

Final Project Report

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Abstract

The invasive thistle *Carduus nutans* has been a management concern for landowners and governments alike, with dispersal in this plant playing an important role in its spread. Given that the terminal velocity of *C. nutans* seeds strongly influences dispersal distance, we seek to better understand how both pappus shape and treatment of the parent plant affect seed terminal velocity. Our analyses show that the width of the seed pappus was the most important factor in determining seed terminal velocity and that the two were negatively correlated, with higher pappus width generally giving rise to a lower terminal velocity. Warming and mowing treatments on the parent plants had significant effects on the pappus width, with warmed and mowed plants having a much smaller pappus width. Regarding the effects of the treatments on seed terminal velocity, early mowing increased terminal velocity, while warming decreased terminal velocity; no significant effect was observed for late mowed plants compared to control plants. Given that mowing was shown to increase pappus width and that increased pappus width was shown to reduce terminal velocity, it is surprising that plants experiencing the early mowing treatment had higher seed terminal velocity. Such a contradiction merits further investigation of additional factors, such as mass, that may affect seed terminal velocity.

Introduction and Data

As the world has become significantly more interconnected, the increased movement of people across the world has also facilitated an increase in the spread of species outside of their native ranges. When species that are introduced outside of their native range become invasive, they can have a detrimental impact on local biodiversity, quantity and quality of valuable ecosystem services, and other aspects of the local ecosystem. Because of this, the study of invasive species has been a large part of the ecology literature in recent decades, ranging from theoretical models of how they spread to investigating possible management practices that keep them at bay.

One particular invasive species that has greatly expanded its range due to human activity is *Carduus nutans*, also known as “musk thistle” or “nodding thistle”. This thistle is native to Europe and Central Asia, but has expanded its range into North America, Australia, and New Zealand, among other parts of the world¹. Within the U.S., this thistle has been reported in all U.S. states except for Alaska, Florida, Hawaii, Maine, and Vermont²; the thistle may even be present in these states, but has not yet been reported. It is also been reported all Canadian provinces except Nunavut, Northwest Territories, and Yukon Territories².

C. nutans is considered to be a noxious weed in many U.S. states for several reasons. Because it can occur in very large numbers and grow to be quite large, this thistle may form dense and often impenetrable stands. The plant is also covered in numerous large spines, making it painful when touched as well as unpalatable to grazing animals. The adverse impacts of this weed on grazing can also lead to substantial economic losses.

Another reason why there is concern over *C. nutans* is because it has a high potential to spread locally when introduced. This thistle is wind dispersed and seeds have a large pappus, or a feathery and lightweight modified calyx, on the seeds that allows them to be transported great distances. Models have been proposed

to model such wind-driven seed dispersal³, and such models have been applied to *C. nutans*⁴, showing significant potential for long-range dispersal events. While there are abiotic and biotic factors that affect how far a seed like those in *C. nutans* can be dispersed, a noteworthy predictor of dispersal distance is seed terminal velocity. For seeds, a higher terminal velocity generally means a decreased dispersal distance; this is because a higher terminal velocity means the seed falls faster and thus spends less time in the air, which means less of an opportunity for wind to carry it further from its source.

However, it is not entirely clear what affects terminal velocity in *C. nutans* seeds, though the most obvious candidates would be physical properties of the seed such as shape and mass. In general, seeds with a larger area perpendicular to the direction of motion will have higher drag and a lower terminal velocity. Seeds with a higher mass will have a higher downward force (mg) from gravity and thus a higher air resistance force that must equal it to achieve terminal velocity, which leads to a higher terminal velocity since said resistance force is proportional to that velocity. However, the physical properties of the seed may be affected by the morphology and physiology of the parent plant; abiotic and biotic factors can affect the parent plant in such a way that may ultimately influence the terminal velocity of its seeds.

Given that there may be a link between abiotic influences on *C. nutans* and the terminal velocity of its seeds, we wish to investigate whether certain treatments applied to the plant before it flowers have any effect on seed dispersal capabilities. Any treatment effects that can reduce the dispersal capability of these thistles may then be used to inform management decisions. By using mowing treatments as well as warming treatments (and combinations of the two), we will examine the effects of said treatments on seed terminal velocity and thus on dispersal capability.

The data used to assess the effects of mowing and warming on seed terminal velocity were collected during a field experiment that involved applying these treatments to parent plants and collecting their flower heads after they had set seed. The experimental setup involves ten blocks, with each block containing two plots: one plot with a warming treatment and one without. Within each plot there are three positions: one with an early mowing treatment, one with a late mowing treatment, and one with no mowing at all. Overall, this yields six unique combinations of warming and mowing, with 10 replicates for each combination. Ten seeds were planted at each position, and one flower head was harvested from all individuals that survived to harvest date. Seeds were collected from individual flower heads and their terminal velocities were determined by timing the amount of time it took them to fall through a 1.25-m drop chamber. For each seed, these drop tests were repeated until two consecutive drop times were recorded within 0.1 seconds of each other.

Using the data from the experiment described above, we seek answers to three research questions of interest:

1. Is seed terminal velocity predicted by the shape of the seed pappus?
2. To what extent do warming and mowing treatments change seed terminal velocity?
3. Can changes in terminal velocity by treatment be explained by changes in seed shape parameters?

Research Question 1

First, we begin by investigating how pappus dimensions affect seed terminal velocity, with a total of six different measurements on the pappus used as predictors. The first two measurements are **SeedWidth** and **SeedLength**, which are respectively the basal width and slant height of a cone-shaped pappus. These measurements are empirically derived and are both measured in millimetres. Four additional measurements are calculated using **SeedWidth** and **SeedLength** and are as follows:

1. ‘SeedAngle’, the vertex angle of the conical pappus (radians)
2. ‘SeedArea’, the basal area of the conical pappus (mm^2)
3. ‘SeedHeight’, the height of the conical pappus (mm)
4. ‘SeedVol’, the volume of the conical pappus (mm^3)

Before any models are fit to the data, we first examine correlations between the various pappus measurements given above. A `pairs` plot of terminal velocity and the various physical properties can be seen in Figure 1. Linear associations between the variables can also be seen in Figure 2, with both the strength and directions of the correlations shown.

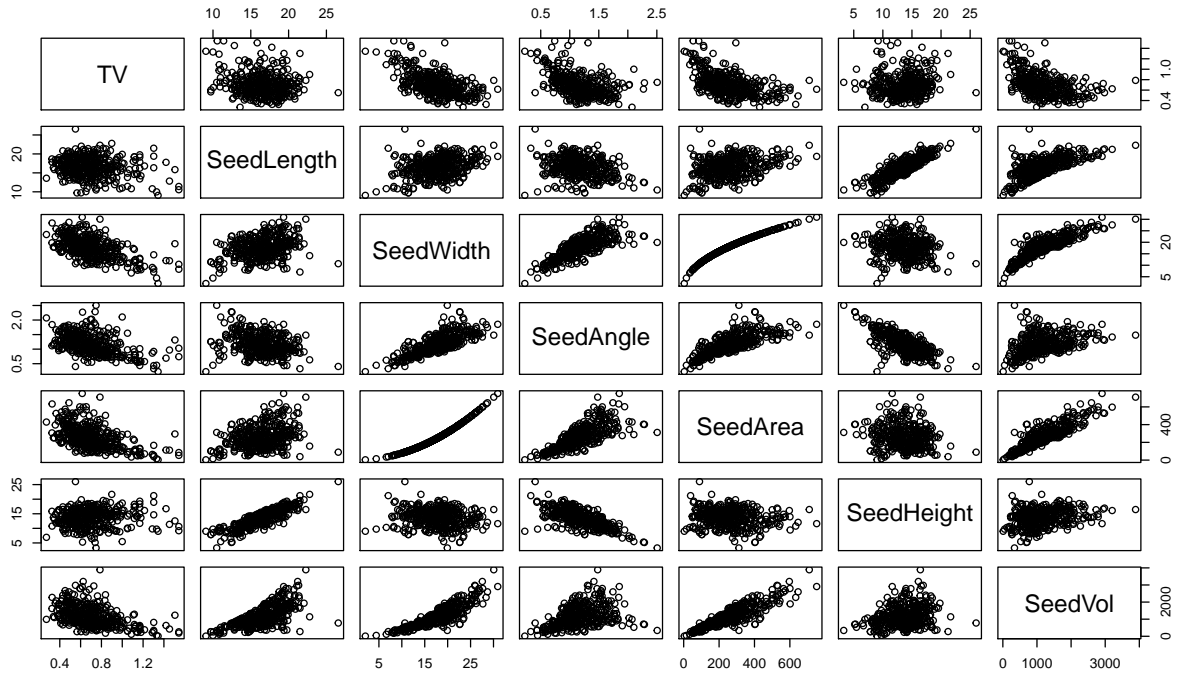


Figure 1: Plot of relationships between variables considered in this analysis.

