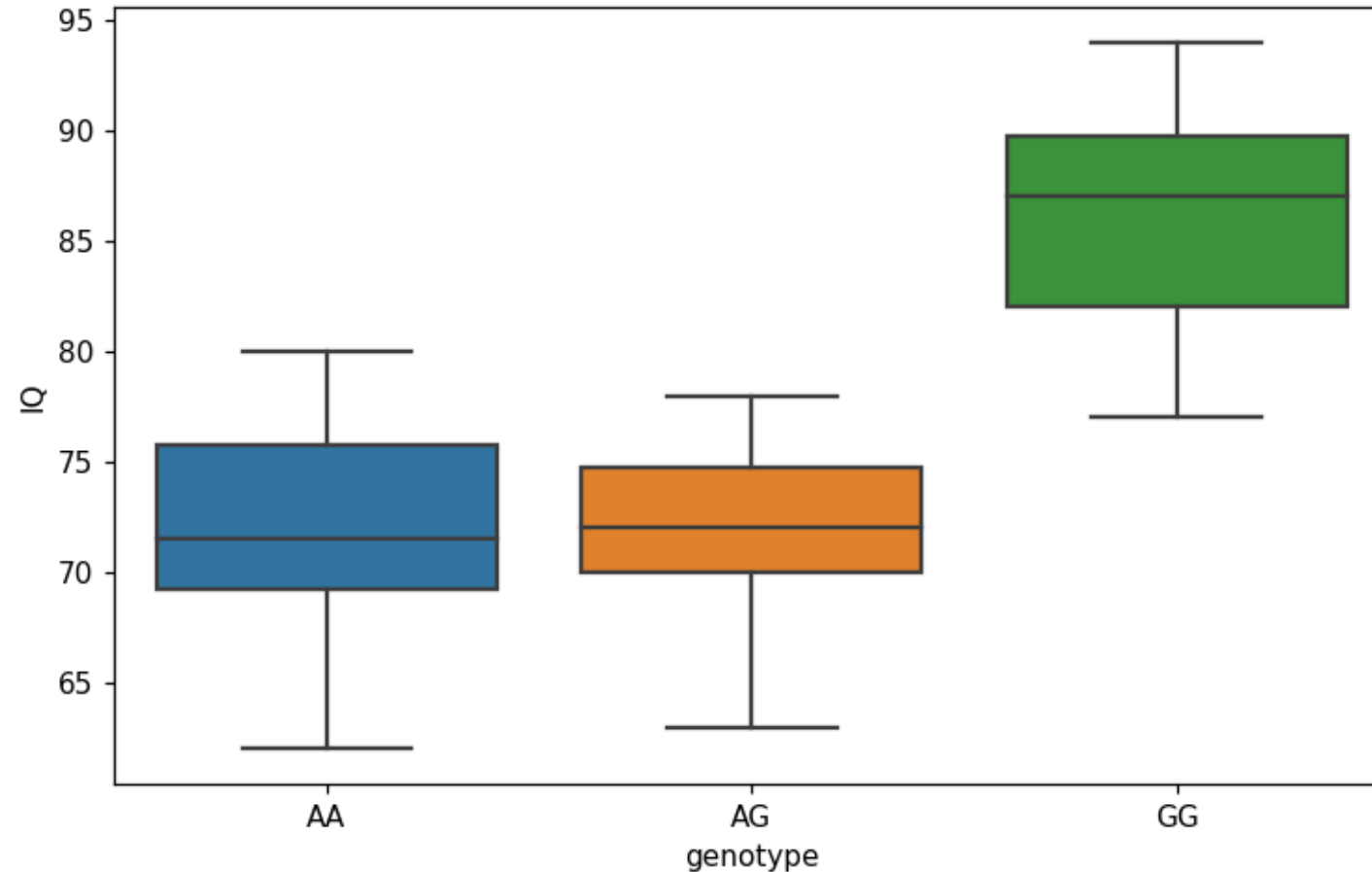
One-Way ANOVA

- When there is only **one categorical variable** which denotes the groups and only **one measurement variable** (quantitative), a one-way ANOVA is carried out
- For a one-way ANOVA the observations are divided into I mutually exclusive categories, giving the one-way classification

One-Way ANOVA Assumptions

- Assumptions
 - Each of the populations is **Normally distributed** with the **same variance** (homogeneity of variance)
 - The observations are **sampled independently**, the groups under consideration are independent
- ANOVA is robust to moderate violations of its assumptions, meaning that the probability values (P-values) computed in an ANOVA are sufficiently accurate even if the assumptions are moderately violated

One-Way ANOVA Example



- 54 observations
- 18 AA observations
mean IQ for AA = 71.9
- 18 AG observations
mean IQ for AG = 72.2
- 18 GG observations
mean IQ for GG = 86.1

Introduction of Notation

- Consider I groups, whose means we want to compare
- Let $n_i, i = 1, 2, \dots, I$ be the sample size of group i
- For the simulated verbal IQ and genotype data, ($I = 3$), representing the three possible genotypes at the particular location in a gene of interest. Each person in this data set, as well as having a genotype, also has a verbal IQ score

One-Way ANOVA Example

- Want to examine if the mean verbal IQ score is the same across the 3 genotype groups

Null hypothesis is that the mean verbal IQ is the same in the three genotype groups:

$$H_0 : \mu_1 = \mu_2 = \mu_3$$

One-Way ANOVA Example

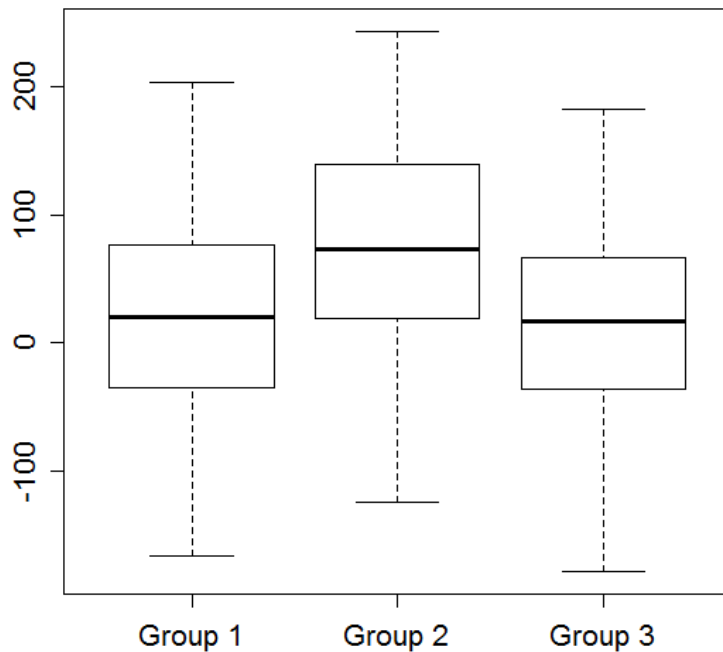
Within-Groups Variance

- Remember assumption that the population variances of the three groups is the same
- Under this assumption, the three variances of the three groups all estimate this common value, σ^2

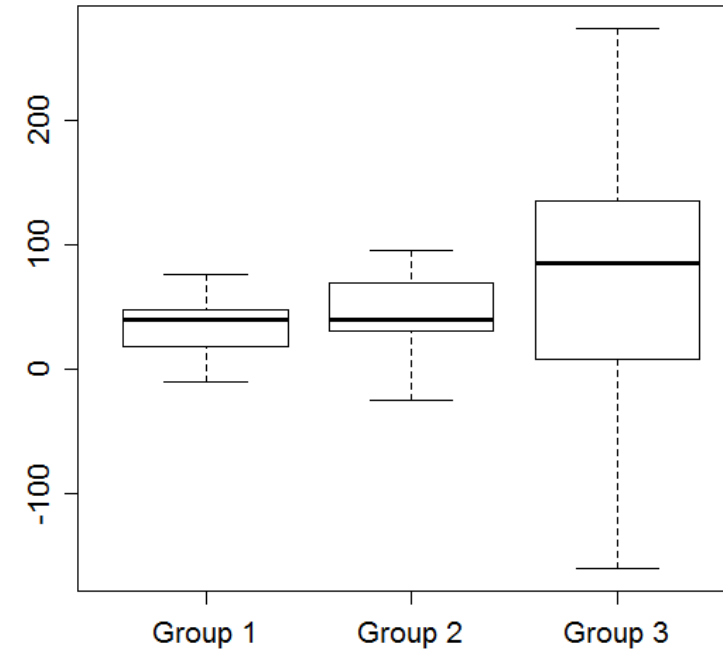
One-Way ANOVA Example

Within-Groups Variance

- Since the population variances are assumed to be equal the estimate of the population variance, derived from the separate within-group estimates, is valid whether or not the null hypothesis is true



EQUAL



NOT EQUAL

Within-Groups Variance

- For groups with equal sample size this is defined as the average of the variances of the groups

$$s_w^2 = \frac{1}{I} \sum_{i=1}^I s_i^2 = \frac{1}{I} \sum_{i=1}^I \sum_{j=1}^{n_i} \left(\frac{(x_{ij} - \bar{x}_i)^2}{n_i - 1} \right)$$

x_{ij} = observation j in group i

$\bar{x}_1, \bar{x}_2, \bar{x}_3$ = sample means of the genotype groups AA, AG, GG

One-Way ANOVA Example

Between-Groups Variance

If the null hypothesis is true (which we assume):

- the groups can be considered as random samples from the same population
- assumed equal variances, and because the null hypothesis is true, then the population means are equal
- The means are observations from the same sampling distribution of the mean

Between-Groups Variance

- The sampling distribution of the mean has variance

$$\sigma^2/n$$

- This gives a second method of obtaining an estimate of the population variance (n = number of observations in each group)
- The observed variance of the treatment means is an estimate of σ^2/n and is given by:

$$\frac{s^2}{n} = \sum_{i=1}^I \frac{(\bar{x}_i - \bar{x})^2}{I - 1}$$

Unequal Sample Sizes

- There are adjustments to these formulae when the sample sizes are not all equal in the groups:
 - Within Groups variance:

$$s_w^2 = \sum_{i=1}^I \frac{(n_i - 1)s_i^2}{N - I}$$

- Between Groups Variance:

$$s_b^2 = \sum_{i=1}^I n_i \frac{(\bar{x}_i - \bar{x})^2}{I - 1}$$

One-Way ANOVA Example

- If the null hypothesis IS TRUE then
 - the between-groups variance S_b^2
 - and the within-groups variance S_w^2
 - are both estimates of the population variance σ^2
- If the null hypothesis is NOT TRUE then
 - the population means are not all equal
 - then S_b^2 will be greater than the population variance, σ^2
 - it will be increased by the *treatment* (genotype) differences

