

Investigation 7

How much is too much?

Investigating the effect of salt on seed germination



Background information, photos, data,
and instructions

In this investigation, we will conduct an experiment to answer the question, “Does salt affect the germination of lentils?”



a field of lentil plants



lentils are produced in pods

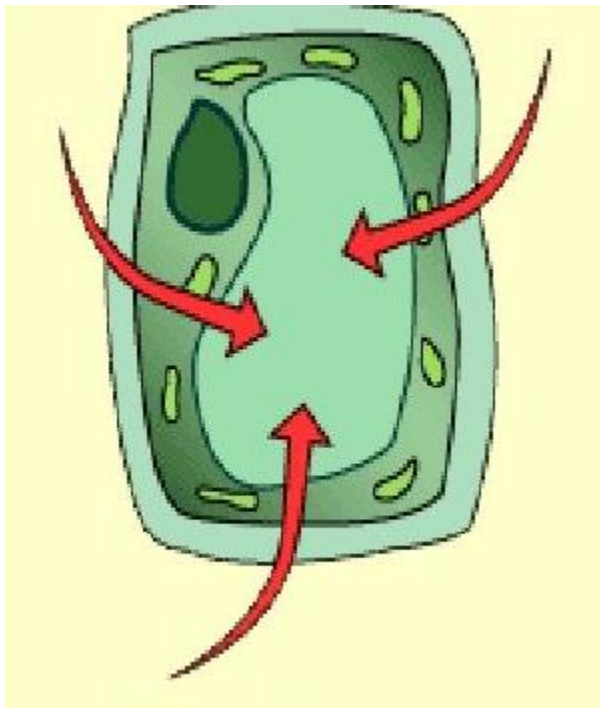


each lentil is a tiny bean

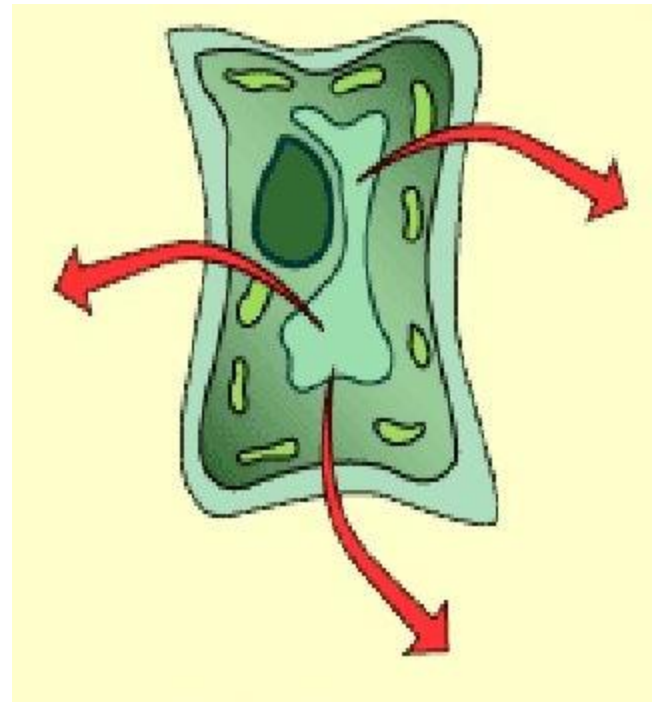
Seed germination requires warmth and water



For a seed to sprout, its cells must absorb water by osmosis (arrows show water movement)



In salty conditions, osmosis causes water to flow OUT of cells instead of into them



Since you won't be able to set up this investigation yourself, your lab instructors set it up. As you view the following photos, refer to the corresponding pages in your lab manual.