Between the `pairs` plot and the correlation matrix/heatmap, it is clear that multicollinearity will be an issue since several of the variables are highly correlated with each other (e.g. `SeedWidth` and `SeedAngle`, `SeedWidth` and `SeedArea`, etc.). Given that several predictors are calculated from `SeedWidth` and `SeedHeight`, this is not surprising.

Our first model uses the terminal velocity `TV` as a response and all six of the pappus measurements as predictors. The resulting coefficients and their significance are listed in Table ?? . It is clear that this model is a poor fit for two main reasons. First, many of the terms are not significant to $\alpha = 0.05$ and add no significant additional predictive power to the model. Second, the VIF for all of the coefficients is quite high, as was predicted.

Table 1: Table of the full model.

	Estimate	Std. Error	t value	Pr(> t)	VIF	Significance
(Intercept)	1.7939	0.2872	6.2465	0		***
SeedWidth	-0.0902	0.0294	-3.0629	0.0024	231.7271	**
SeedLength	0.2323	0.1394	1.6664	0.0965	1672.8695	.
SeedArea	-0.0012	0.0021	-0.5918	0.5544	918.6289	
SeedAngle	-0.2099	0.4397	-0.4774	0.6334	291.9196	
SeedHeight	-0.2336	0.1536	-1.5214	0.129	2709.2157	
SeedVol	4e-04	2e-04	2.2275	0.0265	124.7234	*

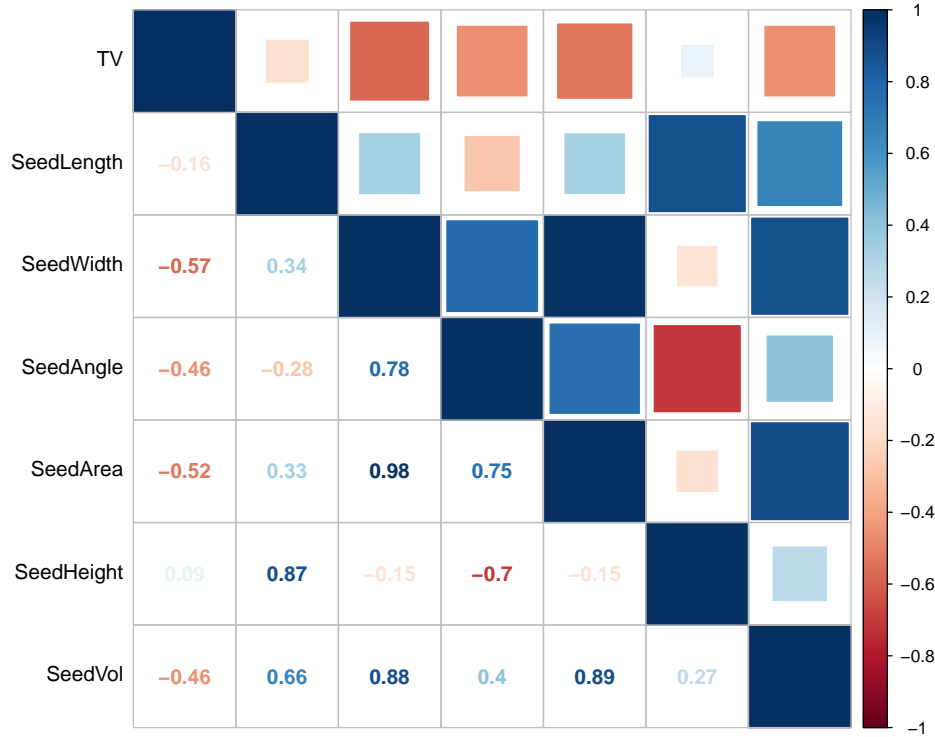


Figure 2: Correlations between variables considered in this analysis.

Given that some of these variables must be removed to improve the fit of the model, we now perform variable selection on the full model to reduce clutter. The first method of selection is backwards selection starting from the full model above, which involves removing subsequent variables based on their p -value; the variable with the highest p -value is removed, the resulting model re-examined, and the process repeated until all predictors were significant at $\alpha = 0.05$. The second method of selection involves backwards selection from the full model using AIC, where terms are removed until the AIC of the model is minimised. The third method of selection starts from the full model and uses the `step` selection function in both directions, while the fourth method starts from the null model and added terms; again, in both cases, the selection continues until the AIC of the model is minimised. The results from the model selection can be found in Table ??, along with the VIF for each term as well as both the AIC and R^2 for each resulting model. All coefficients are significant to $\alpha < 0.001$.

Table 2: Table of the full model.

	Method 1	Method 2	Method 3	Method 4	Method 1,2,3 VIF	Method 4 VIF
(Intercept)	1.6699	1.6699	1.6699	1.6695		
SeedWidth	-0.1027	-0.1027	-0.1027	-0.0865	37.3816	29.508
SeedLength	0.1536	0.1536	0.1536		69.9871	
SeedArea				-0.1471		29.508
SeedAngle						
SeedHeight	-0.1471	-0.1471	-0.1471		72.0249	
SeedVol	0.0003	0.0003	0.0003		14.8502	
AIC	-1293.6000	-1293.6000	-1293.6000	-1288.4000		
R^2	0.3921	0.3921	0.3921	0.3803		

As we can see in the table, the first three selection methods produce the exact same result (which will henceforth be referred to as Model 1), while the fourth method produces a result with only two terms (henceforth Model 2). While Model 1 has a higher R^2 and lower AIC than Model 2, it suffers greatly from high VIF. Because the VIF on Model 2 is much lower than that of Model 1 and the difference in R^2 is very small, we will proceed with Model 2, effectively trading a small increase in the amount of unexplained variance for a large decrease in multicollinearity.

