

Mysis Preservation – Length-Weight Analysis for Manuscript

January 5, 2008

1 Length Analysis

1.1 Initialization

1.1.1 Data entry and cleaning

```
> mysis <- read.xls("Mysis_Data_Master.xls", sheet = "Mysis", colClasses = c("logical",  
+   "logical", "factor", "logical", "logical", "factor", "factor", "numeric",  
+   "factor", "factor", "isodate", "isotime", "factor", "factor", "factor",  
+   "numeric", "numeric", "numeric", "numeric", "numeric", "numeric",  
+   "numeric", "numeric", "factor"))  
> mysisL <- mysis[mysis$include.L, ]  
> mysis.len <- mysisL[mysisL$use.l, ]  
> mysis.len$tx <- factor(mysis.len$tx)  
> mysis.len$stage <- factor(mysis.len$stage)  
> mysis.len$group <- mysis.len$stage:mysis.len$tx  
> mysis.len$loss.p2w <- mysis.len$len.p2w - mysis.len$len.fr  
> mysis.len$loss.p2m <- mysis.len$len.p2m - mysis.len$len.fr  
> mysis.len$ploss.p2w <- mysis.len$loss.p2w/mysis.len$len.fr  
> mysis.len$ploss.p2m <- mysis.len$loss.p2m/mysis.len$len.fr
```

Getting rid of juveniles – decided this from length-weight work – see below.

```
> mysis.len2 <- subset(mysis.len, stage != "Juv")  
> mysis.len2$stage <- factor(mysis.len2$stage)  
> mysis.len2$group <- factor(mysis.len2$group)
```

1.2 Sample Summaries

These results form Table R1 in the manuscript.

```
> attach(mysis.len2)  
> ftable(xtabs(~tx + stage))
```

	stage	Fem	Male
tx			
8BF		24	18
8SBF		15	8

```
> tapply(len.fr, stage:tx, Summary, na.rm = T)
```

```
$`Fem:8BF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
9.200000	10.000000	11.300000	12.200000	14.400000	16.000000	2.458964

```
$`Fem:8SBF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
8.300000	9.200000	9.600000	10.290000	10.400000	16.100000	1.974287

```
$`Male:8BF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
10.400000	12.830000	13.150000	13.110000	14.170000	14.500000	1.277815

```
$`Male:8SBF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
12.200000	13.020000	13.550000	13.310000	13.700000	13.900000	0.5792544

```
> tapply(loss.p2w, stage:tx, Summary, na.rm = T)
```

```
$`Fem:8BF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
-0.6000000	-0.3250000	-0.2000000	-0.2125000	-0.1000000	0.2000000	0.1872281

```
$`Fem:8SBF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
-0.3000000	-0.2000000	-0.1000000	-0.1000000	0.0000000	0.1000000	0.1362770

```
$`Male:8BF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
-0.3000000	-0.2000000	-0.1000000	-0.1000000	0.0000000	0.1000000	0.1283378

```
$`Male:8SBF`
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	StDev
-0.5000000	-0.2000000	0.0000000	-0.1000000	0.0250000	0.2000000	0.2618615

1.3 Results on Length After 2 Weeks Preservation

Most (63.1%) Mysids were shorter after two weeks of , while a small number (10.8%) actually appeared to be longer after preservation.

```
> p <- prop.table(table(round(loss.p2w, 1)))
```

```
> p
```

-0.6	-0.5	-0.4	-0.3	-0.2	-0.1	0	0.1
0.01538462	0.04615385	0.06153846	0.13846154	0.20000000	0.16923077	0.26153846	0.07692308
0.2	0.03076923						

```
> sum(p[1:6])
```

```
[1] 0.6307692
```

```
> sum(p[8:9])
```

```
[1] 0.1076923
```