One-Way ANOVA Example

- To test the null hypothesis we compare the ratio of s_b^2 and s_w^2 using an F-test

- F statistic is given by:
$$F = \frac{s_b^2}{s_w^2}$$

with $I - 1$ and $I(n-1) = N - I$ degrees of freedom

- Can also think of the F statistic as:

$$F = \frac{\text{variability due to treatment effect and variability due to chance}}{\text{variability due to chance}}$$

F Distribution and F-test

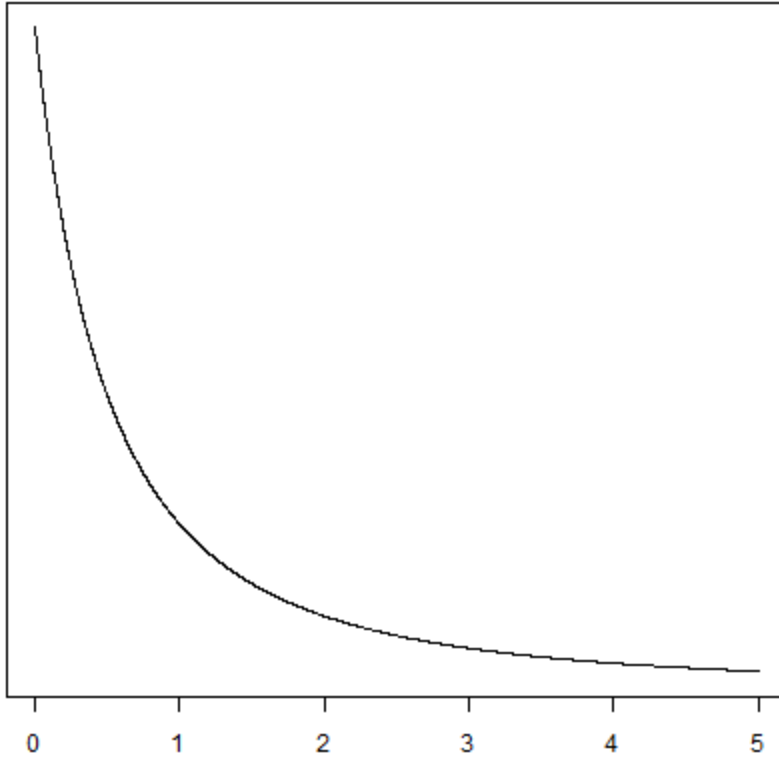
- The F distribution is the continuous distribution of the ratio of two estimates of variance
- The F distribution has two parameters: degrees of freedom numerator (top) and degrees of freedom denominator (bottom)
- The F-test is used to test the hypothesis that two variances are equal

F Distribution and F-test

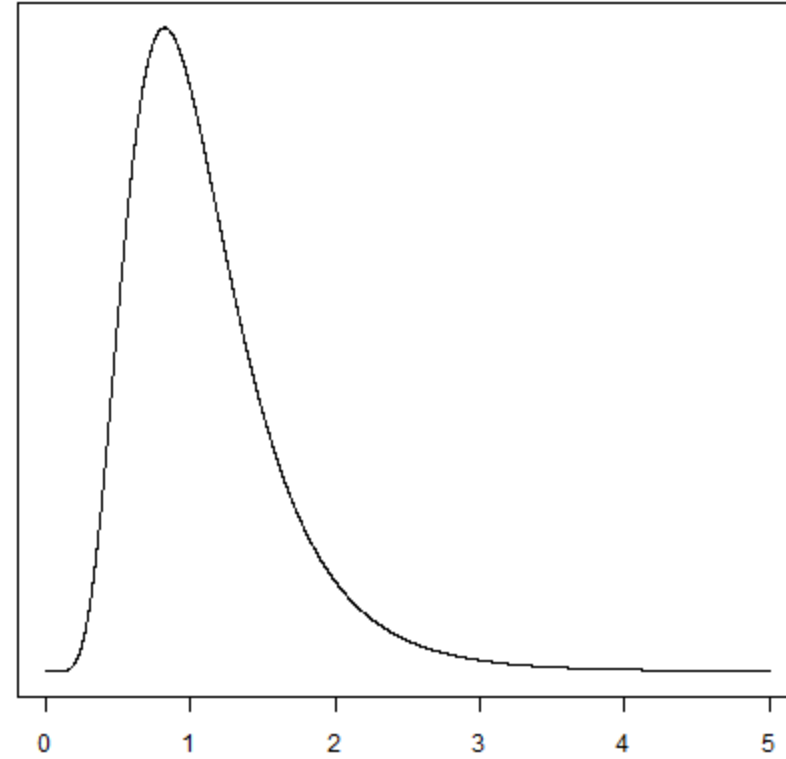
- The validity of the F-test is based on the requirement that the populations from which the variances were taken are Normal
- In the ANOVA, a one-sided F-test is used

F Distribution

F Distribution with $df1 = df2 = 2$



F Distribution with $df1 = df2 = 20$



One-Way ANOVA Example

```
df.head()
```

```
Out[5]:
```

```
  genotype  IQ
0      AA  63.0
1      AA  67.0
2      AA  75.0
3      AA  76.0
4      AA  70.0
```

```
#create lm model output and then use anova on it, create using ols formula
```

```
from statsmodels.formula.api import ols
```

```
model= ols('IQ ~ C(genotype)', data=df).fit()
```

```
#or get dummy variables to get same results
```

```
df[["geno_AG","geno_GG"]]=pd.get_dummies(df["genotype"])[["AG","GG"]]
```

```
X = df[["geno_AG","geno_GG"]]
```

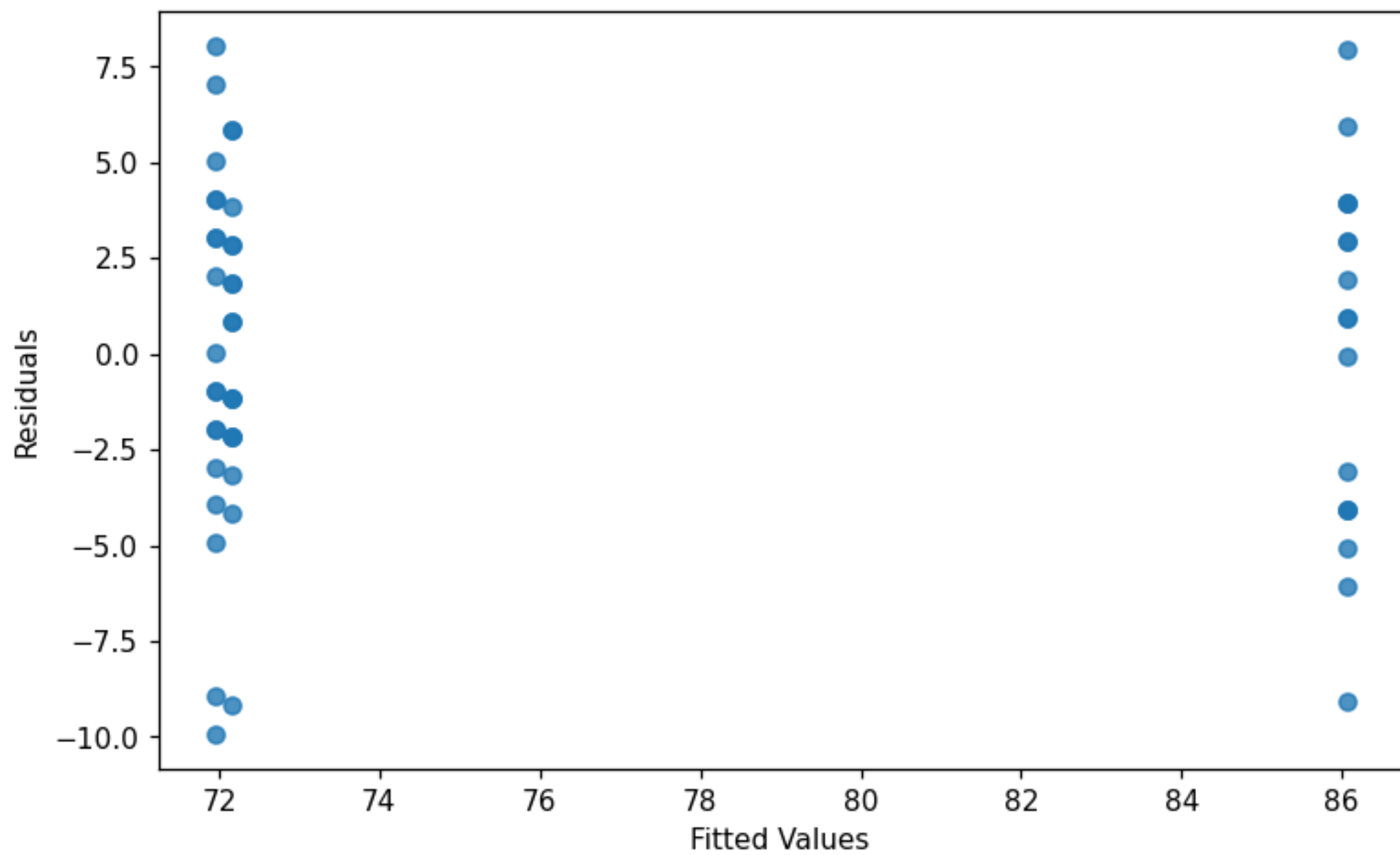
```
X = sm.add_constant(X)
```

```
y = df['IQ']
```

```
model1 = sm.OLS(y, X).fit()
```