Step 1: Prepare your seeds for testing



counting out 80 lentils
(10 per salt concentration)



soaking lentils in bleach
solution to kill mold spores
that can interfere with
germination

Step 2: Prepare germination chambers



cutting out three layers of paper towels for an absorbent surface



placing paper towel layers in each germination chamber

Step 3: Prepare test solutions

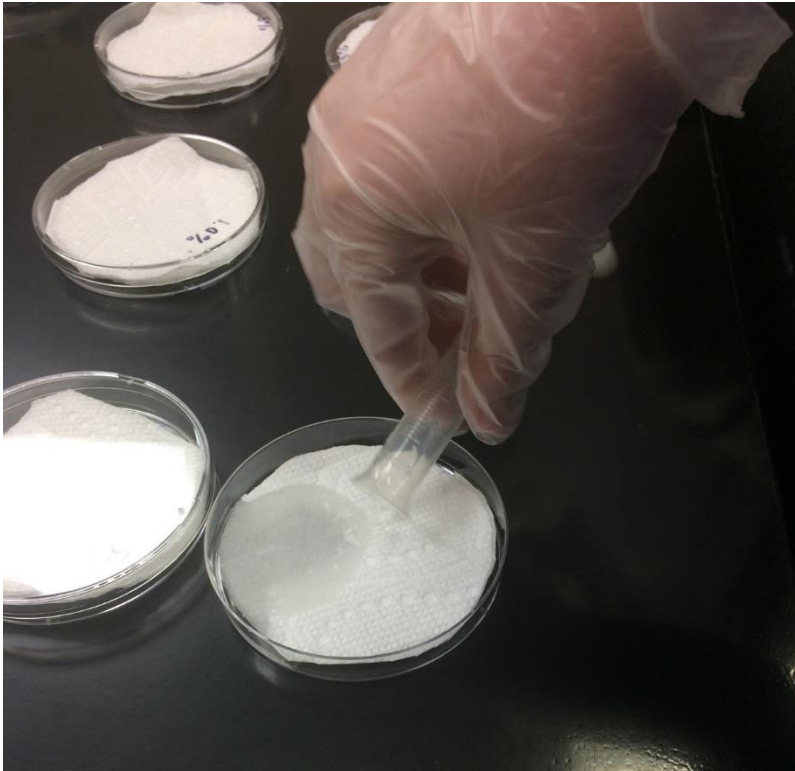


using graduated cylinder to measure 100 ml of water for each salt solution



using electronic balance to weigh the exact amount of salt needed for each salt solution (0%-3.5%)