However, even in Model 2, we still see a rather high VIF for the **SeedWidth** and **SeedArea** terms. While the

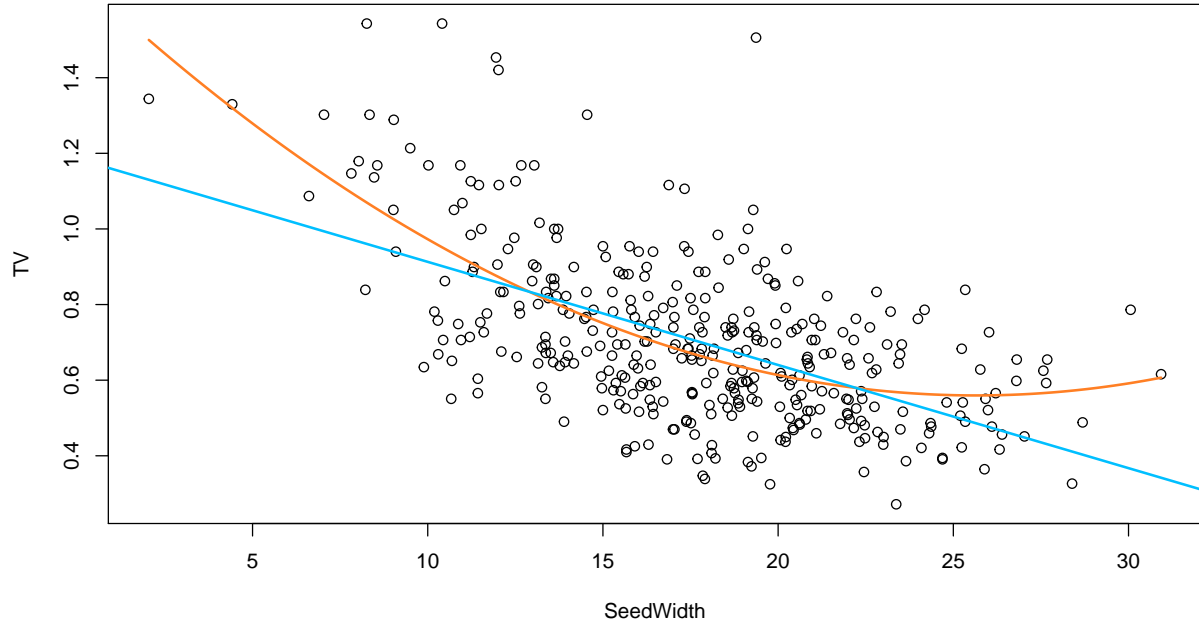


Figure 3: Plot of seed terminal velocity as a function of seed width. The two models are also plotted: the quadratic model (orange) and the linear model (blue).

Research Question 2

Not only are we interested in predicting seed terminal velocity based on the physical properties of the seed, but we are also interested in quantifying the extent to which two environmental variables, warming and mowing, affect seed terminal velocity. Figure 7 shows variation in terminal velocity across all six treatments. Using this plot, we can generate a preliminary hypothesis that warming increases terminal velocity, while mowing decreases it. The following analyses will determine whether such differences are statistically significant. Figure 7 also shows that we have less data for late mowed plants, as many of them died.

Because we are fitting a continuous response variable to categorical predictor variables, we will use an ANOVA model to characterize changes in terminal velocity by treatment. We log-transformed terminal velocity to compensate for skewed right distribution of the terminal velocity data and significant deviations from error normality, as seen in Figure 8.

We used two methods to identify the most appropriate model fit for the log-transformed terminal velocity data. First, we fit the following four models:

- ‘Mowing Only’: $TerminalVelocity = \beta_0 + \beta_1 Mow$
- ‘Warming Only’: $TerminalVelocity = \beta_0 + \beta_1 Warm$
- ‘Warming and Mowing’: $TerminalVelocity = \beta_0 + \beta_1 Mow + \beta_2 Warm$
- ‘Interaction’: $TerminalVelocity = \beta_0 + \beta_1 Mow + \beta_2 Warm + \beta_3 Warm \cdot Mow$

We used model p-values for the **Mowing Only** and **Warming Only** models to establish which provided a better fit for the data. Then, with the better fitting model as a base, we performed a partial F-test to identify whether additional variables significantly improved model fit. Second, to validate the results of the partial F-test, we evaluated both the AIC and BIC for each model.

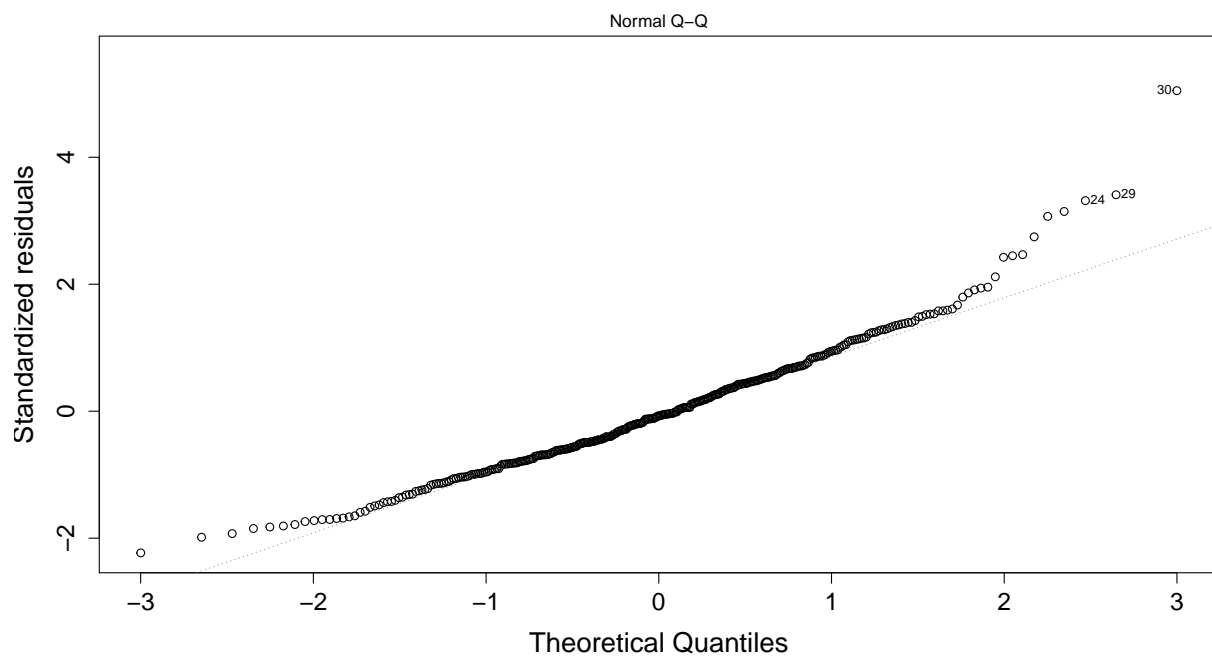
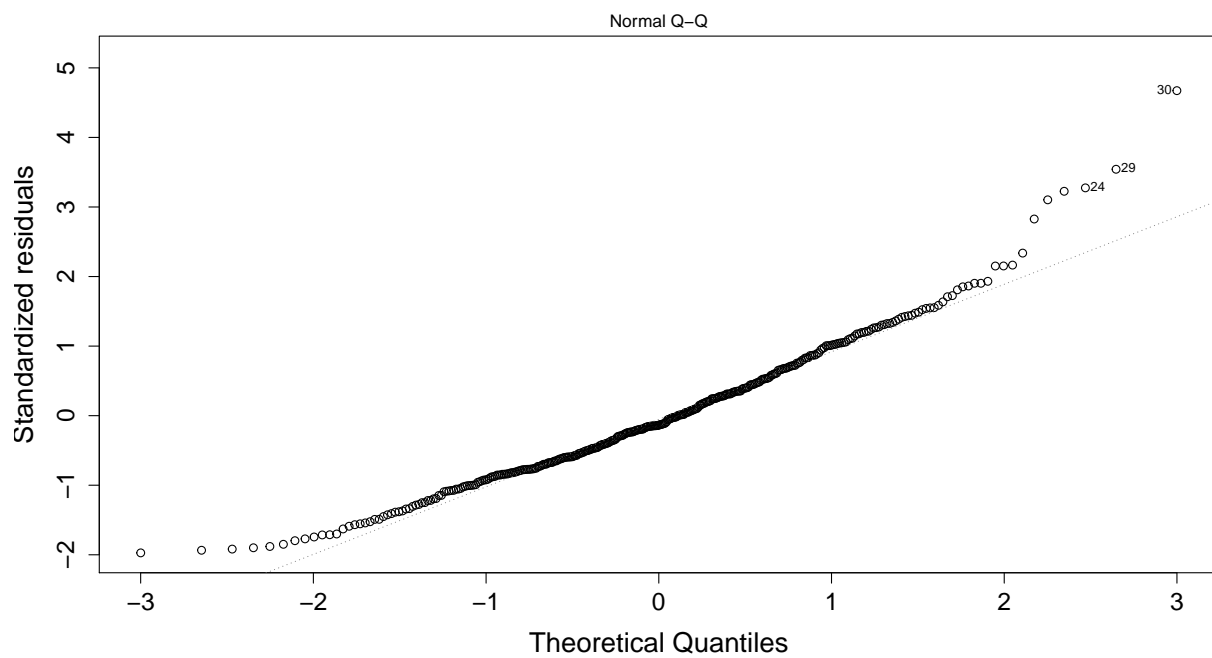


Figure 4: Q-Q Plots comparing normality of the residuals in the linear model (left) and quadratic model (right).