One-Way ANOVA Assumption Checking

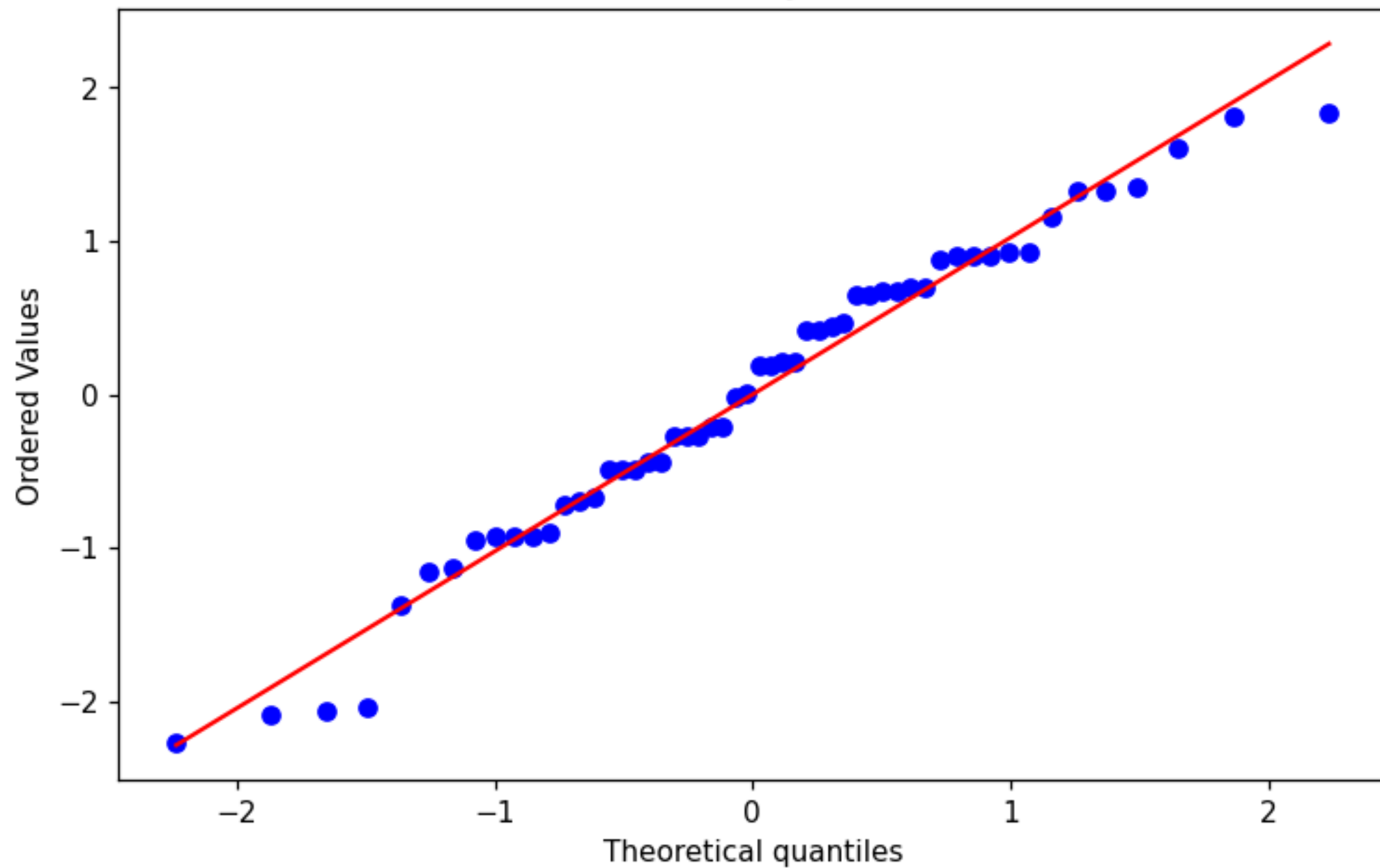
- **Homogeneity of variance = homoscedasticity**
 - The dependent variable (quantitative measurement) should have the same variance in each category of the independent variable (qualitative variable)
- ANOVA is robust for small to moderate departures from homogeneity of variance, especially with equal sample sizes for the groups
- Rule of thumb: the ratio of the **largest to the smallest group variance should be 3:1 or less**, but be careful, the more unequal the sample sizes the smaller the differences in variances which are acceptable
- Testing for homogeneity of variance
 - Examine **boxplots** of the data by group, will highlight visually if there is a large difference in variability between the groups
 - Plot **residuals versus fitted values** and examine scatter around zero,
residuals = observations – group mean
group mean = fitted value



One-Way ANOVA Assumption Checking

- **Normality Assumption**
 - The dependent variable (measurement, quantitative variable) should be Normally distributed in each category of the independent variable (qualitative variable)
- Again ANOVA is robust to moderate departures from Normality
- Checking the Normality assumption
 - **Boxplots of the data**
 - **Quantile-Quantile plots (QQ plots) of the residuals**, which should give a 45-degree line on a plot of observed versus expected values,

Probability Plot



One-Way ANOVA Example

```
from statsmodels.stats.anova import anova_lm
```

```
anovaResults = anova_lm(model, typ=1)
```

```
anovaResults
```

```
Out[18]:
```

	df	sum_sq	mean_sq	F	PR(>F)
C(genotype)	2.0	2352.444444	1176.222222	57.658568	8.112339e-14
Residual	51.0	1040.388889	20.399782	NaN	NaN

Between-Groups
Between treatments
= Treatment SS/df

P-Value

F Statistic

$I-1 = 3-1 = 2$, since
3 genotype groups,
AA, AG, GG

$I(n-1) = 3(18-1)$
= 51
 $N-I = 54 - 3 = 51$

Within-Groups
Residual Variation =
Residual SS/df

OLS Regression Results

```

=====
Dep. Variable:          IQ    R-squared:          0.693
Model:                  OLS   Adj. R-squared:      0.681
Method:                 Least Squares   F-statistic:      57.66
Date:                   Wed, 01 Jan 2020   Prob (F-statistic): 8.11e-14
Time:                   00:00:00   Log-Likelihood:     -156.50
No. Observations:       54   AIC:                319.0
Df Residuals:           51   BIC:                325.0
Df Model:                2
Covariance Type:        nonrobust
=====
  
```

$SS_{\text{between_treatment}}/SS_{\text{Total}}$

Mean of
baseline/control group

Estimate for differences
between the mean of each
group and the control group

T-test for
differences
between these
means

```

=====
               coef      std err      t    P>|t|    [0.025    0.975]
-----
Intercept      71.9444      1.065    67.580    0.000    69.807    74.082
C(genotype)[T.AG]  0.2222      1.506     0.148    0.883    -2.800     3.245
C(genotype)[T.GG] 14.1111      1.506     9.373    0.000    11.089    17.134
=====
  
```

```

=====
Omnibus:          1.387    Durbin-Watson:          1.943
Prob(Omnibus):    0.500    Jarque-Bera (JB):        1.402
Skew:             -0.325    Prob(JB):                0.496
Kurtosis:         2.553    Cond. No.                 3.73
=====
  
```

What to do with a Significant ANOVA Result (F-test)

- If the ANOVA is significant and the null hypothesis is rejected, the only valid inference that can be made is that at least one population mean is different from at least one other population mean
- The ANOVA does not reveal which population means differ from which others

What to do with a Significant ANOVA Result (F-test)

- Only think about investigating differences between individual groups when the overall comparison of groups (ANOVA) is significant, or that you had intended particular comparisons at the outset
- Need to consider whether the groups are ordered or not

Two Way ANOVA

- Two way analysis of variance (ANOVA) without interactions is the same as a regression with two categorical explanatory variables.
- Two way analysis of variance (ANOVA) with interactions is the same as a regression with two categorical explanatory variables plus a third categorical explanatory variable for the interaction.
- When the explanatory variable is categorical, conceptually it is recoded using dummy or indicator variables.

Two Way ANOVA

- We can test more hypothesis in a two-way ANOVA:
- There is no difference in the means of factor A
- There is no difference in means of factor B
- There is no interaction between factors A and B
- The alternative hypothesis for the first two is: the means are not equal.
- The alternative hypothesis for the last one is: there is an interaction between A and B.

Assumptions of a Two-Way ANOVA

- The dependent variable should be continuous and the two independent variables should be in categorical, independent groups.
- Observations are sampled independently – that each sample has been drawn independently of the other samples
- Equality of Variance (homogeneity of variance) – That the variance of data in the different groups should be the same
- Normality – That each sample is taken from a normally distributed population

Example: ToothGrowth

- Tooth Growth dataset in R contains data from a study evaluating the effect of vitamin C on tooth growth in Guinea pigs.
- The experiment has been performed on 60 pigs, where each animal received one of three dose levels of vitamin C (0.5, 1, and 2 mg/day) by one of two delivery methods - orange juice or ascorbic acid (a form of vitamin C and coded as VC).