Step 4: Expose seeds to test solutions

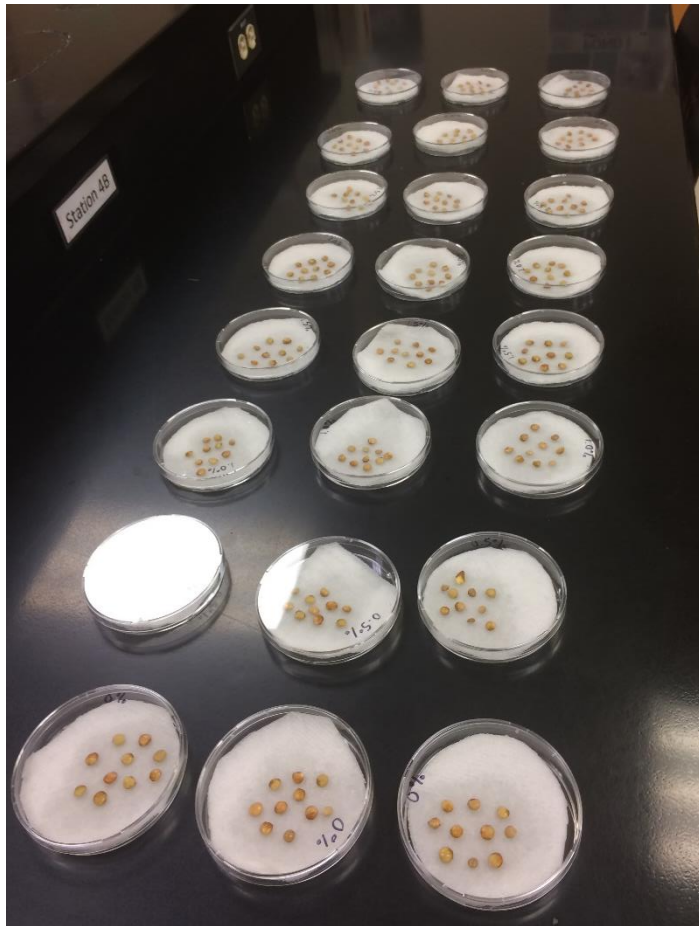


pouring 8 ml of each test solution into corresponding germination chamber



placing 10 lentils in each chamber

We set up the investigation for six lab tables.



germination chambers for
three lab tables



each table's germination
chambers stored in a Ziploc bag
to hold moisture...waiting
several days to observe results

Before viewing the results of this experiment, you should examine the **practice data at the bottom of p. 79 in your lab manual.**

The table on p. 79 shows the number of seeds that germinated, out of 10 seeds that were incubated in various salt concentrations at room temperature for several days. (This is a **sample** data set so that you can become familiar with how to analyze the data.)

You will use online resources to calculate mean (average) seed germination and determine if salt has a significant effect on germination.

Your online tool for calculating means and comparing them statistically is at <https://uca.edu/biology/biology-1400-01-02/>. When you open that page, click on ANOVA:Analysis of Variance Between Groups.

The screenshot shows a web browser window with the URL <https://goodcalculators.com/one-way-anova-calculator/>. The interface includes three data entry fields for Group 1, Group 2, and Group 3. Red arrows point from text boxes to the input fields and the 'Calculate' button. The text boxes provide instructions on how to use the calculator, including deleting demo data and entering specific values for each group. The 'Calculate' button is highlighted in blue.

Group 1
5, 1, 11, 2, 8

Delete the demo data and enter data for Group 1 (0% salt):
8, 7, 8, 7, 8, 7

Group 2
0, 1, 4, 6, 3

Delete the demo data and enter data for Group 2 (0.5% salt)

Group 3
13, 9, 8, 15, 7

After entering Group 3 (1.0% salt) data, click "add group." Continue adding groups so that you can enter data for all 8 salt concentrations.

+ Add Group - Delete Group

Calculate

Click "Calculate" to conduct the statistical test and view results

Scroll down below the blue “Calculate” button to view your results.

There are lots of results, but you should focus on:

- 1) the mean germination for each salt concentration (from the “Data Summary” chart). Copy the mean germination into the bottom row of your **practice** data table on p. 79
- 2) the P-value right above the “Data Summary” chart. If the number that follows “P-value = “ is greater than 0.05, the effect of salt on seed germination is NOT statistically significant. If the number is 0.05 or below, the effect of salt on seed germination is statistically significant.

A very, very small P-value may round to 0
(in that case, the P-value is far below 0.05, so the difference is statistically significant)

After analyzing the **practice** data, you're ready to view the actual results from this experiment and analyze them (exactly as you did for the practice data).

After four days, we counted the number of lentils that had sprouted in each germination chamber and recorded the data for the first five lab tables.