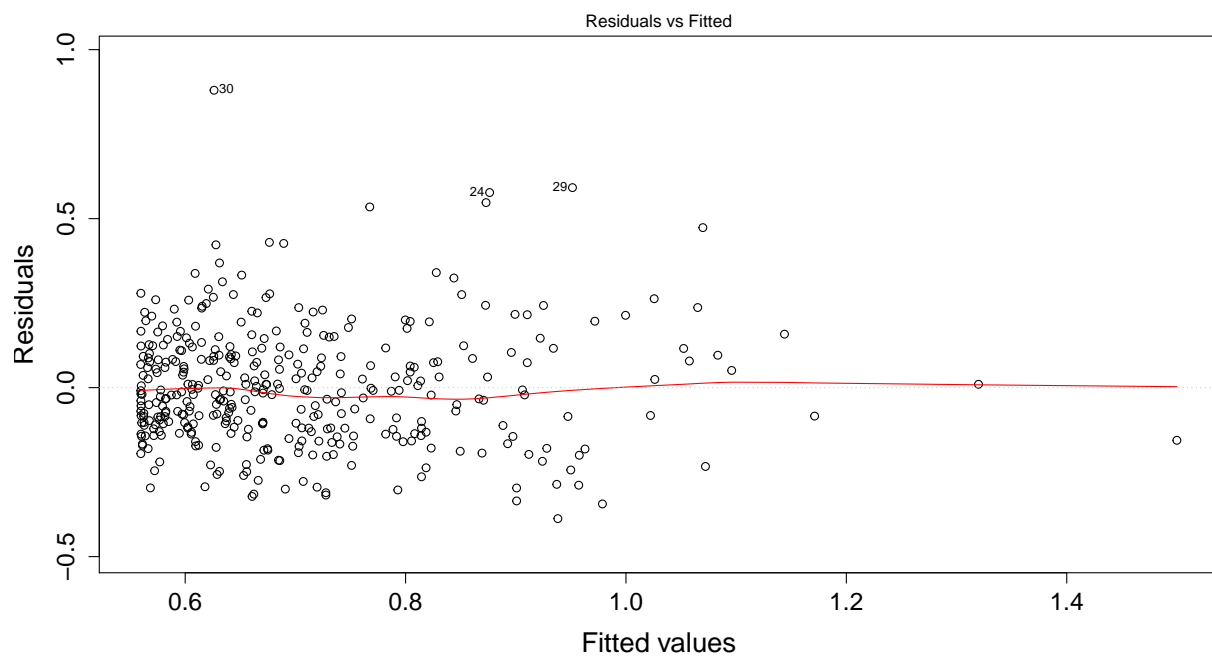
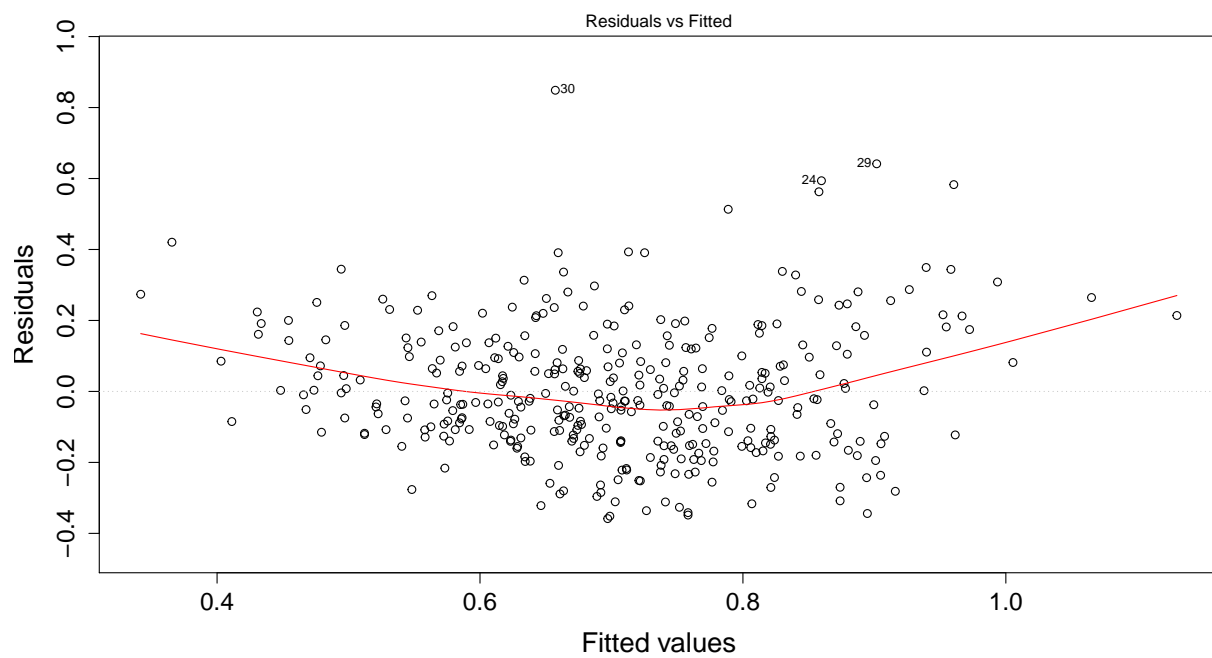


Figure 5: Comparison of residual distributions as a function of fitted values in the linear model (left) and quadratic model (right).