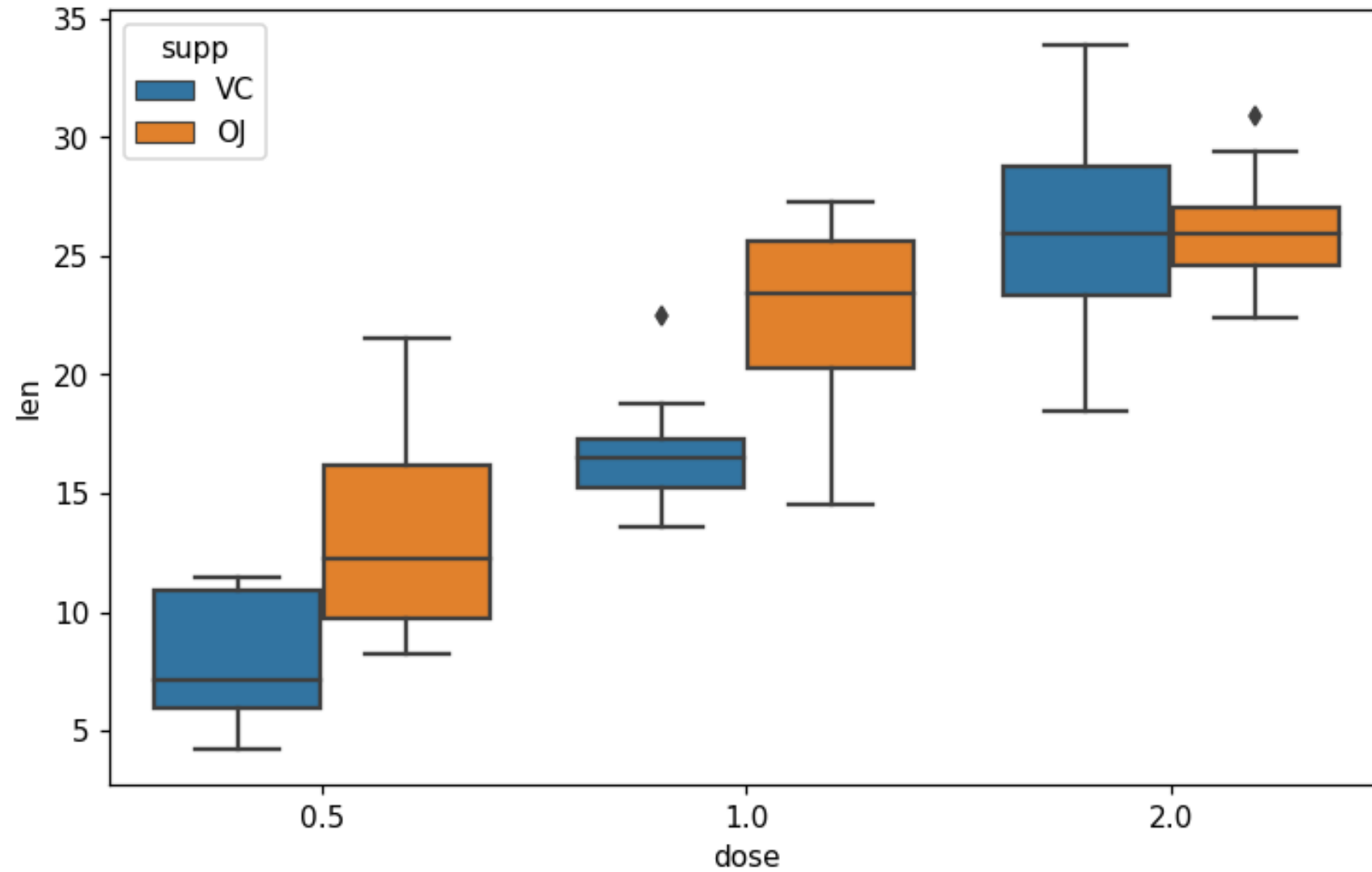
```
ToothGrowth.info()
<class 'pandas.core.frame.DataFrame'>
RangeIndex: 60 entries, 0 to 59
Data columns (total 3 columns):
#   Column  Non-Null Count  Dtype
---  -
0   len     60 non-null         float64
1   supp    60 non-null         object
2   dose    60 non-null         float64
dtypes: float64(2), object(1)
memory usage: 1.5+ KB
```

```
ToothGrowth.head()
```

```
Out[57]:
```

	len	supp	dose
0	4.2	VC	0.5
1	11.5	VC	0.5
2	7.3	VC	0.5
3	5.8	VC	0.5
4	6.4	VC	0.5

```
sns.boxplot(x="dose", y="len", hue='supp', data=ToothGrowth)
```



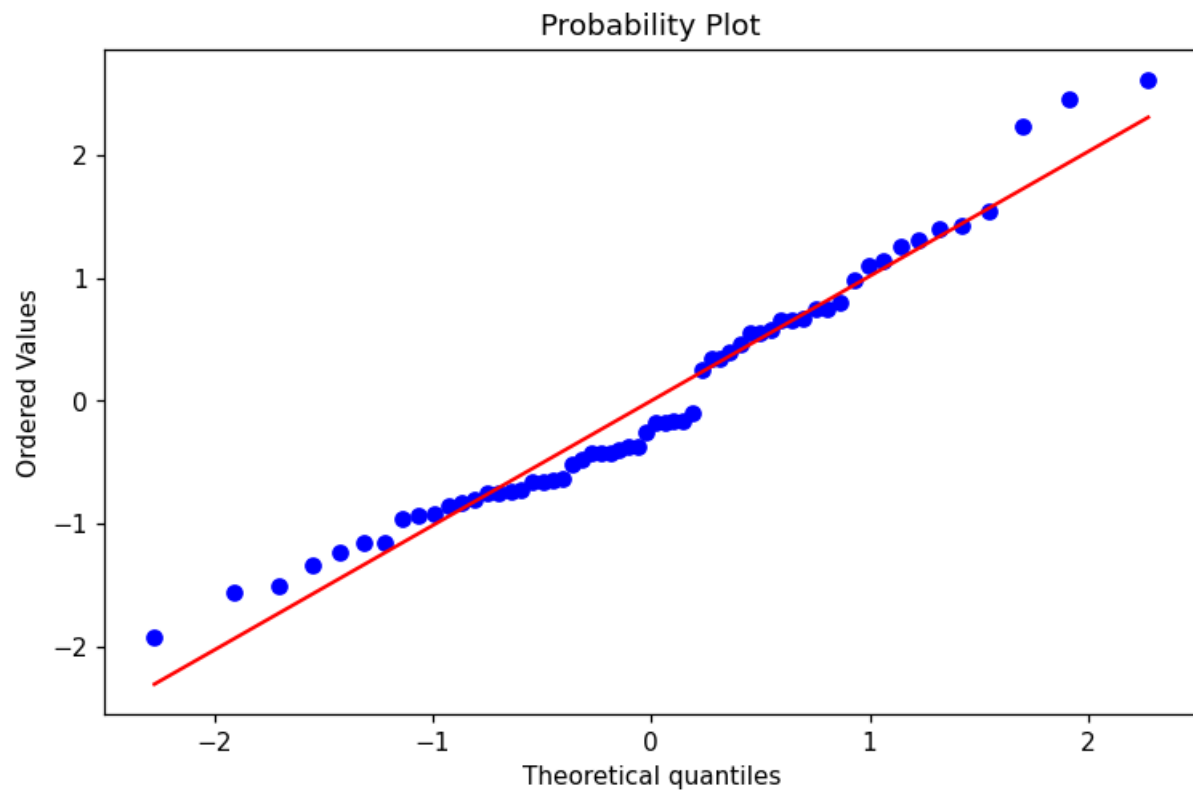
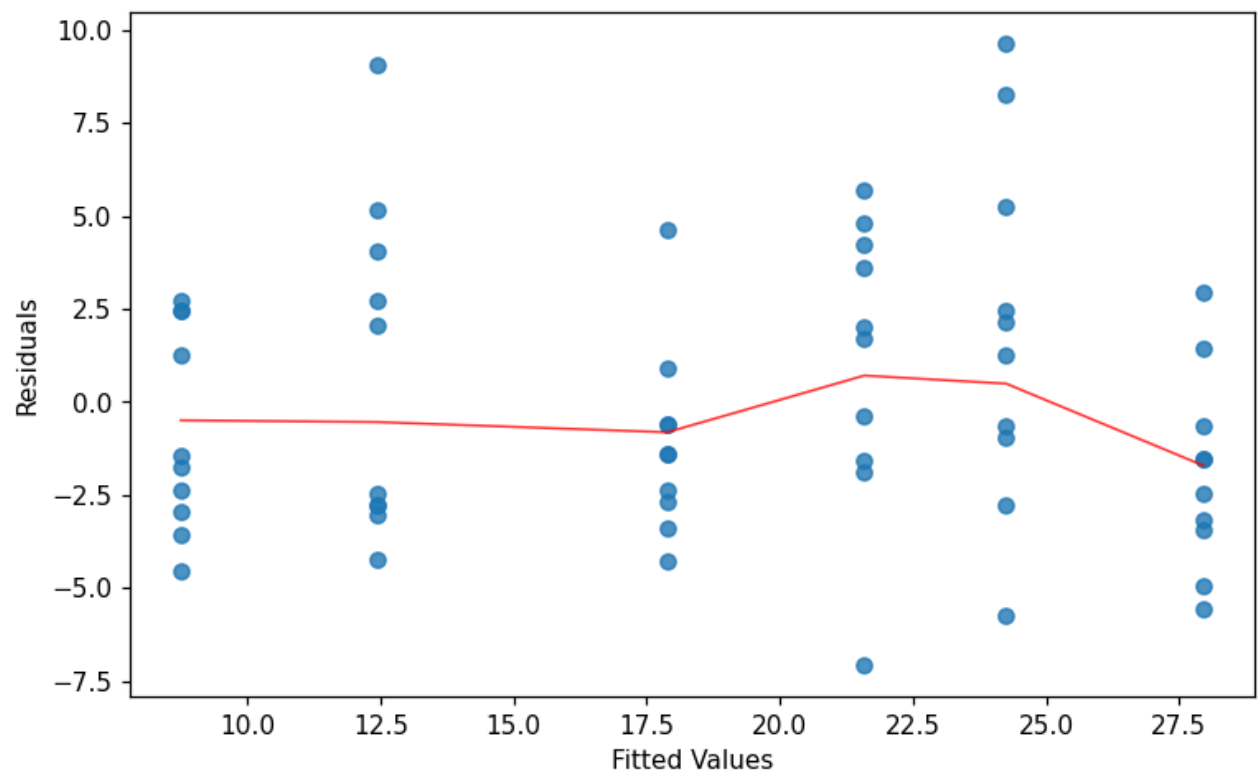
Example

```
model2= ols('len ~ C(supp)+C(dose)', data=ToothGrowth).fit()

#fitted values
model_fitted_vals = model2.fittedvalues
#model residuals
model_residuals = model2.resid
#standardised residuals
model_norm_residuals = model2.get_influence().resid_studentized_internal

sns.regplot(x=model_fitted_vals,y=model_residuals,
            ci=False,lowess=True,
            line_kws={'color': 'red', 'lw': 1, 'alpha': 0.8})
plt.xlabel("Fitted Values")
plt.ylabel("Residuals")

stats.probplot(model_norm_residuals, plot=sns.mpl.pyplot)
plt.show()
```