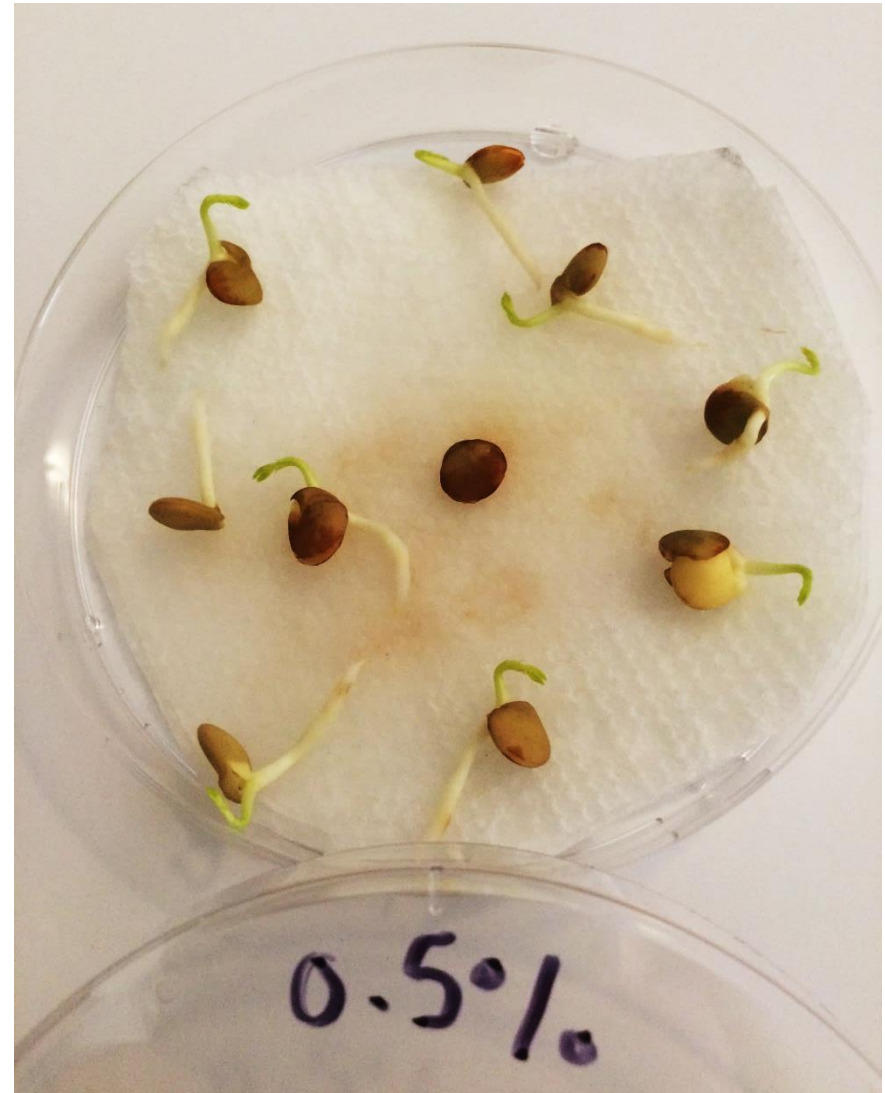
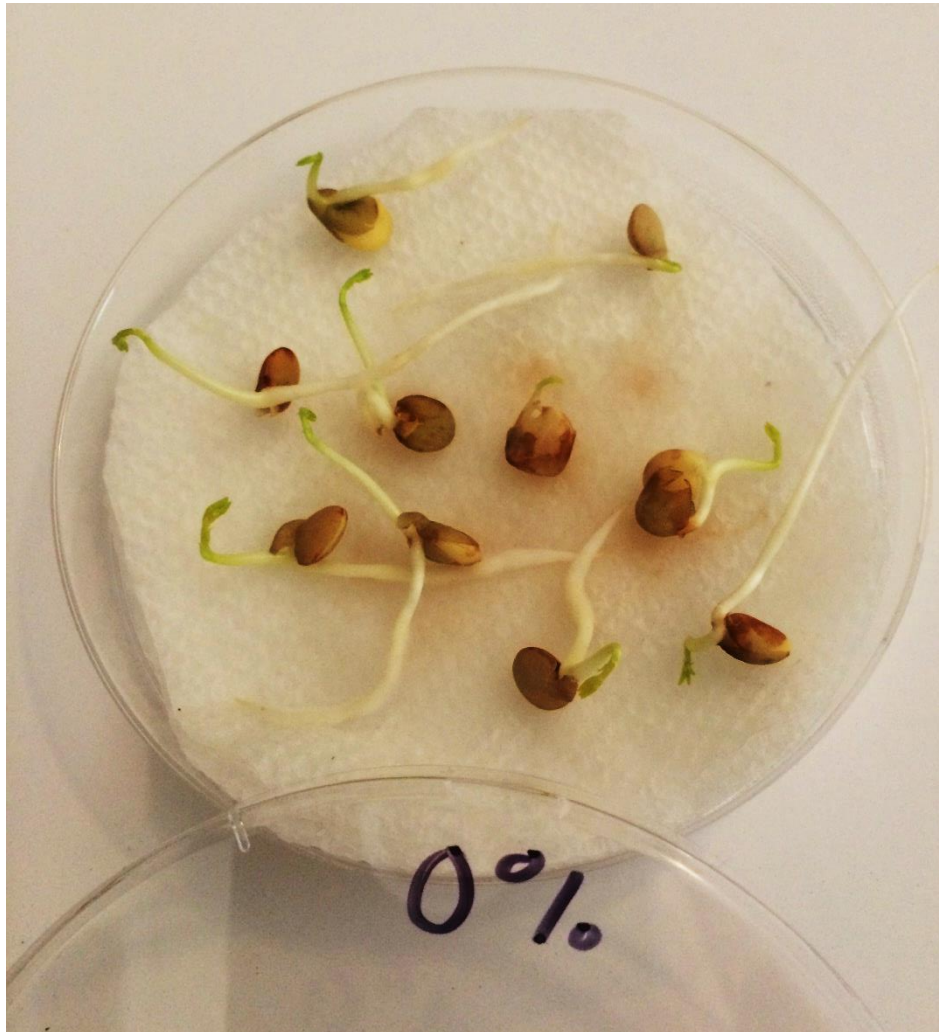
You should enter these data into the blank chart on p. 81 of your lab manual.