The change in standard length after two weeks of preservation was not significantly related to the fresh standard length measurement ($p=0.2001$) nor did it differ between male and female Mysids ($p=0.1732$). There was weak but insignificant evidence ($p=0.0638$) that Mysids preserved in 8% buffered formalin lost more length than Mysids preserved in 8% sugar-buffered formalin.

```
> lm2 <- lm(loss.p2w ~ len.fr * stage * tx)
> Anova(lm2)
```

Anova Table (Type II tests)

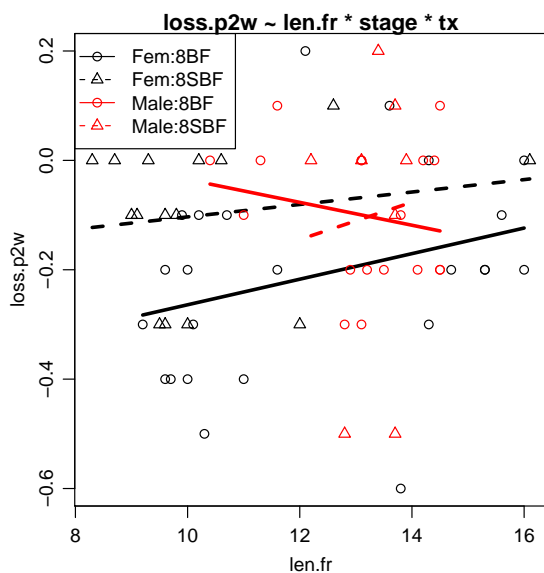
Response: loss.p2w

	Sum Sq	Df	F value	Pr(>F)
len.fr	0.05080	1	1.6752	0.20078
stage	0.05768	1	1.9023	0.17321
tx	0.10837	1	3.5740	0.06378
len.fr:stage	0.03731	1	1.2305	0.27196
len.fr:tx	0.00300	1	0.0988	0.75445
stage:tx	0.03503	1	1.1551	0.28701
len.fr:stage:tx	0.00920	1	0.3036	0.58381
Residuals	1.72835	57		

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

The following plot is NOT in the manuscript.

```
> fit.plot(lm2, legend = "topleft")
```



Thus, regardless of fresh standard length, life stage or preservation type, Mysids lost an average of 0.14 mm (95% CI: 0.10,0.19).

```
> t.test(loss.p2w)

One Sample t-test

data:  loss.p2w
t = -6.4265, df = 64, p-value = 1.865e-08
alternative hypothesis: true mean is not equal to 0
95 percent confidence interval:
 -0.18553658 -0.09754035
sample estimates:
mean of x
-0.1415385

> detach(mysis.len2)
```

2 Length-Weight Analysis

2.1 Initialization

2.1.1 Data entry and cleaning

```
> mysis <- read.xls("Mysis_Data_Master.xls", sheet = "Mysis")
> mysis$c.wt <- mysis$c.wt * 1000
> mysis1 <- mysis[mysis$include.LW, ]
> mysis2 <- mysis1[mysis1$use.lw == TRUE, ]
> mysis2$logwt <- log10(mysis2$c.wt)
> mysis2$loglen <- log10(mysis2$len)
> mysis2$treat <- mysis2$tx:mysis2$stage
```

mysis2 is now the cleaned and ready data frame.

2.1.2 declare some labels

```
> xlbl1 <- "Standard Length (mm)"
> xlbl2 <- "log(Standard Length (mm))"
> ylbl1 <- "Dry Weight (mg)"
> ylbl2 <- "log(Dry Weight (mg))"
```

2.2 EDA

2.2.1 Original data

None of this section is in the manuscript. Note (1) that gravid-8BF are missing and (2) increased variance at small lengths on log-log scale. This led to the exclusion of juveniles – problem with weights of small specimens.