Example

```
supp_dose= pd.crosstab(index=ToothGrowth['supp'], columns=ToothGrowth["dose"], margins=True)
```

```
supp_dose
```

```
Out[60]:
```

```
dose 0.5 1.0 2.0 All
```

```
supp
```

```
OJ   10  10  10  30
```

```
VC   10  10  10  30
```

```
All  20  20  20  60
```

```
#balanced design
```

```
anova2way = anova_lm(model2, typ=1)
```

```
anova2way
```

```
Out[61]:
```

	df	sum_sq	mean_sq	F	PR(>F)
C(supp)	1.0	205.350000	205.350000	14.016638	4.292793e-04
C(dose)	2.0	2426.434333	1213.217167	82.810935	1.871163e-17
Residual	56.0	820.425000	14.650446	NaN	NaN

OLS Regression Results

```

=====
Dep. Variable:          len    R-squared:          0.762
Model:                  OLS    Adj. R-squared:      0.750
Method:                 Least Squares    F-statistic:      59.88
Date:                   Wed, 01 Jan 2020    Prob (F-statistic):  1.78e-17
Time:                   00:00:00    Log-Likelihood:     -163.60
No. Observations:       60    AIC:              335.2
Df Residuals:           56    BIC:              343.6
Df Model:                3
Covariance Type:        nonrobust
=====

```

```

=====
               coef      std err      t      P>|t|      [0.025      0.975]
-----
Intercept      12.4550      0.988     12.603     0.000     10.475     14.435
C(supp)[T.VC]   -3.7000      0.988     -3.744     0.000     -5.680     -1.720
C(dose)[T.1.0]   9.1300      1.210      7.543     0.000      6.705     11.555
C(dose)[T.2.0]  15.4950      1.210     12.802     0.000     13.070     17.920
=====

```

```

=====
Omnibus:          3.615    Durbin-Watson:          1.814
Prob(Omnibus):    0.164    Jarque-Bera (JB):        3.366
Skew:             0.575    Prob(JB):                0.186
Kurtosis:         2.853    Cond. No.                 4.22
=====

```

Interpretation

- From the ANOVA table (or coefficient table), we can conclude that both *supp* and *dose* are statistically significant.
- Therefore, this would imply that changing delivery methods (*supp*) or the dose of vitamin C, will impact significantly the mean tooth length.
- Not the above model is called **additive model**. It makes an assumption that the two factor variables are independent.
- If you think that these two variables have interaction effect then replace the plus symbol (+) by an asterisk (*)

Example with interaction term

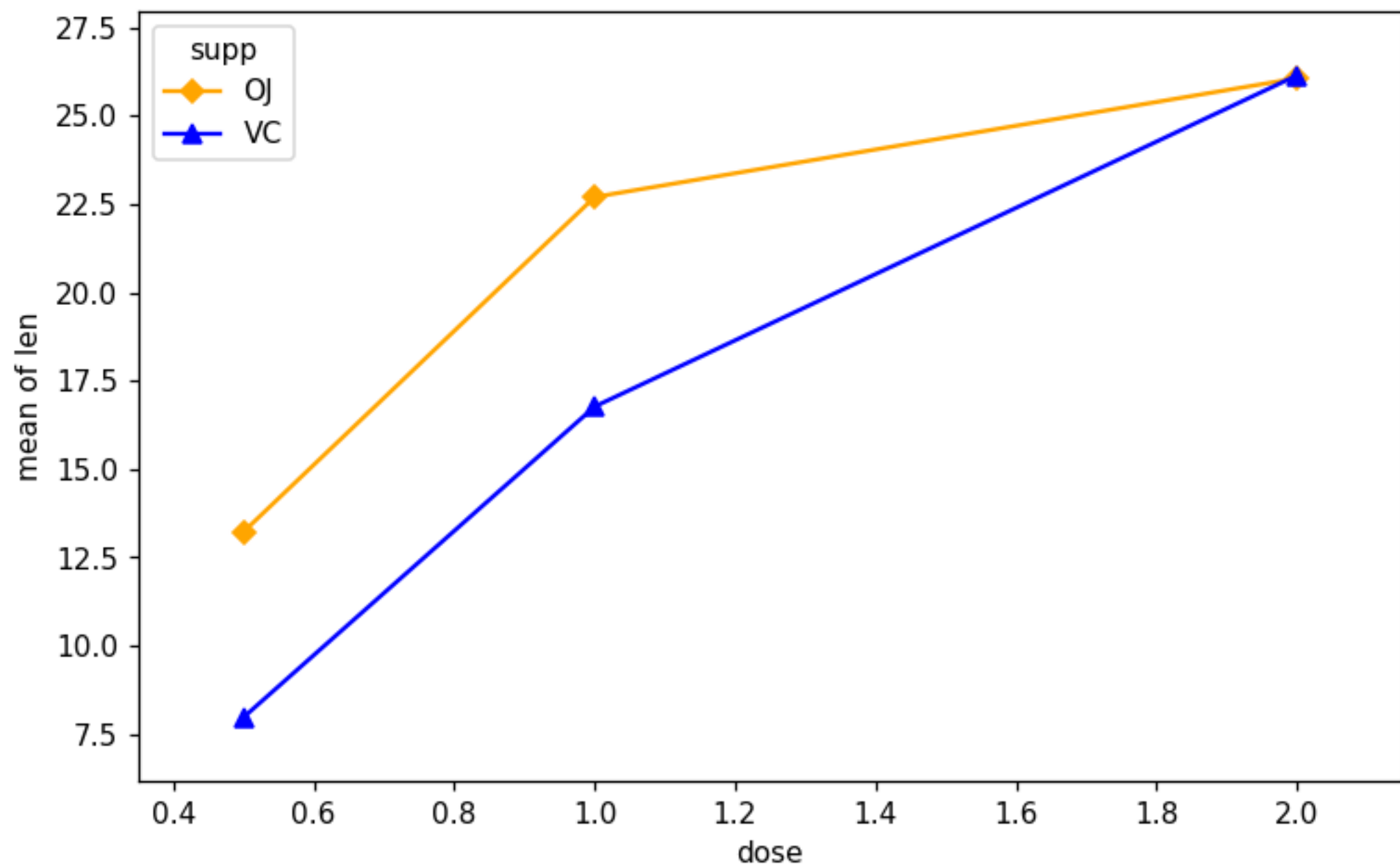
```
model2int= ols('len ~ C(supp)*C(dose)', data=ToothGrowth).fit()  
model2int= ols('len ~ C(supp)+C(dose)+C(supp):C(dose)', data=ToothGrowth).fit()  
# These two calls are equivalent
```

Interaction Plot in python

- `Interaction_plot` command in `statsmodels` package
- Parameters inside that package:
 - **x** : the factor to be plotted on x axis.
 - **trace**: the factor to be plotted as lines
 - **response**: a numeric variable giving the response

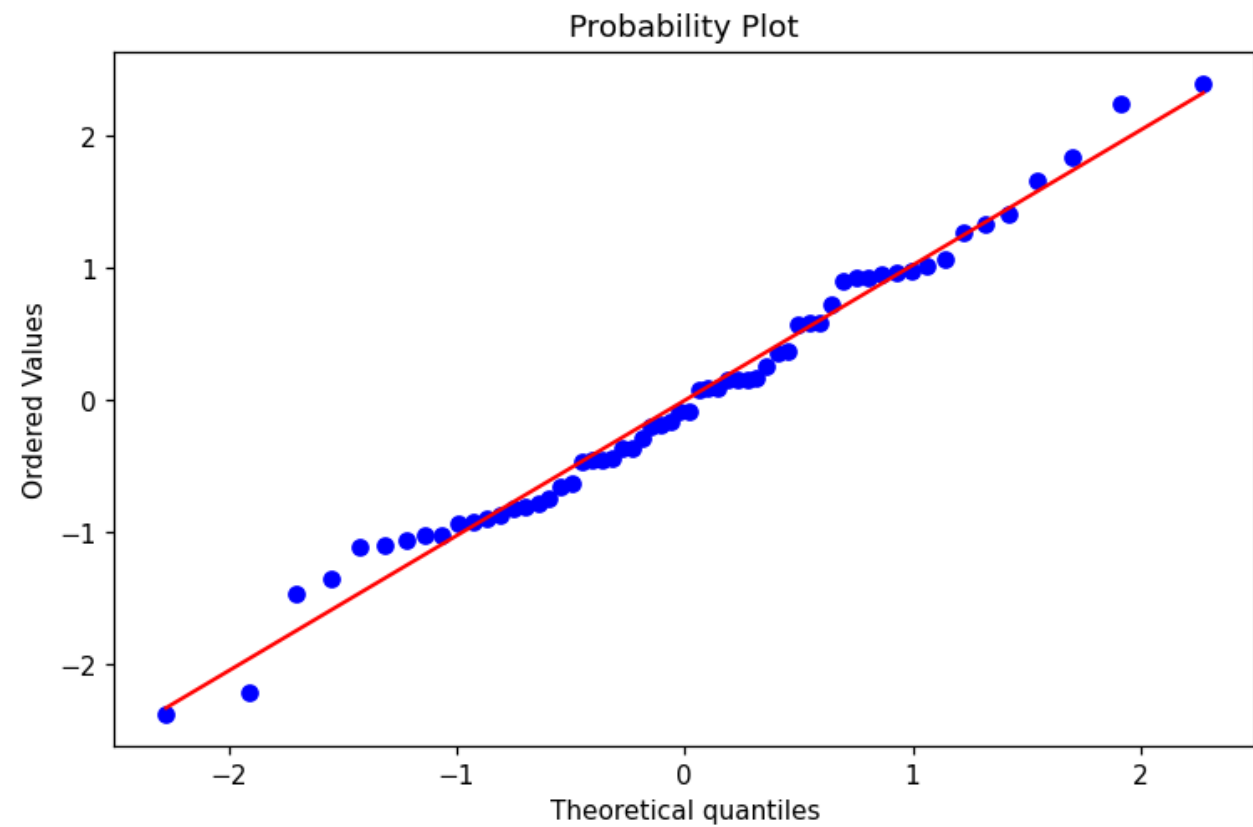
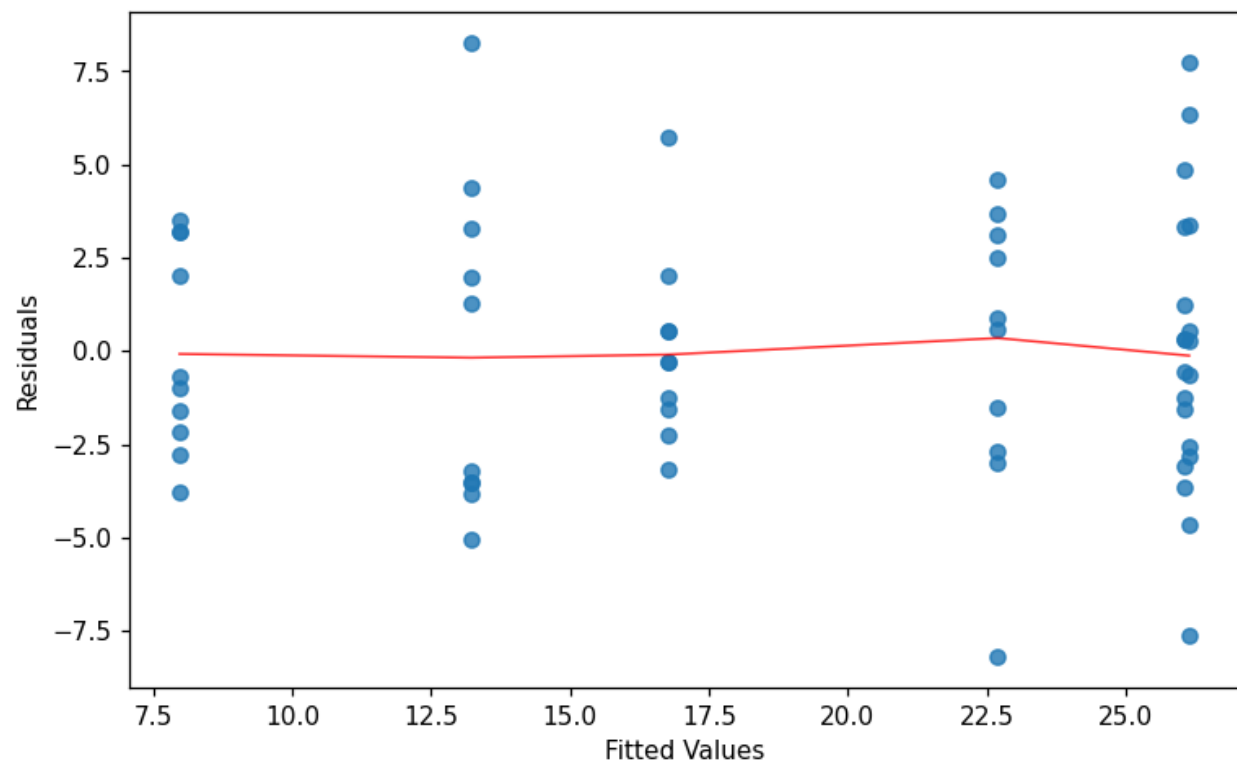
```
from statsmodels.graphics.factorplots import interaction_plot

interaction_plot(ToothGrowth['dose'], ToothGrowth['supp'], ToothGrowth['len'],
                 colors=['orange', 'blue'], markers=['D', '^'])
plt.show()
```