Lab table	0% (A)	0.5% (B)	1.0% (C)	1.5% (D)	2.0 % (E)	2.5% (F)	3.0% (G)	3.5% (H)
1	9	9	9	9	0	0	0	0
2	9	10	9	6	0	0	0	0
3	10	10	7	6	0	0	0	0
4	10	9	9	3	4	0	0	0
5	10	9	8	8	0	0	0	0
6	Table 6 data will come from the following photos							



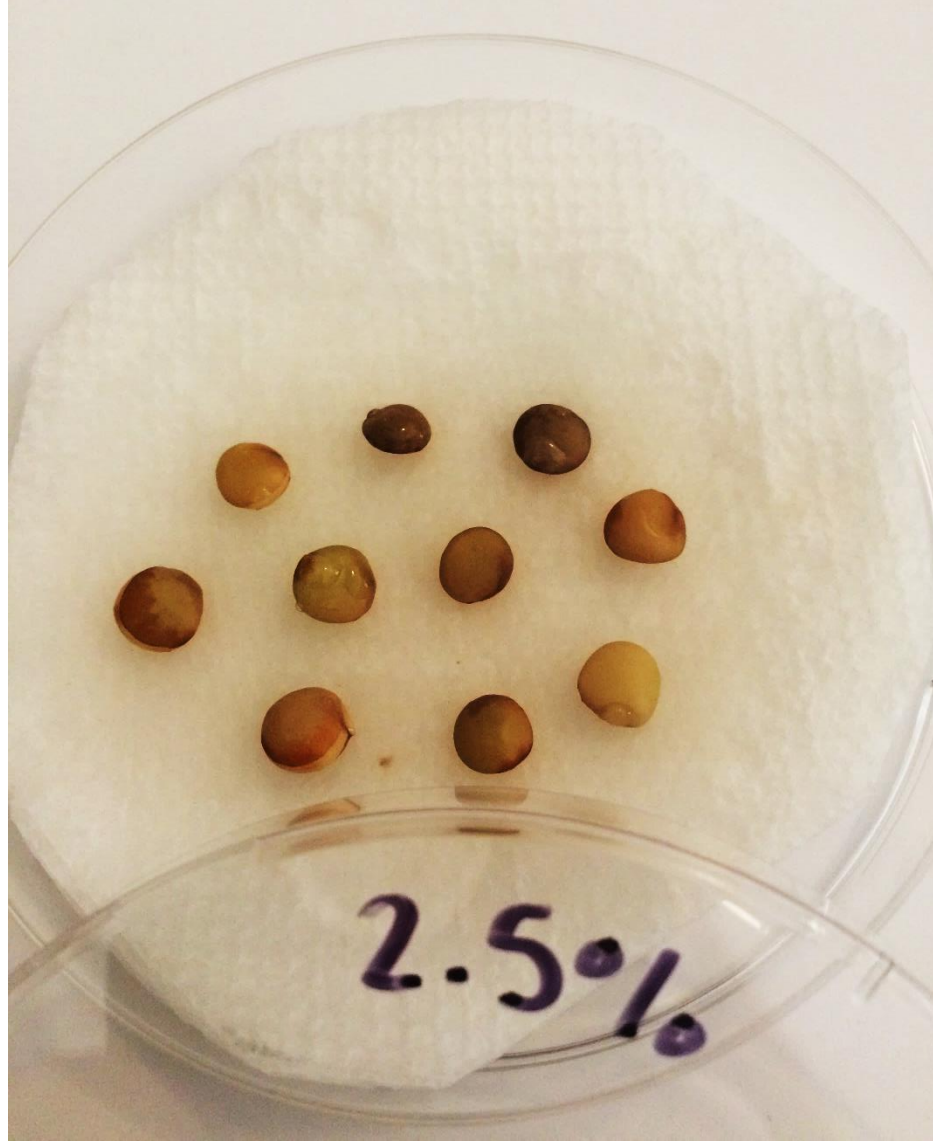