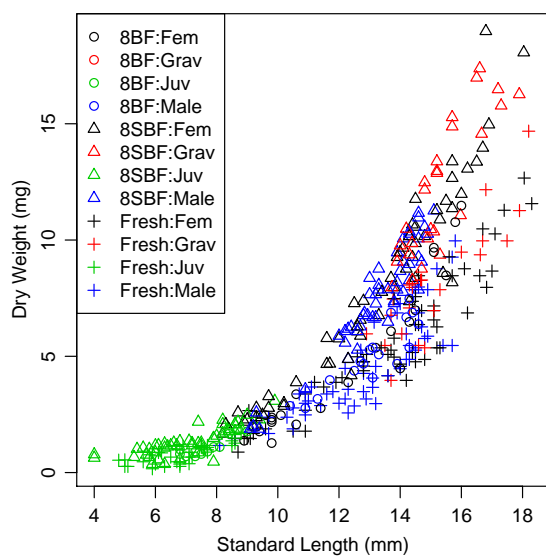
```
> attach(mysis2)
> ftable(xtabs(~tx + stage + len.bin))
```

		len.bin	3	5	7	9	11	13	15	17
tx	stage									
8BF	Fem		0	0	1	11	2	5	5	0
	Grav		0	0	0	0	0	0	0	0
	Juv		0	2	9	1	0	0	0	0
	Male		0	0	0	2	5	11	0	0
8SBF	Fem		0	0	4	10	12	19	14	1
	Grav		0	0	0	0	0	12	12	3
	Juv		2	20	23	3	0	0	0	0
	Male		0	0	0	5	12	32	1	0
Fresh	Fem		0	0	3	11	10	19	13	5
	Grav		0	0	0	0	1	12	6	3
	Juv		1	19	16	0	0	0	0	0
	Male		0	0	2	9	15	23	5	0

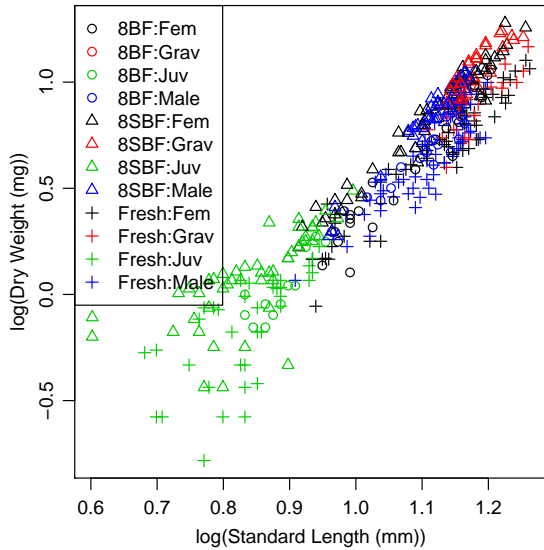
```
> ftable(xtabs(~tx + stage))
```

	stage	Fem	Grav	Juv	Male
tx					
8BF		24	0	12	18
8SBF		60	27	48	50
Fresh		61	22	37	54

```
> plot(c.wt ~ len, pch = as.numeric(tx), col = as.numeric(stage), xlab = xlbl1,
+      ylab = ylbl1)
> legend(x = "topleft", legend = levels(tx:stage), col = rep(seq(1, 4),
+      3), pch = rep(seq(1, 4), each = 4))
```



```
> plot(logwt ~ loglen, pch = as.numeric(tx), col = as.numeric(stage),
+       xlab = xlbl2, ylab = ylbl2)
> legend(x = "topleft", legend = levels(tx:stage), col = rep(seq(1, 4),
+       3), pch = rep(seq(1, 4), each = 4))
> detach(mysis2)
```



2.2.2 Permanently Remove Juveniles

```
> mysis3 <- mysis2[mysi2$stage != "Juv", ]
> mysis3$stage <- factor(mysis3$stage)
> mysis3$treat <- factor(mysis3$treat)
> attach(mysis3)
```

mysi3 is now the cleaned and ready data frame

These results are not in the manuscript – just double-checking that juveniles have been removed.

```
> ftable(xtabs(~tx + stage + len.bin))
```

		len.bin	7	9	11	13	15	17
tx	stage							
8BF	Fem		1	11	2	5	5	0
	Grav		0	0	0	0	0	0
	Male		0	2	5	11	0	0
8SBF	Fem		4	10	12	19	14	1
	Grav		0	0	0	12	12	3
	Male		0	5	12	32	1	0
Fresh	Fem		3	11	10	19	13	5
	Grav		0	0	1	12	6	3
	Male		2	9	15	23	5	0