Interpreting Interaction Plots

- Parallel lines - No interaction occurs.
- Nonparallel lines - An interaction occurs. The more nonparallel the lines are, the greater the strength of the interaction.
- Can see that the lines are parallel between D0.5 and D1 for two different supplement types with OJ leading to longer teeth but then there is an interaction when the dose is increased to 2 and both supplements perform the same.



Example with interaction term

```
model2int= ols('len ~ C(supp)*C(dose)', data=ToothGrowth).fit()  
model2int= ols('len ~ C(supp)+C(dose)+C(supp):C(dose)', data=ToothGrowth).fit()  
# These two calls are equivalent
```

```
anova2wayint = anova_lm(model2int, typ=1)
```

```
anova2wayint
```

	df	sum_sq	mean_sq	F	PR(>F)
C(supp)	1.0	205.350000	205.350000	15.571979	2.311828e-04
C(dose)	2.0	2426.434333	1213.217167	91.999965	4.046291e-18
C(supp):C(dose)	2.0	108.319000	54.159500	4.106991	2.186027e-02
Residual	54.0	712.106000	13.187148	NaN	NaN

OLS Regression Results

```

=====
Dep. Variable:          len    R-squared:                0.794
Model:                  OLS    Adj. R-squared:           0.775
Method:                 Least Squares    F-statistic:        41.56
Date:                  Wed, 01 Jan 2020    Prob (F-statistic):   2.50e-17
Time:                  00:00:00    Log-Likelihood:      -159.35
No. Observations:      60    AIC:                330.7
Df Residuals:          54    BIC:                343.3
Df Model:              5
Covariance Type:       nonrobust
=====

```

```

=====
               coef      std err      t      P>|t|    [0.025    0.975]
-----
Intercept          13.2300     1.148    11.521    0.000    10.928    15.532
C(supp)[T.VC]       -5.2500     1.624    -3.233    0.002    -8.506    -1.994
C(dose)[T.1.0]       9.4700     1.624     5.831    0.000     6.214    12.726
C(dose)[T.2.0]      12.8300     1.624     7.900    0.000     9.574    16.086
C(supp)[T.VC]:C(dose)[T.1.0] -0.6800     2.297    -0.296    0.768    -5.285     3.925
C(supp)[T.VC]:C(dose)[T.2.0]  5.3300     2.297     2.321    0.024     0.725     9.935
=====

```

- From the ANOVA results and based on a significance level of 0.05, you can conclude that:
 - the p-value of supp is 2.311828×10^{-4} , which indicates that the levels of supp are associated with significant different tooth length.
 - the p-value of dose is 4.046291×10^{-18} , which indicates that the levels of dose are associated with significant different tooth length.
 - the p-value for the interaction between supp*dose is 0.02, which indicates that the relationships between dose and tooth length depends on the supp method.


```
ToothGrowth.groupby('dose').mean()
```

```
Out[99]:
```

```
len
```

```
dose
```

```
0.5  10.605
```

```
1.0  19.735
```

```
2.0  26.100
```

```
ToothGrowth.groupby('supp').mean()
```

```
Out[100]:
```

```
len
```

```
dose
```

```
supp
```

```
OJ      20.663333  1.166667
```

```
VC      16.963333  1.166667
```

```
ToothGrowth.groupby(['dose', 'supp']).mean()
```

```
Out[101]:
```

```
len
```

```
dose
```

```
supp
```

```
0.5    OJ      13.23
```

```
VC      7.98
```

```
1.0    OJ      22.70
```

```
VC      16.77
```

```
2.0    OJ      26.06
```

```
V      C      26.14
```

Unbalanced two-way ANOVA

- Balanced designs correspond to the situation where we have equal sample sizes within levels of our independent grouping levels.
- In experimental data, the experimenter will often set up the data with equal number of observations per cell.
- An **unbalanced design** has unequal numbers of samples in each group.
- `typ = "2"` can be used to compute two-way ANOVA test for unbalanced designs.

Unbalance ANOVA

```
anova2wayint = anova_lm(model2int, typ=2)
```

```
anova2wayint
```

```
Out[102]:
```

	sum_sq	df	F	PR(>F)
C(supp)	205.350000	1.0	15.571979	2.311828e-04
C(dose)	2426.434333	2.0	91.999965	4.046291e-18
C(supp):C(dose)	108.319000	2.0	4.106991	2.186027e-02
Residual	712.106000	54.0	NaN	NaN

Unbalanced vs Balanced

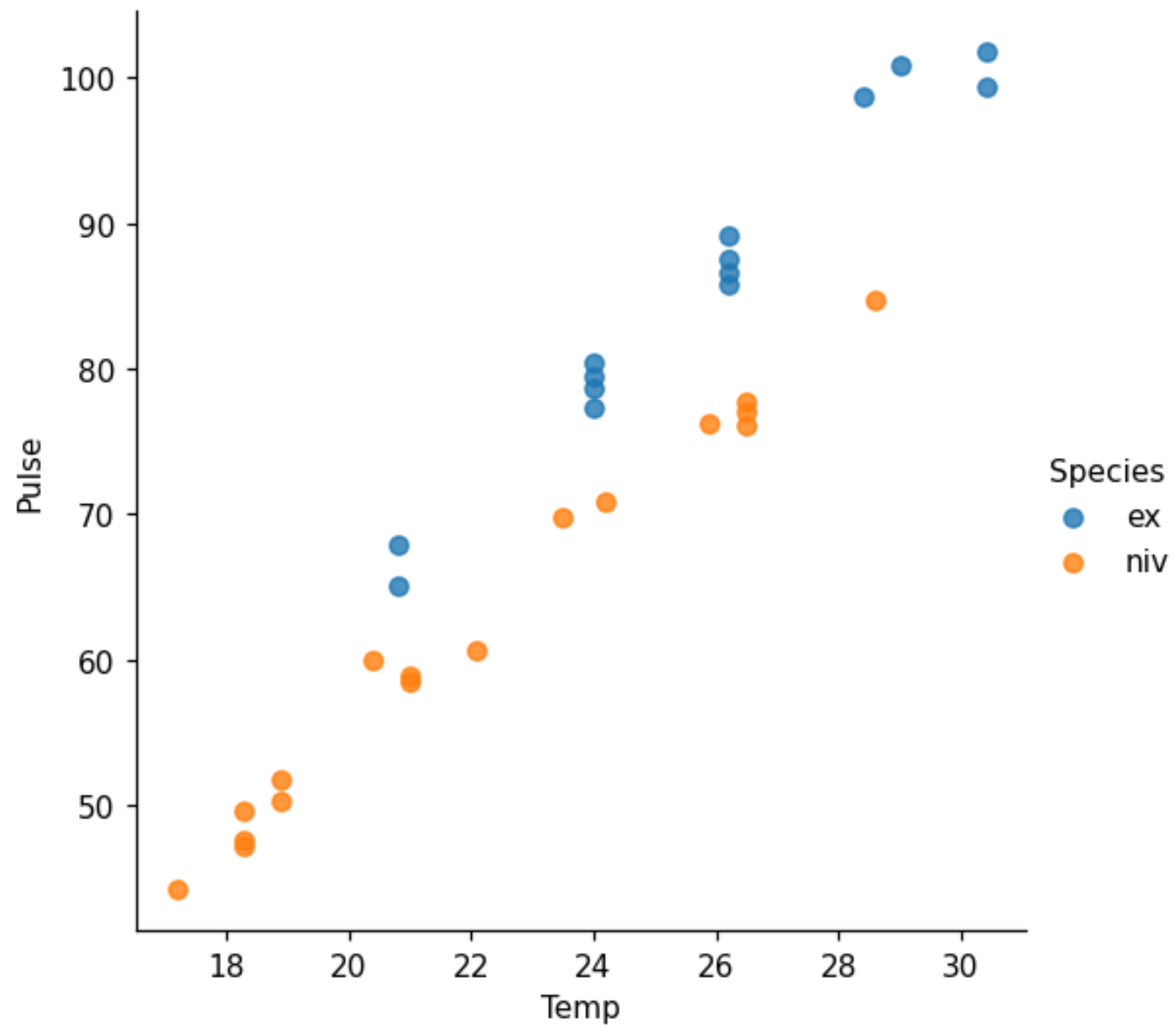
- Unbalanced design will always impact the power - the ability to detect significant differences. The power is limited by the size of the smallest cell.
- Unbalanced design usually impacts the ability to cleanly divide up the sums of squares and may end up with unexplained variance that is due to an effect but unable to say which effect. This is different than unexplained (residual) variance.
- Also it may mask an important relationship in the data.