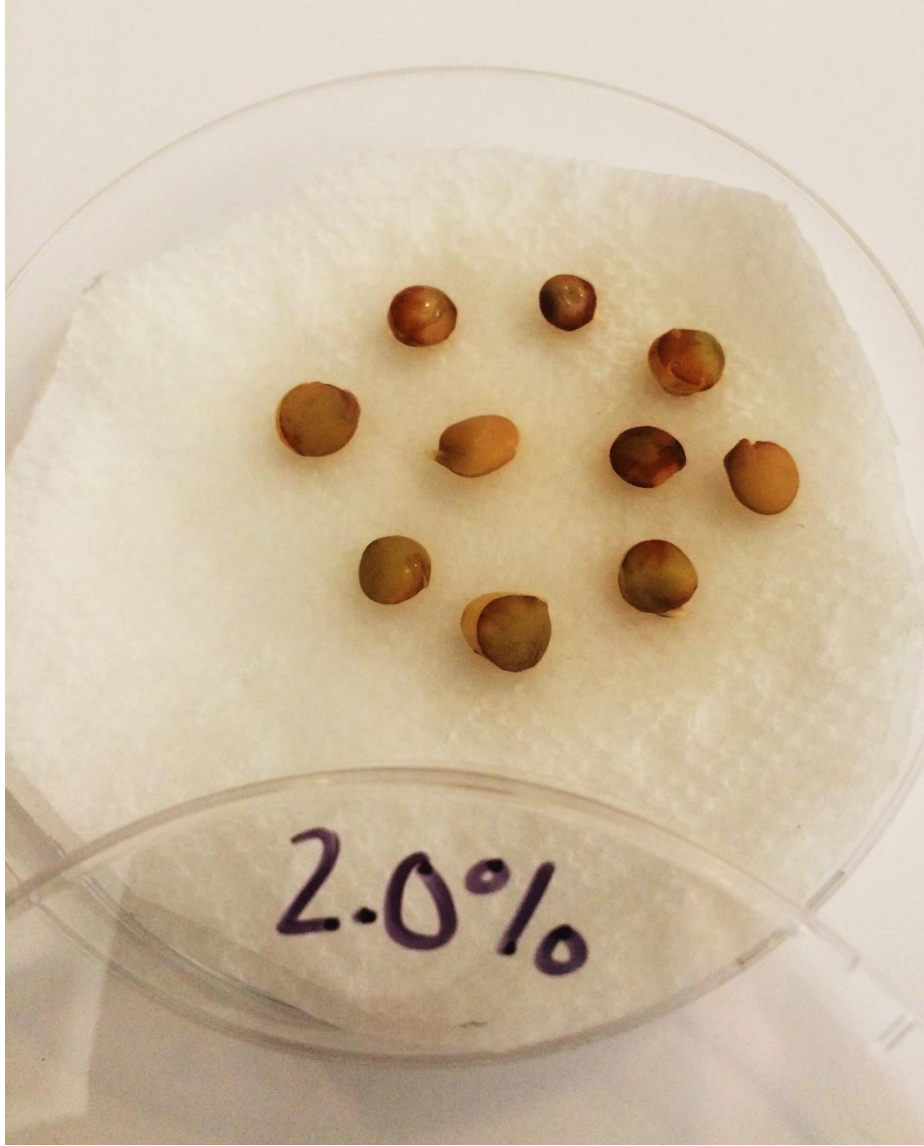
To complete the bottom row of this data table, use the following pictures to count and record the number of seeds that sprouted in each salt concentration, enter those data into the blank chart, and include all six tables in your analysis.