The following results form Table R2 in the manuscript.

```
> ftable(xtabs(~tx + stage))
```

	stage	Fem	Grav	Male
tx				
8BF		24	0	18
8SBF		60	27	50
Fresh		61	22	54

```
> tapply(len, stage:tx, Summary, na.rm = T)
```

```
$`Fem:8BF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
8.900000 9.800000 11.000000 11.990000 14.350000 16.000000 2.522752

$`Fem:8SBF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
8.200000 11.600000 13.850000 13.150000 14.900000 18.040000 2.541817

$`Fem:Fresh`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
8.600000 11.500000 13.800000 13.420000 15.100000 18.300000 2.611002

$`Grav:8BF`
NULL

$`Grav:8SBF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
13.730000 14.390000 15.100000 15.280000 15.840000 17.890000 1.125171

$`Grav:Fresh`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
12.900000 14.110000 14.680000 15.160000 16.460000 18.200000 1.517133

$`Male:8BF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
10.400000 12.550000 13.100000 13.010000 14.120000 14.600000 1.257396

$`Male:8SBF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
9.100000 12.620000 13.300000 13.180000 14.250000 15.100000 1.473647

$`Male:Fresh`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
8.100000 11.450000 13.100000 12.870000 14.470000 15.800000 1.890991
```

```
> tapply(c.wt, stage:tx, Summary, na.rm = T)
```

```
$`Fem:8BF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
1.270000 2.145000 3.070000 4.644000 6.605000 11.480000 3.246756

$`Fem:8SBF`
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   StDev
1.665000 4.544000 8.073000 7.877000 10.590000 18.980000 4.141679
```

```

$`Fem:Fresh`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 0.880000  3.880000  5.080000  5.456000  7.165000 12.660000  2.783356

$`Grav:8BF`
NULL

$`Grav:8SBF`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 7.965000  9.615000 11.060000 12.070000 14.720000 17.380000  2.952271

$`Grav:Fresh`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 3.965000  6.165000  7.915000  8.081000  9.451000 14.680000  2.512076

$`Male:8BF`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 2.770000  4.155000  4.930000  5.045000  6.130000  7.480000  1.407255

$`Male:8SBF`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 1.880000  6.269000  7.715000  7.445000  9.040000 11.270000  2.344641

$`Male:Fresh`
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.    StDev
 1.165000  3.005000  4.215000  4.628000  5.840000  9.965000  2.098266

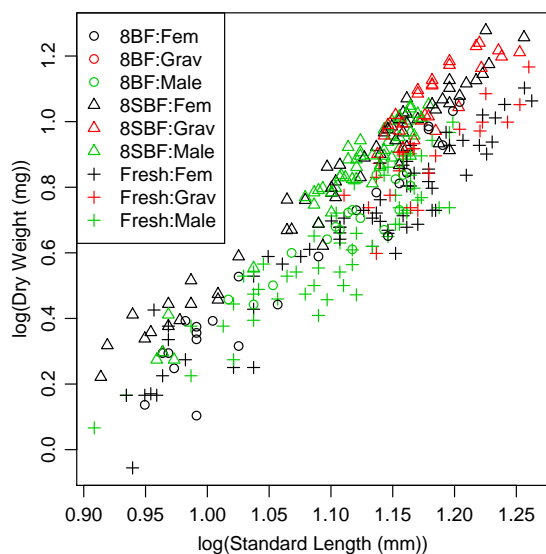
```

The following figure is NOT in the manuscript.

```

> plot(logwt ~ loglen, pch = as.numeric(tx), col = as.numeric(stage),
+      xlab = xlbl2, ylab = ylbl2)
> legend(x = "topleft", legend = levels(tx:stage), col = rep(seq(1, 3),
+      3), pch = rep(seq(1, 3), each = 3))
> detach(mysis3)

```



2.3 First Length-Weight Model Analysis

The first model examined the effect of all three preservation treatments and two life stages (males and females) on the L-W relationship.