ANCOVA

- Analysis of covariance (ANCOVA) is the same as a regression with one categorical and one continuous explanatory variables.
- ANCOVA evaluates whether the means of a dependent variable are equal across levels of a categorical independent variable, while statistically controlling for the effects of other continuous variables that are not of primary interest, known as covariates.
- The categorical variable divides the regressions into two or more sets.

Example: crickets

- Walker (1962) studied the mating songs of male tree crickets. Each wingstroke by a cricket produces a pulse of song, and females may use the number of pulses per second to identify males of the correct species.
- Walker wanted to know whether the chirps of the crickets *Oecanthus exclamationis* and *Oecanthus niveus* had different pulse rates.
- Measure the pulse rate of the crickets at a variety of temperatures from both species



Example: crickets - unbalanced

```
data['Species'].value_counts()
```

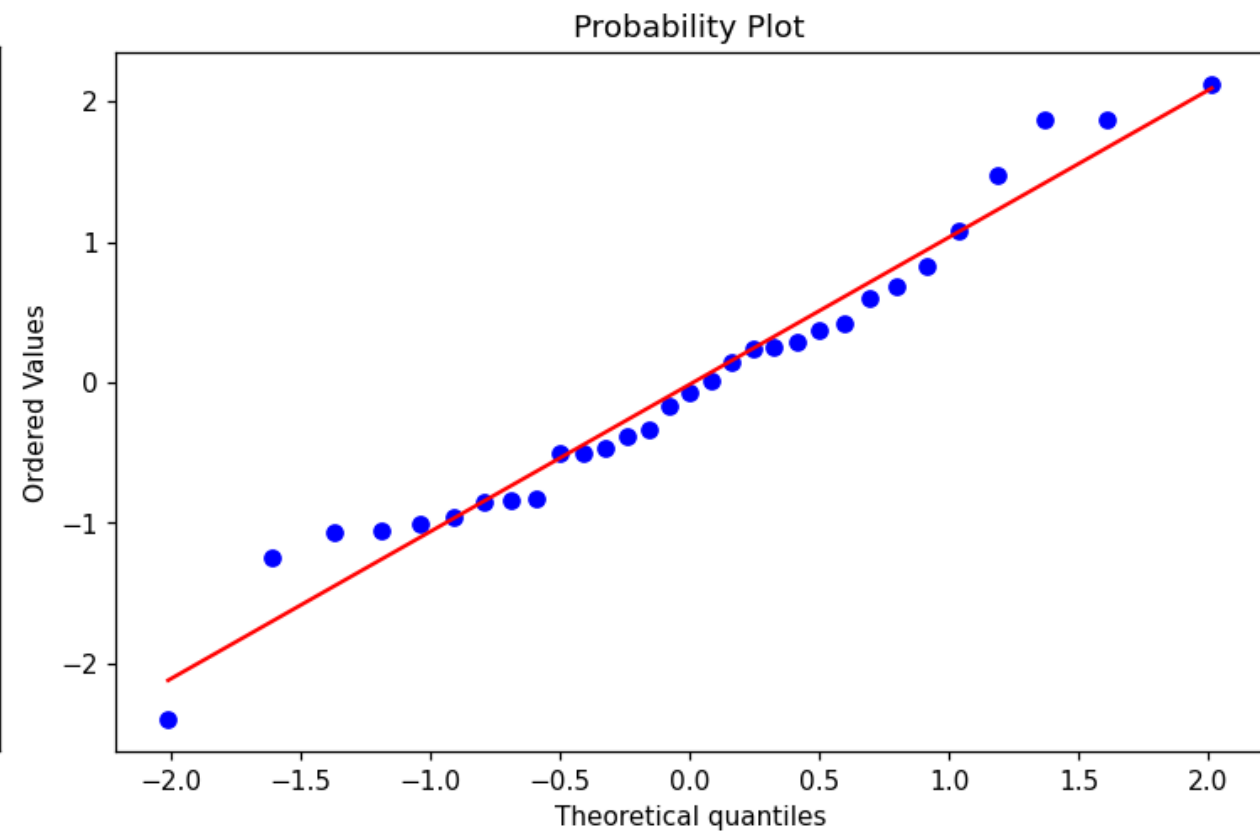
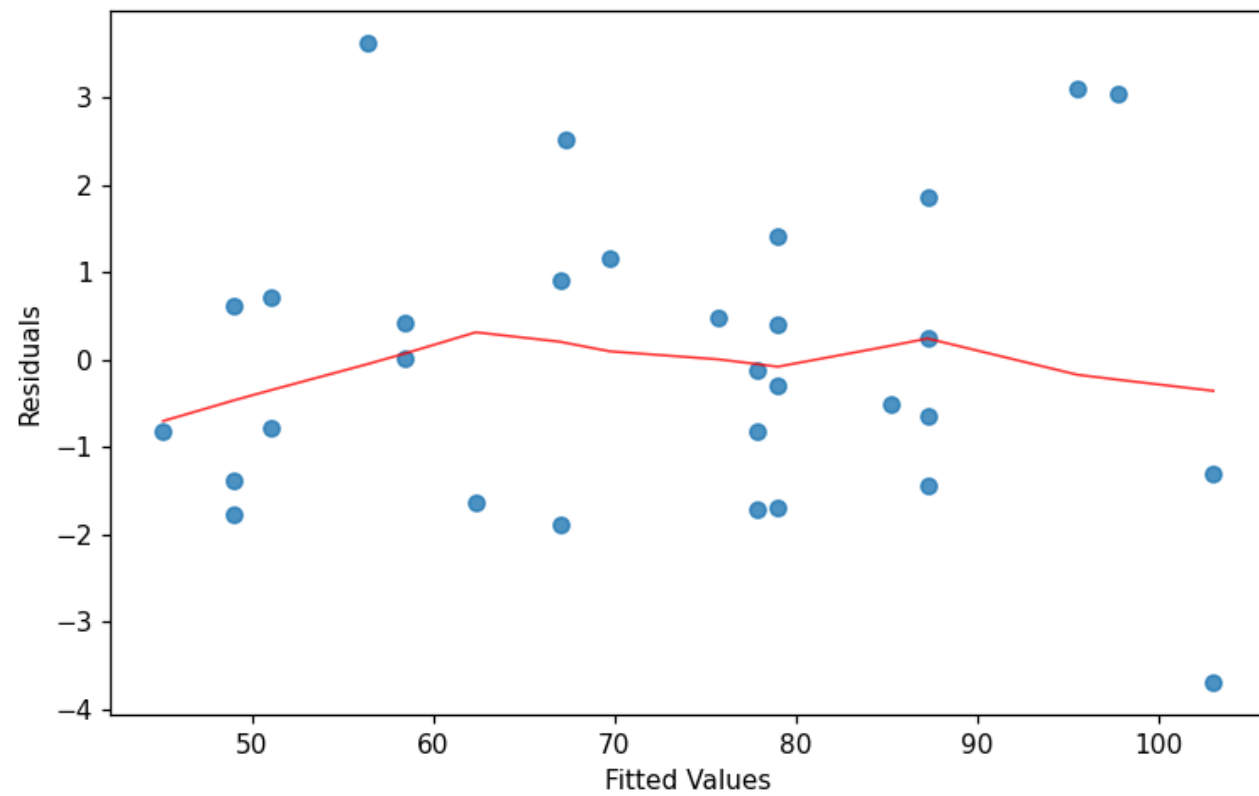
```
Out[115]:
```

```
niv  17
```

```
ex   14
```

```
Name: Species, dtype: int64
```

```
model= ols('Pulse ~ Temp * C(Species)', data=data).fit()
```

OLS Regression Results

```
=====
Dep. Variable:          Pulse  R-squared:                0.990
Model:                  OLS    Adj. R-squared:            0.989
Method:                 Least Squares  F-statistic:        898.9
Date:                  Wed, 01 Jan 2020  Prob (F-statistic):  3.77e-27
Time:                  00:00:00  Log-Likelihood:       -59.635
No. Observations:      31      AIC:                  127.3
Df Residuals:          27      BIC:                  133.0
Df Model:              3
Covariance Type:       nonrobust
=====
```

```
=====
               coef      std err      t      P>|t|      [0.025      0.975]
-----
Intercept      -11.0408     4.151    -2.659     0.013    -19.559     -2.523
C(Species)[T.niv] -4.3484     4.962    -0.876     0.389    -14.529      5.832
Temp           3.7514     0.160    23.429     0.000      3.423      4.080
Temp:C(Species)[T.niv] -0.2340     0.201    -1.165     0.254     -0.646      0.178
=====
```

```
=====
Omnibus:          0.829  Durbin-Watson:           1.623
Prob(Omnibus):    0.661  Jarque-Bera (JB):           0.615
Skew:             0.334  Prob(JB):                 0.735
Kurtosis:         2.828  Cond. No.                 531.
=====
```