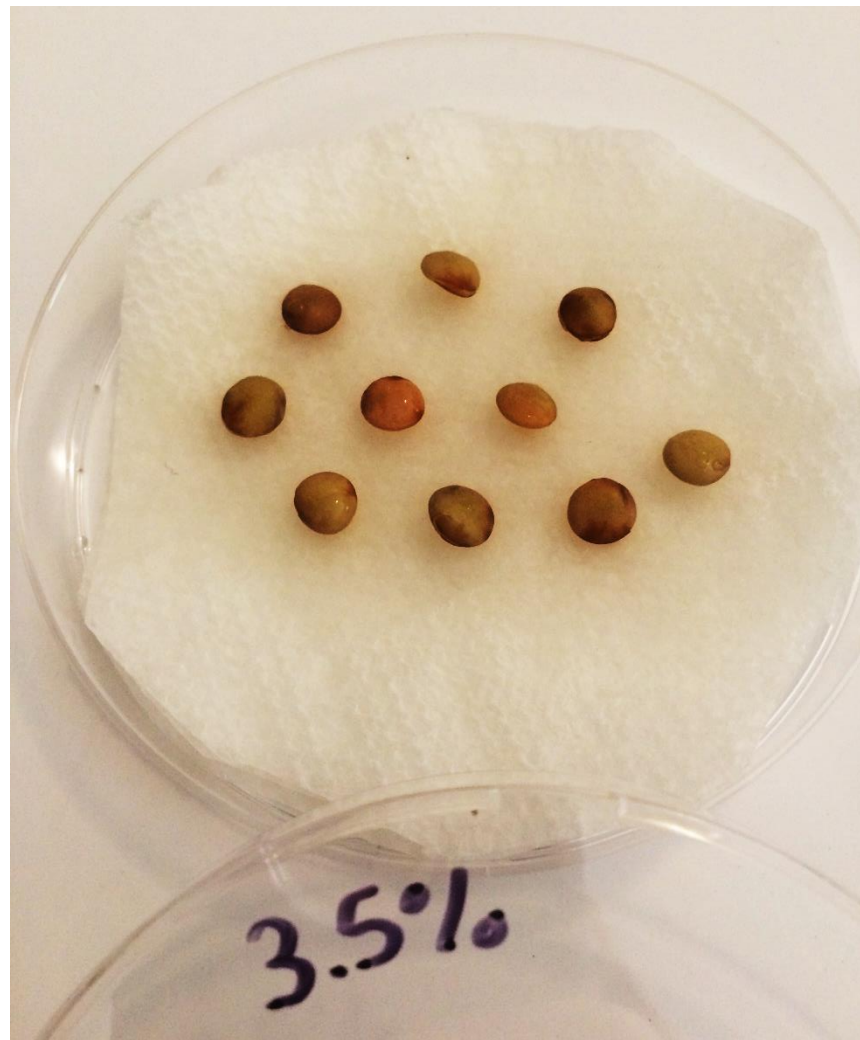
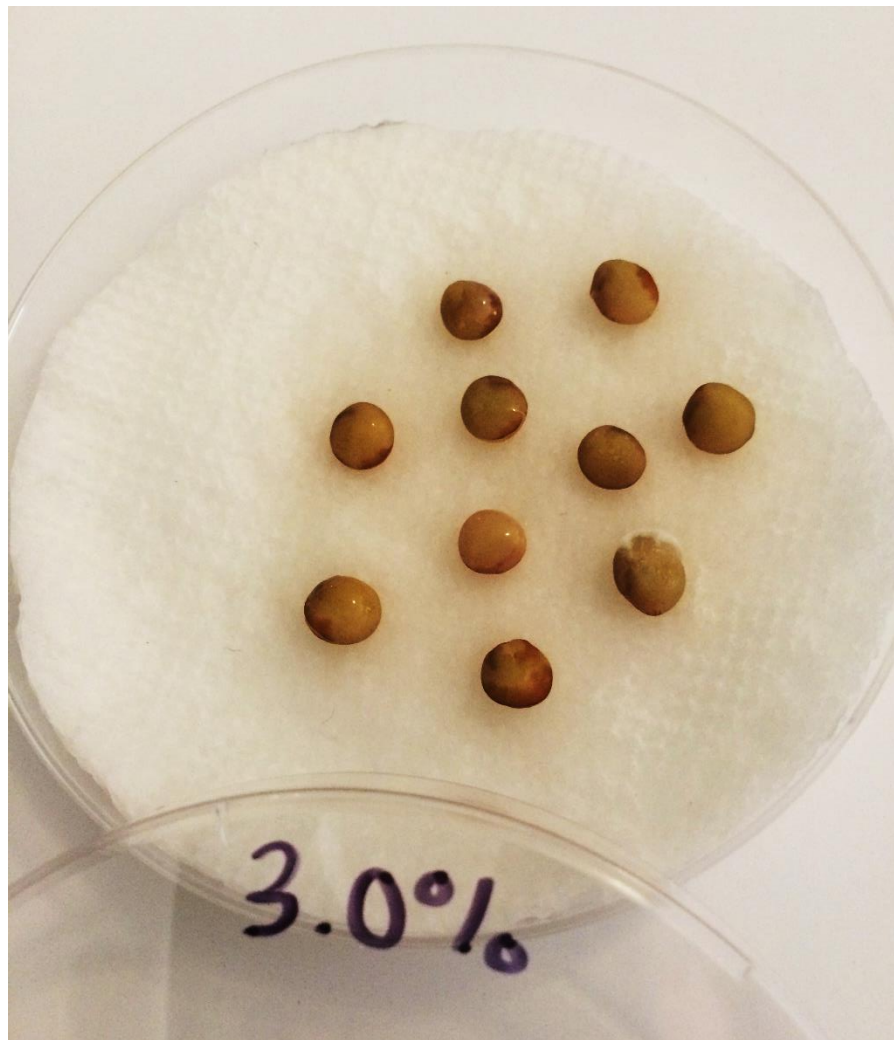
**You should count any lentils that have sprouted,
even if the “sprouts” are small.**



Even though these
"sprouts" are small, they
still count







Now, you should analyze the results from the table on p. 81 exactly as you did for the **practice** data (using the ANOVA link, entering the data, and interpreting the results).

If the P-value is a very, very small number, it may be rounded to 0 (in that case, the P-value is far below 0.05, so the difference is statistically significant)

Complete the pre-lab assignment on p. 77 and the data sheets on pp. 79-82 in your lab manual.

You do **NOT** need to answer any questions on p. 83.

You will turn in your completed work at the **start** of your next in-person lab.