2.3.1 Temporarily excludes gravids

None of these results are in the manuscript.

```
> mysis3.nograv <- mysis3[mysis3$stage != "Grav", ]
> mysis3.nograv$stage <- factor(mysis3.nograv$stage)
> mysis3.nograv$treat <- factor(mysis3.nograv$treat)
> attach(mysis3.nograv)
> ftable(xtabs(~tx + stage + len.bin))
```

		len.bin	7	9	11	13	15	17
tx	stage							
8BF	Fem		1	11	2	5	5	0
	Male		0	2	5	11	0	0
8SBF	Fem		4	10	12	19	14	1
	Male		0	5	12	32	1	0
Fresh	Fem		3	11	10	19	13	5
	Male		2	9	15	23	5	0

2.3.2 Fitting The Model

```
> lm1 <- lm(logwt ~ loglen * tx * stage)
```

Assumption Checking

None of these results are in the manuscript. The normality test and non-constant variance test both indicate assumption violations. However, the residual plot shows a general homoscedasticity (with the exception of two minor outliers and a small narrowing at large lengths) and the histogram is largely symmetric.

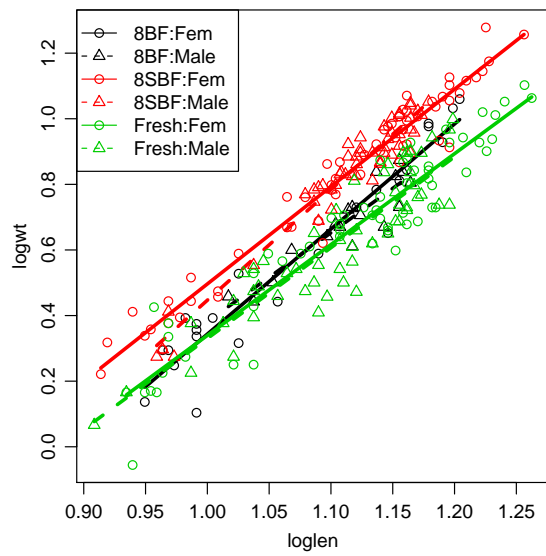
```
> fit.plot(lm1, leg = "topleft", main = "")
> ad.test(lm1$residuals)
```

Anderson-Darling normality test

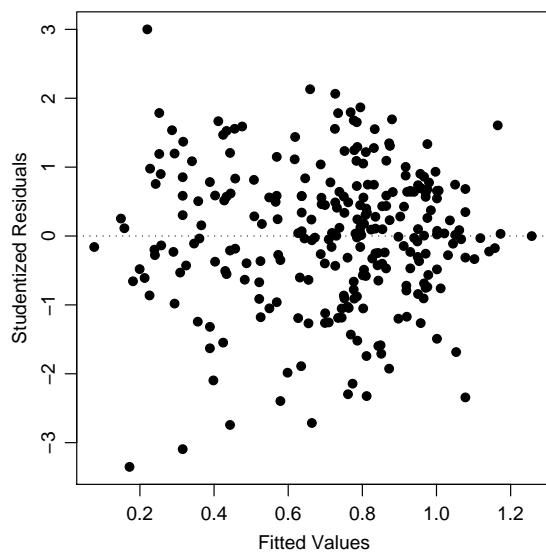
```
data: lm1$residuals
A = 0.8453, p-value = 0.02923
```

```
> ncv.test(lm1)
```