Example: crickets - unbalanced

```
anova_model= anova_lm(model, typ=2)
```

```
anova_model
```

```
Out[120]:
```

	sum_sq	df	F	PR(>F)
C(Species)	598.003953	1.0	189.788769	9.906686e-14
Temp	4376.082568	1.0	1388.839184	9.350847e-25
Temp:C(Species)	4.275779	1.0	1.357006	2.542464e-01
Residual	85.074090	27.0	NaN	NaN

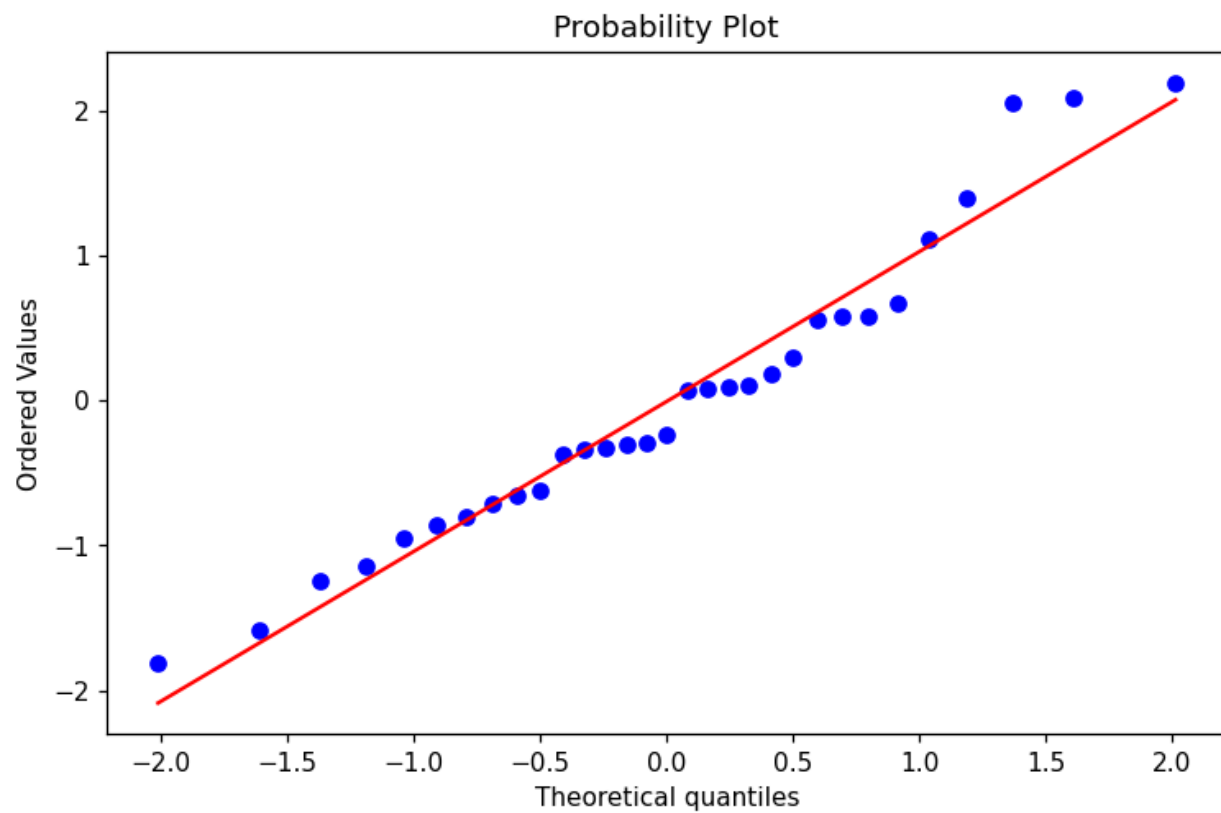
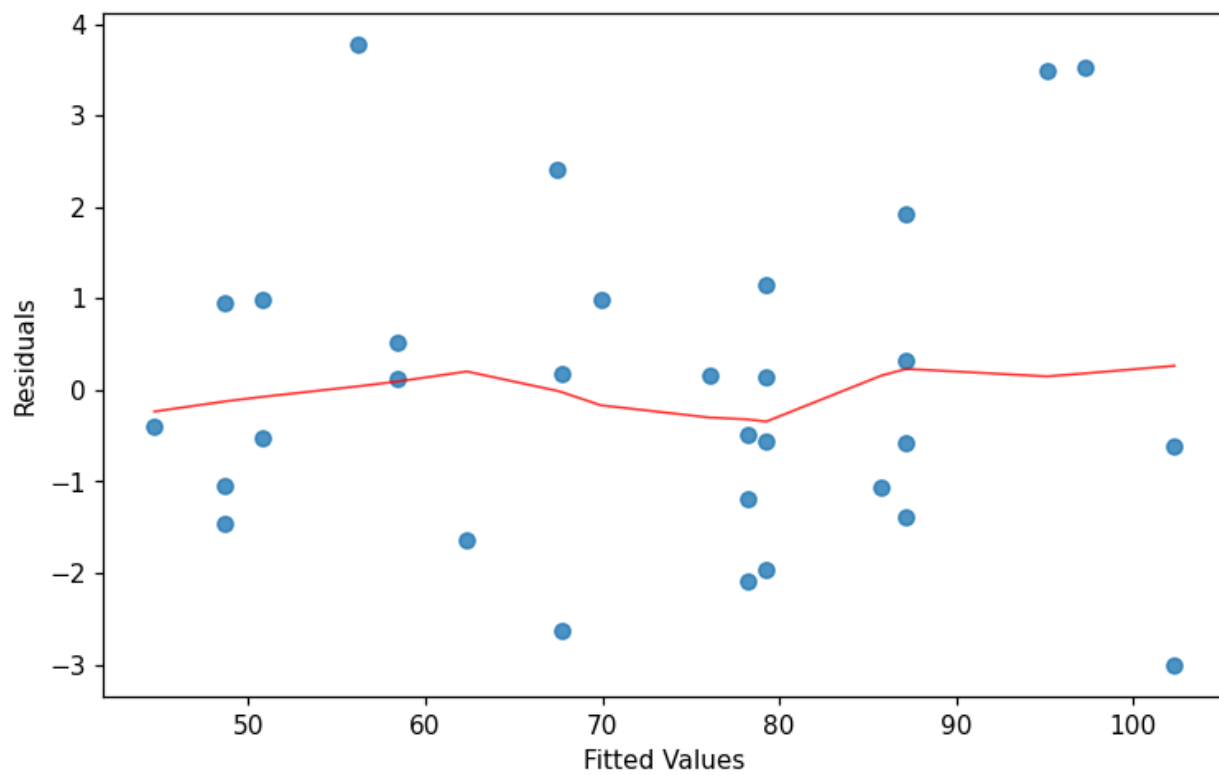
Example: crickets

- From the ANOVA results and based on a significance level of 0.05, you can conclude that:
 - the p-value of temp is 9.351×10^{-25} , which indicates that temperature and pulse have a significant relationship.
 - the p-value of Species is 9.907×10^{-14} , which indicates that the levels of species are associated with significant different pulse rates.
 - the p-value for the interaction between temp*Species is 0.2542, which indicates that the interaction term is not significant, so the slope across species levels is not different.

Example: crickets

- Rerun without interaction term:

```
model2= ols('Pulse ~ Temp + C(Species)', data=data).fit()
```



OLS Regression Results

```

=====
Dep. Variable:          Pulse  R-squared:                0.990
Model:                  OLS    Adj. R-squared:            0.989
Method:                 Least Squares  F-statistic:        1331
Date:                  Wed, 01 Jan 2020  Prob (F-statistic):  1.76e-28
Time:                  00:00:00  Log-Likelihood:      -60.395
No. Observations:      31      AIC:                  126.8
Df Residuals:          27      BIC:                  131.1
Df Model:              3
Covariance Type:       nonrobust
=====

```

```

=====
               coef      std err      t      P>|t|      [0.025      0.975]
-----
Intercept      -7.2109      2.551     -2.827     0.009    -12.436     -1.986
C(Species)[T.niv]  -10.0653      0.735    -13.689     0.000    -11.571     -8.559
Temp           3.6028      0.097     37.032     0.000      3.403      3.802
=====

```

```

=====
Omnibus:          2.343  Durbin-Watson:           1.509
Prob(Omnibus):    0.310  Jarque-Bera (JB):           1.789
Skew:             0.586  Prob(JB):                   0.409
Kurtosis:         2.892  Cond. No.                   195.
=====

```

Example: crickets - unbalanced

```
anova_model2= anova_lm(model2, typ=2)
```

```
anova_model2
```

```
Out[124]:
```

	sum_sq	df	F	PR(>F)
C(Species)	598.003953	1.0	187.399388	6.271533e-14
Temp	4376.082568	1.0	1371.354138	2.487732e-25
Residual	89.349869	28.0	NaN	NaN

Example: crickets

- From the ANOVA results and based on a significance level of 0.05, you can conclude that:
 - the p-value of temp is 2.488×10^{-25} , which indicates that temperature and pulse have a significant relationship.
 - the p-value of Species is 6.272×10^{-14} , which indicates that the levels of species are associated with significantly different pulse rates.
 - This leads to different intercepts among the groups

ANCOVA – three different models

1. Common slope and intercept

$$y = \beta_0 + \beta_1 x + \varepsilon$$

Here x is Temp and y is Pulse rate

2. Common slope – different intercepts

$$y = \beta_0 + \beta_1 x + \beta_2 z_2 + \varepsilon$$

Here z_2 is an indicator variable for niv species

3. Separate lines – different intercepts and different slopes

$$y = \beta_0 + \beta_1 x + \beta_2 z_2 + \beta_3 x \times z_2 + \varepsilon$$

Example: crickets

2. Common slope – different intercepts

$$y = -7.2109 + 3.6028 x - 10.0653z_2 + \varepsilon$$

- When $z_2 = 0$:

$$y = -7.2109 + 3.6028 x + \varepsilon$$

- When $z_2 = 1$:

$$y = -17.2762 + 3.6028 x + \varepsilon$$

```
sns.lmplot(x='Temp', y='Pulse', hue='Species', data=data, fit_reg=False)
x=data['Temp']
b, m = -7.2109, 3.6028,
plt.plot(x, b+ m*x, color='blue', linestyle='dashed')
b, m = -17.2762, 3.6028
plt.plot(x, b+ m*x, color='orange', linestyle='dashed')
```