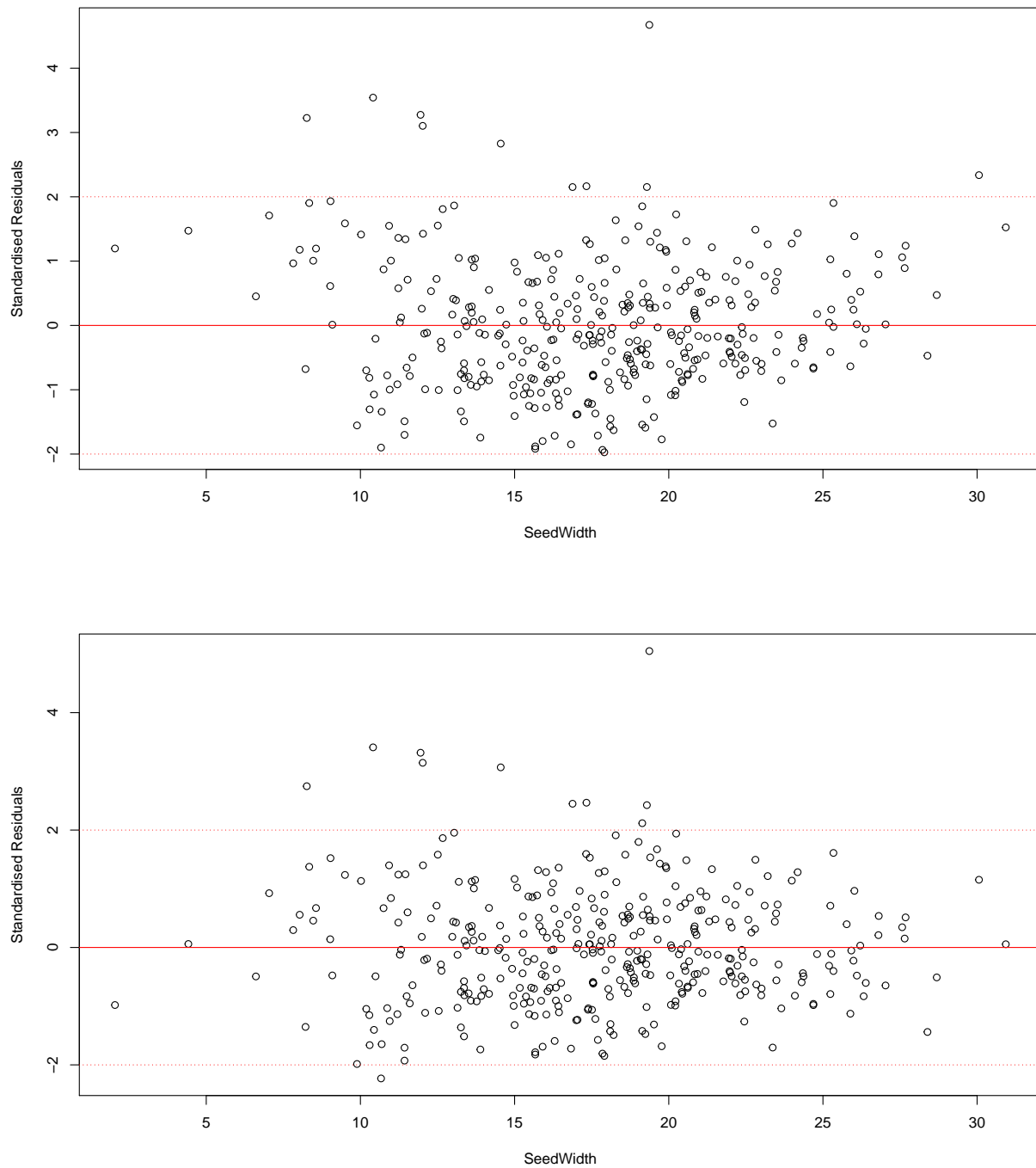


Figure 6: Comparison of standardised residual distributions as a function of observed values in the linear model (left) and quadratic model (right).

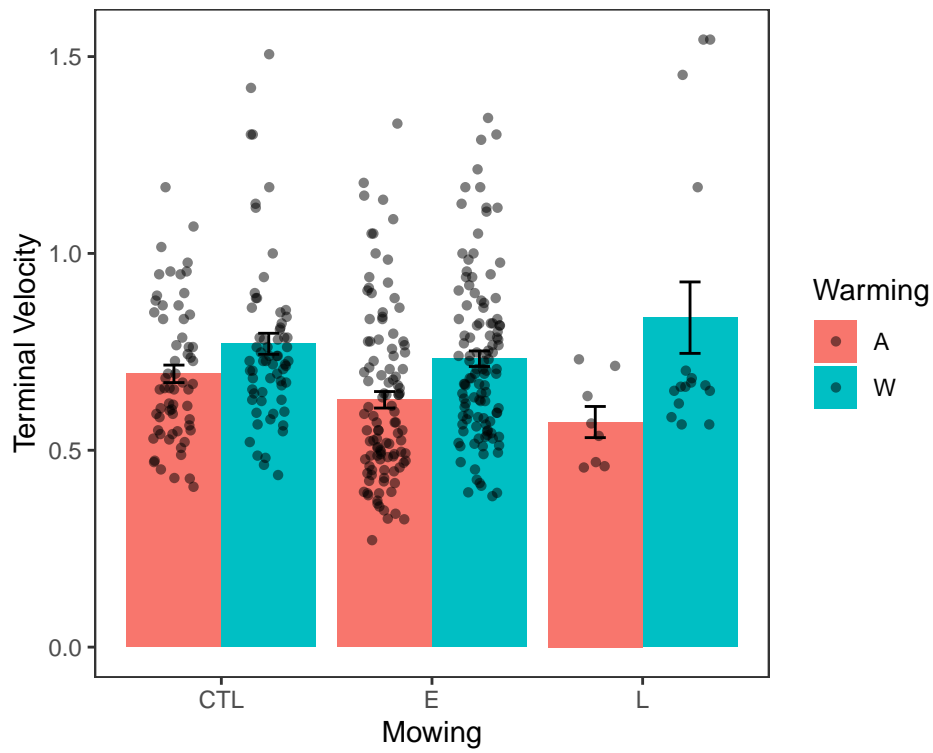


Figure 7: Terminal velocity for plants grown under warming and mowing treatments. Height of bar shows group mean, and error bars show one standard error. Actual data shown with black dots.

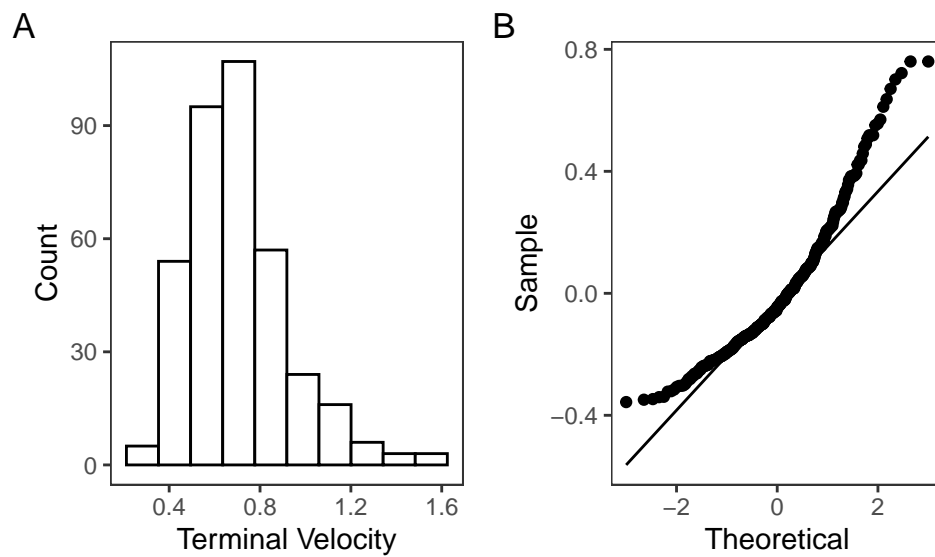


Figure 8: Justification for transformation of terminal velocity. Histogram of terminal velocity (A) and Normal Q-Q plot of linear model with both warming and mowing predictors (B).

Here, the **Warming Only** model has a p-value of 0 whereas the **Mowing Only** model has a p-value of 0.0248. Therefore, we use the **Warming Only** model as the base model in our partial F-test. Results, shown in Table 3, suggest that the **Warming and Mowing** model is the best fit for the data.

Table 3: Results of partial F-test comparing three models for predicting terminal velocity.

Model	P.Value
Warming Only	
Warming and Mowing	0.0217
Interaction	0.2134

Like the partial F-test, both AIC and BIC support the **Warming and Mowing** model (Table 4). Thus, we conclude that $TerminalVelocity = \beta_0 + \beta_1 Mowing + \beta_2 Warming$ is the best model fit for the log-transformed terminal velocity data.

Table 4: AIC and BIC values for each model.

Model	AIC	BIC
Mowing Only	162.1546	169.1052
Warming Only	162.1546	169.1100
Warming and Mowing	138.4409	147.1292
Interaction	139.3011	151.4648

The residuals for the **Warming and Mowing** model are shown in Figure 9. The residuals show approximately random distribution, and though there are a few points with high leverage, they do are not high includence accrodg to Cook’s distance. Lastly, we can see that our log-transformation significantly improved the normality of the errors, as shown in the Normal Q-Q plot.

Model results are shown in Table 5 and treatment level predictions are shown in Figure 10. From these results, we conclude that early mowing significnatly decreases terminal velocity, whereas warming increases terminal velocity.