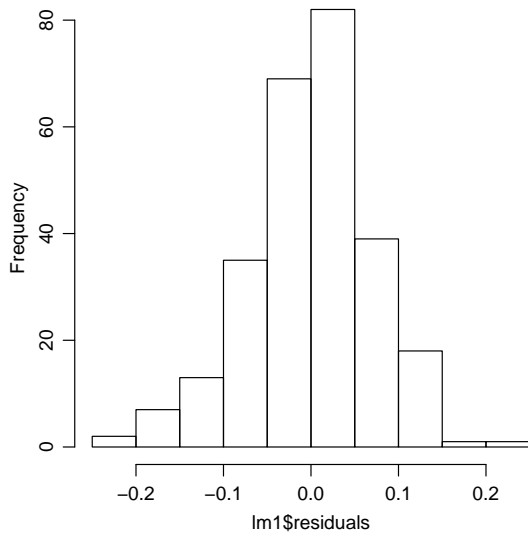
```
Non-constant Variance Score Test
Variance formula: ~ fitted.values
Chisquare = 10.80524    Df = 1    p = 0.00101213
```



```
> residual.plot(lm1, main = "")
```



```
> hist(lm1$residuals, main = "")
```



Results

The first linear model fit to males and females and all three preservation treatments showed that there was a significant difference in slope among the three preservation treatments ($p=0.0213$) but not among the two life-stages ($p=0.4457$).

The mean log(weight) adjusted for log(length) did not differ among male and females ($p=0.4038$).

```
> Anova(lm1)
```

Anova Table (Type II tests)

Response: logwt

	Sum Sq	Df	F value	Pr(>F)
loglen	13.5407	1	2593.5986	< 2e-16 ***
tx	1.9385	2	185.6499	< 2e-16 ***
stage	0.0037	1	0.6992	0.40382
loglen:tx	0.0408	2	3.9089	0.02128 *
loglen:stage	0.0030	1	0.5835	0.44567
tx:stage	0.0010	2	0.0943	0.91005
loglen:tx:stage	0.0215	2	2.0626	0.12923
Residuals	1.3313	255		

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Post hoc multiple comparisons indicated that the 8% buffered formalin and 8% sugar-buffered formalin treatments had statistically similar slopes ($p=0.7974$) that were significantly steeper than the slope of the fresh preserved treatment ($p<0.0798$).

```
> comp.slopes(lm1)
```

Multiple comparison control procedures used: fdr

Multiple Slope Comparisons

	comparison	diff	lwr	upr	raw.p	adj.p
1	8SBF-8BF	-0.04466	-0.38690	0.29758	0.79743	0.79743
2	Fresh-8BF	-0.33003	-0.66463	0.00456	0.05318	0.07977
3	Fresh-8SBF	-0.28538	-0.53023	-0.04053	0.02253	0.06759

Slope Information

	level	slopes	lwr	upr	raw.p	adj.p
3	Fresh	2.77462	2.60912	2.94011	0	0
2	8SBF	3.06000	2.87954	3.24045	0	0
1	8BF	3.10465	2.81385	3.39545	0	0

Removed fresh treatment and fit a model with males and females combined to see if there was a difference in intercepts between the 8% buffered formalin and 8% sugar buffered formalin group.

```
> detach(mysis3.nograv)
> mysis3.nograv.nofresh <- mysis3.nograv[mysi3.nograv$tx != "Fresh",
+ ]
> mysis3.nograv.nofresh$tx <- factor(mysis3.nograv.nofresh$tx)
> attach(mysis3.nograv.nofresh)
> lm1a <- lm(logwt ~ loglen + tx)
```

Further comparisons indicated that the 8% sugar-buffered formalin treatment had a larger mean log(weight) than the 8% buffered formalin treatment for all values of log(length) ($p < 0.0005$).

```
> anova(lm1a)
```

Analysis of Variance Table

Response: logwt

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
loglen	1	8.9499	8.9499	2647.80	< 2.2e-16 ***
tx	1	0.5505	0.5505	162.86	< 2.2e-16 ***
Residuals	149	0.5036	0.0034		