Table 5: Estimated coefficients and p-values for **Warming and Mowing** model. 95% confidence interval shown in parenthesis.

	Estimate	Pr(> t)
(Intercept)	-0.42 (-0.48, -0.36)	0
MowE	-0.09 (-0.15, -0.03)	0.01
MowL	-0.04 (-0.17, 0.09)	0.56
WarmingW	0.16 (0.1, 0.21)	0

Research Question 3

Next, we will explore the extent to which the results of Research Question 2, or the changes in terminal velocity by treatment, can be explained by changes in seed width. We are investigating seed width specifically because the results of Research Question 1 show that seed width is a good predictor of terminal velocity. Figure 11 shows the variation in seed width across treatments.

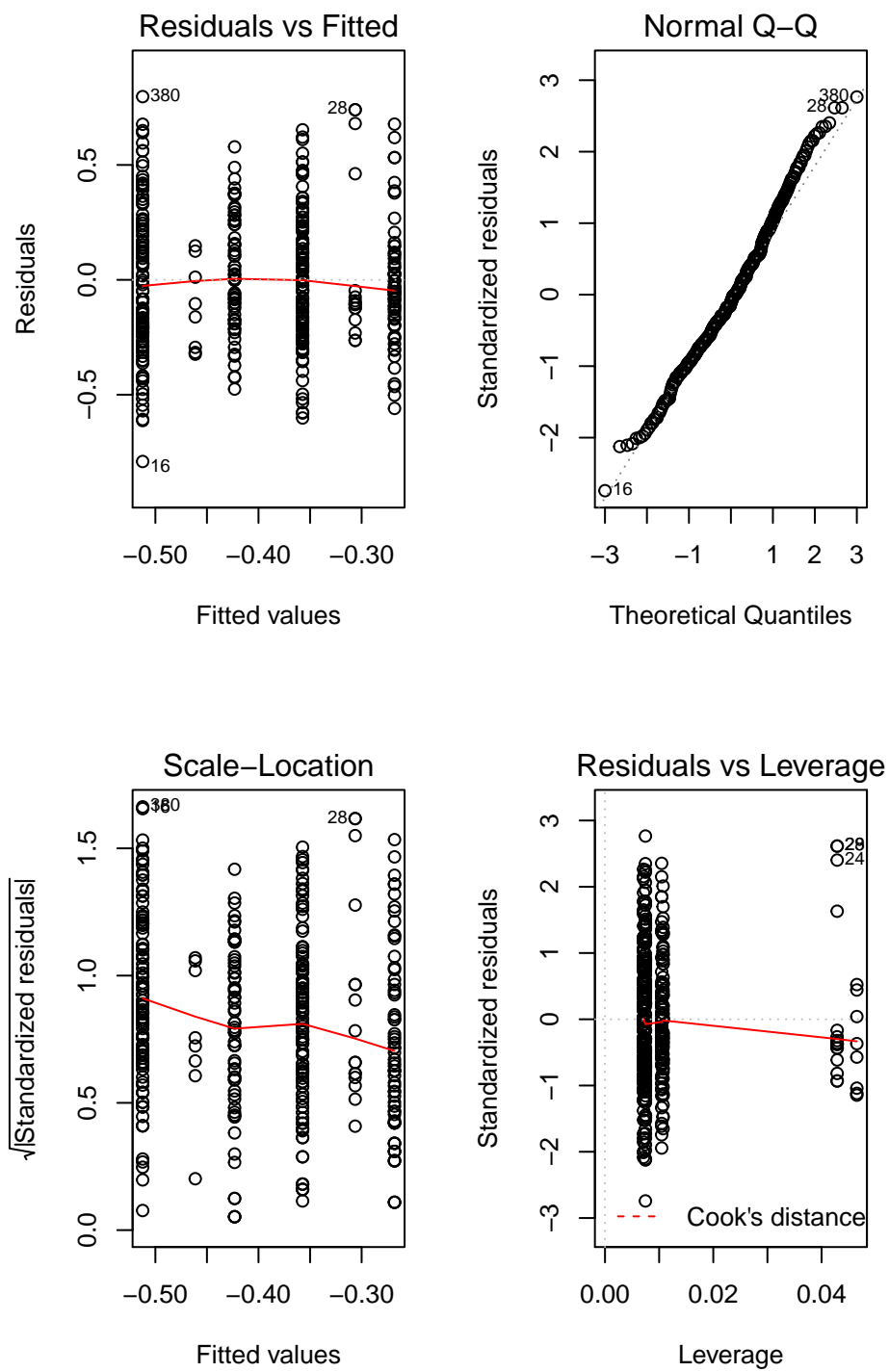


Figure 9: Diagnostic plots for Warming and Mowing model.

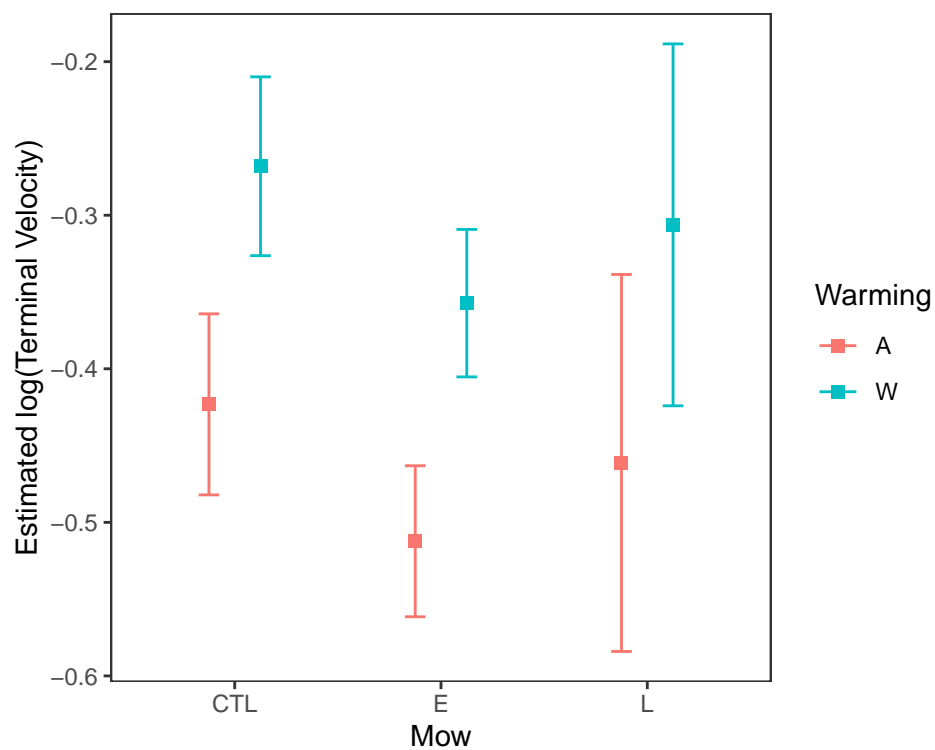


Figure 10: Results of best fitting ANOVA model, which included both warming and mowing as predictors. Square dots show mean predicted value and error bars show 95% confidence intervals.

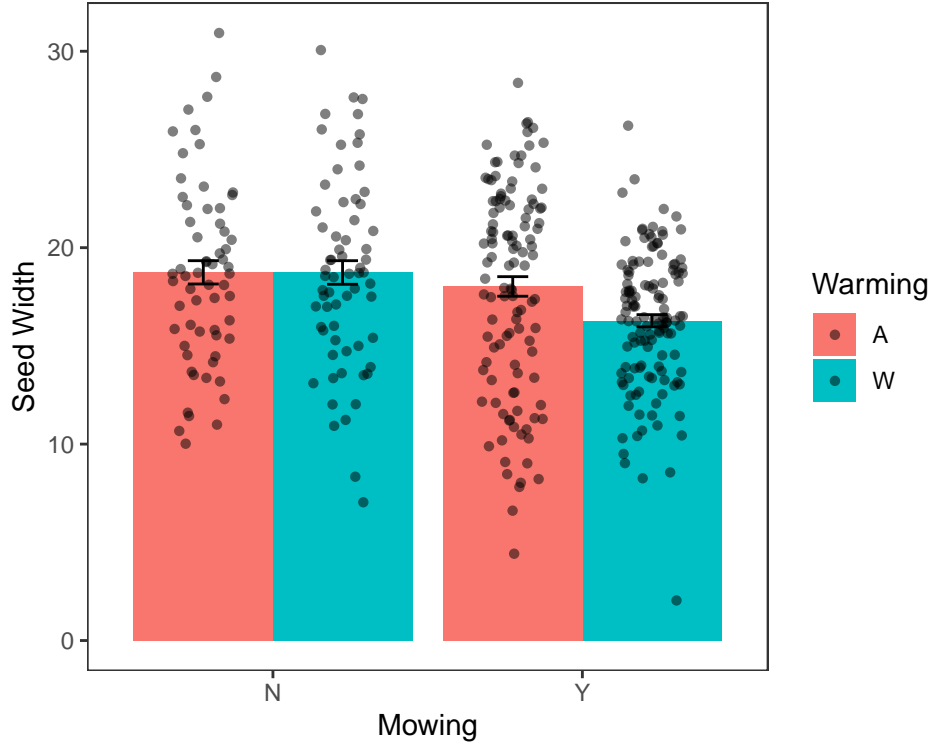


Figure 11: Seed width by warming and mowing treatments. Height of bar shows group mean, and error bars show one standard error. Actual data shown with black dots.

Our data fitting methods are similar to those of Research Question 2. We use partial F-tests and AIC/BIC analyses for model selection, but did not transform our response variable. First, we analyze the fit of the four basic models - mowing only, warming only, warming and mowing, and interaction - to identify which model is the best fit.

Here, the **Warming Only** model has a p-value of 0.0152 whereas the **Mowing Only** model has a p-value of 0.0031. Therefore, we use the **Mowing Only** model as the base model in our partial F-test. Results suggest that the **Warming and Mowing** model is again the best fit for the data (Table 6).

Table 6: Results of partial F-test for models predicting seed width by treatment.

Model	P.Value
Mowing Only	
Warming and Mowing	0.0198
Interaction	0.2272

Again, we use AIC and BIC model selection criteria to validate these results. Table 7 shows similar results to those in Research Question 2, suggesting that the **Warming and Mowing** model is the best fit.

Table 7: AIC and BIC values for each of the four models predicting seed width.

Model	AIC	BIC
Mowing Only	2185.078	2192.028
Warming Only	2188.787	2194.000
Warming and Mowing	2181.592	2190.280
Interaction	2182.579	2194.743

Thus, both the partial F-test and the AIC/BIC values support the **Warming and Mowing** model, whose model diagnostic plots are shown in Figure 12. These residuals suggest the errors are approximately normally distributed. Again, there may be a few leverage points, though they do not have high influence.

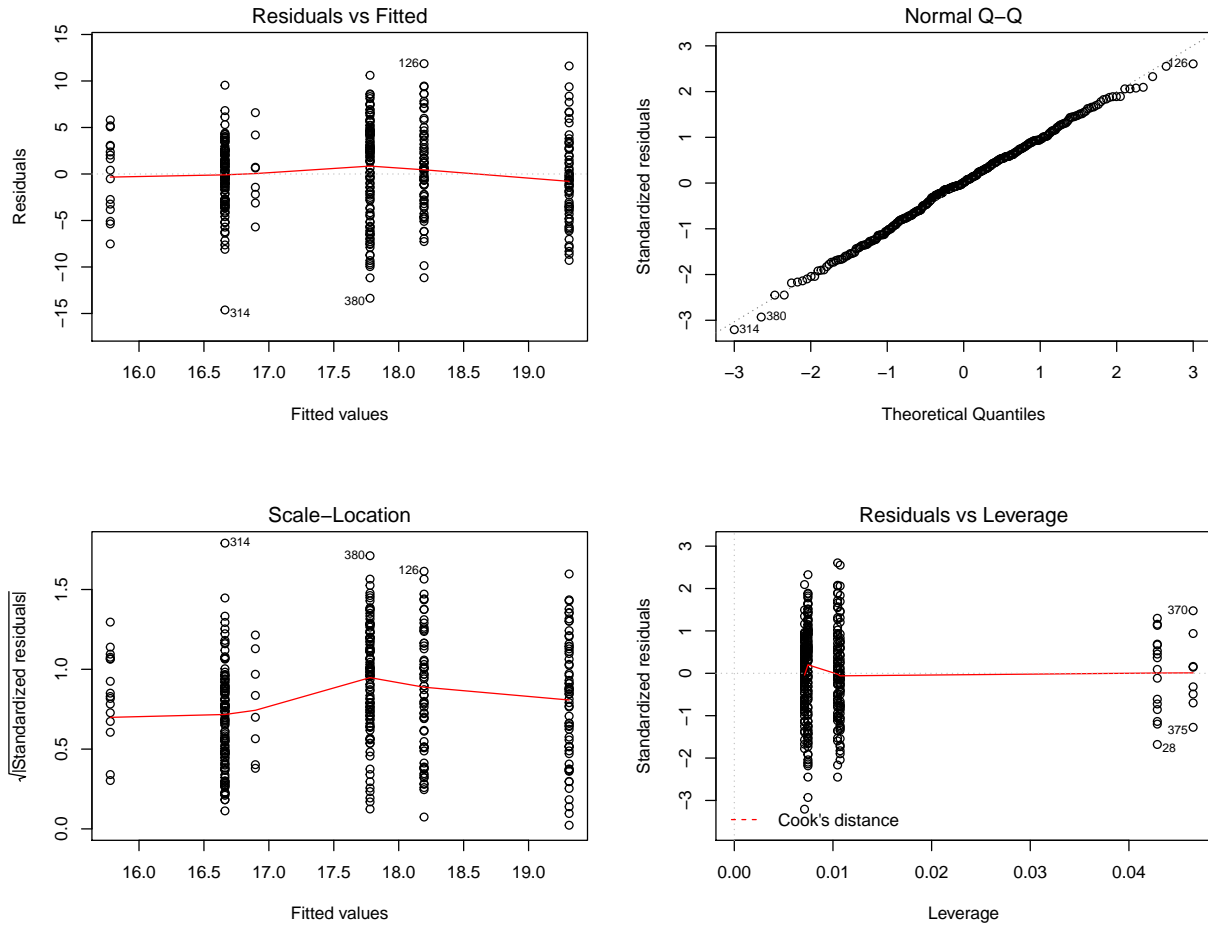


Figure 12: Diagnostic plots for predicting seed width using Warming and Mowing model.

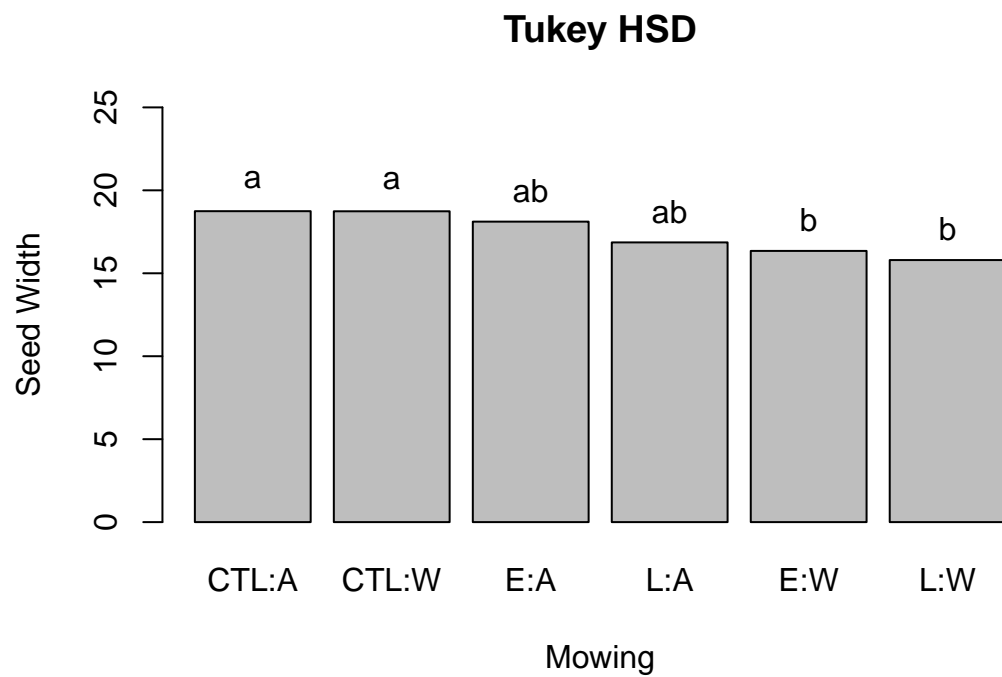


Figure 13: Results of post-hoc Tukey HSD test. Letters indicate groups which are significantly different, where 0.05 is the significance level. Notation is as follows: CTL - no mowing, E - early mowing, L - late mowing; A - ambient (i.e. no warming), W - warming

Table 9: Results of partial F-test performed for models predicting seed width using warming and binary mowing predictors.

Model	P.Value
Warming Only	
Warming and Mowing	0.0161
Interaction	0.0828

Table 10: AIC and BIC for models predicting seed width using warming and binary mowing predictors.

Model	AIC	BIC
Mowing Only	2184.213	2189.426
Warming Only	2188.787	2194.000
Warming and Mowing	2180.400	2187.351
Interaction	2179.354	2188.042

Discerning between the **Warming and Mowing** and the **Interaction** models is less clear in this case. On one hand, the partial F-test suggests the **Interaction** model may provide a slightly better fit (Table 9) and the **Interaction** model has the lowest AIC value (Table 10), but the **Warming and Mowing** model has the lowest BIC value (Table 10). This result is consistent with our expectations, as a slight improvement in model fit may not be enough to overcome the BIC penalty for an additional term in the **Interaction** model.

The results of both models are summarized in Table 11. In the **Warming and Mowing** model, both treatments decrease seed width; whereas when an interaction term is added, neither the warming nor mowing terms are significant, but the interaction term itself becomes significant at a significance level of 0.1.

Table 11: Results of **Warming and Mowing** and **Interaction** models, respectively.

	Warming and Mowing Estimate	Pr(> t)	Interaction Estimate	Pr(> t)
Intercept	19.3284 (18.4, 20.26)	0	18.7442 (17.6, 19.88)	0
Mow	-1.6198 (-2.61, -0.63)	0.0013	-0.718 (-2.13, 0.7)	0.3194
Warm	-1.1485 (-2.09, -0.21)	0.0164	-0.0071 (-1.6, 1.59)	0.993
Mow:Warm			-1.7398 (-3.71, 0.23)	0.0828

These two results suggest that plants which are both warmed and mowed have smaller seed width. Plants which are both warmed and mowed are likely driving the significant mowing and warming coefficients we see in the **Warming and Mowing** model. Another post-hoc Tukey test, shown in Figure 14, supports this conclusion.

Discussion

The results of this experiment contribute to our understanding of how *Carduus nutans* may spread under future climates, and further how mowing management may mediate these changes. We found that, as predicted by physical models⁴, the width of the seed is a good predictor of terminal velocity. Smaller seeds fall faster, and thus disperse shorter distances.

Theoretical⁵ and experimental⁶ results suggest that plants under stress will disperse farther, and our results are consistent with this idea. We showed that early mowing decreased terminal velocity, which means, with

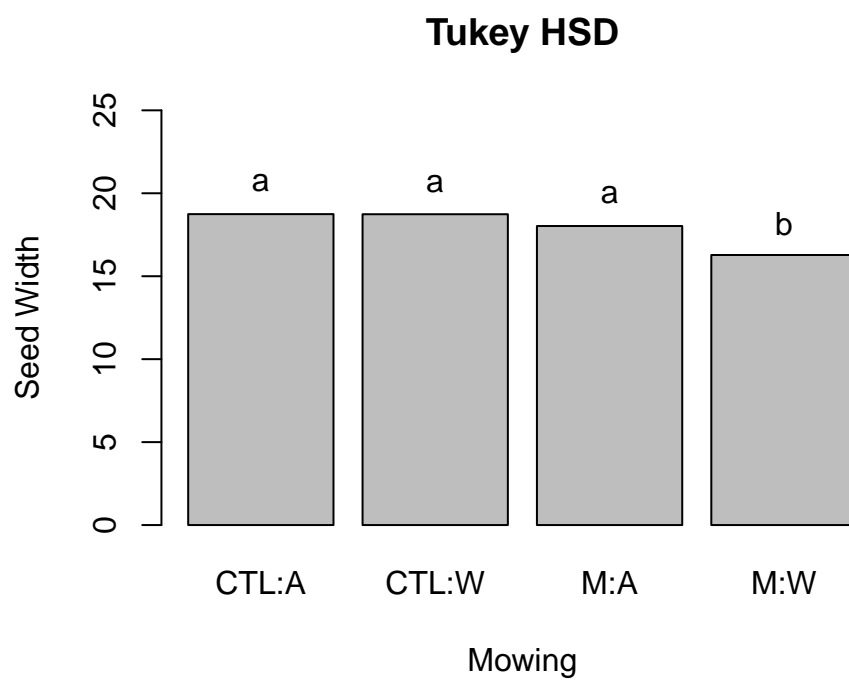


Figure 14: Results of post-hoc Tukey HSD test. Letters indicate groups which are significantly different, where 0.05 is the significance level. Notation is as follows: CTL - no mowing, M - mowing; A - ambient (i.e. no warming), W - warming

all else equal, these mowed plants will disperse further. Understanding the effect of late mowing on seed terminal velocity was difficult because many of the late mowed plants died. Fewer data points paired with small effect sizes for late mowed plant terminal velocity created wide confidence intervals and high p-values. Quantifying such terminal velocity changes in late-mowed plants may require a study with more replication to ensure sufficient statistical power after plant mortality due to late mowing.

We also found that warming increased terminal velocity. In this case, all else being equal, we would expect seeds to disperse less far. However, previous work in our lab⁷ showed that warmed thistles reach taller heights. Given a fixed terminal velocity, seeds of taller plants are expected to disperse further. Thus, an increase in terminal velocity of warmed plants does not necessarily translate directly into less dispersal and spread. Further modeling, that includes changes in plant height as well as changes in terminal velocity, will quantify overall changes in thistle spread.

We showed that plants which were both warmed and mowed had smaller seed width. Based on our model which describes terminal velocity as a function of seed width (Research Question 1), we expect these warmed and mowed plants to have higher terminal velocity and, therefore, to disperse less far. Yet, in our analysis of changes in terminal velocity by treatment (Research Question 2), we showed mowing decreased terminal velocity. These discrepancies suggest that treatment level changes in seed terminal velocity cannot be fully explained by changes in seed width.

This data set is lacking seed mass data, which may be responsible for the inconsistencies we observed. Seed mass is a second physical property needed to predict terminal velocity⁴ and may also change with warming and mowing. Lacking seed mass data limits our ability to explain changes in seed terminal velocity through physical properties of the seed. However, our results which quantify the effect of the treatments on terminal velocity are unaffected. Further studies which measure both seed width and seed mass will be necessary to explain physical and biological mechanisms which explain the observed changes in terminal velocity.

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