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
> comp.intercepts(lm1a)
```

Tukey HSD on adjusted means assuming parallel lines.

	comparison	diff	lwr	upr	p.adj
1	8SBF-8BF	0.1360738	0.1153065	0.1568411	0

Mean adjusted values at a covariate value of 1.10621751474566

	8BF	8SBF
	0.6738179	0.8098917

```
> detach(mysis3.nograv.nofresh)
```

2.4 Second Length-Weight Model Analysis

The second model examined the effect of two preservation treatments (fresh preserved and 8

2.4.1 Temporarily Excludes 8% Buffered Formaling Treatment

```
> mysis3.no8BF <- mysis3[mysis3$tx != "8BF", ]
> mysis3.no8BF$stage <- factor(mysis3.no8BF$stage)
> mysis3.no8BF$tx <- factor(mysis3.no8BF$tx)
> mysis3.no8BF$treat <- factor(mysis3.no8BF$treat)
> attach(mysis3.no8BF)
> ftable(xtabs(~tx + stage + len.bin))
```

		len.bin	7	9	11	13	15	17
tx	stage							
8SBF	Fem		4	10	12	19	14	1
	Grav		0	0	0	12	12	3
	Male		0	5	12	32	1	0
Fresh	Fem		3	11	10	19	13	5
	Grav		0	0	1	12	6	3
	Male		2	9	15	23	5	0

2.4.2 Fitting The Model

```
> lm2 <- lm(logwt ~ loglen * tx * stage)
```

Assumption Checking

None of these results are in the manuscript. The normality test and non-constant variance test both indicate assumption violations. However, the residual plot shows a general homoscedasticity (with the exception of two minor outliers and a small narrowing at large lengths) and the histogram is largely symmetric.

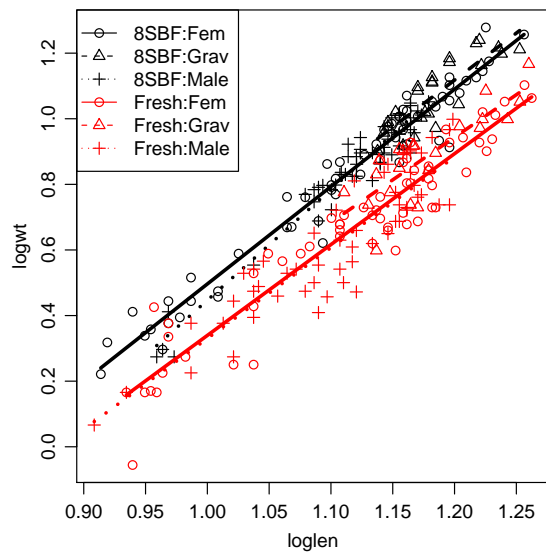
```
> fit.plot(lm2, leg = "topleft", main = "")
> ad.test(lm2$residuals)
```

Anderson-Darling normality test

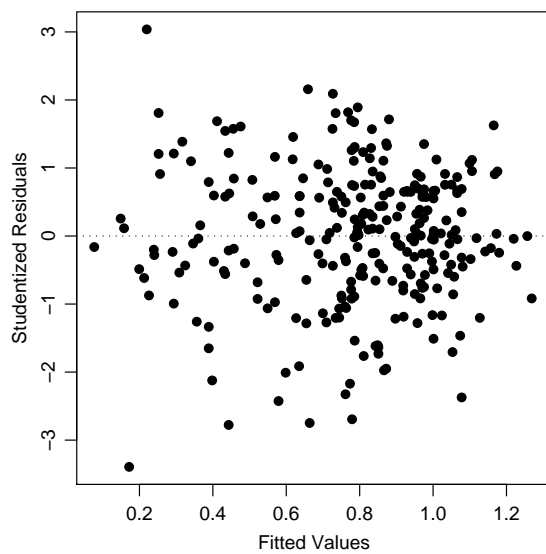
```
data: lm2$residuals
A = 0.6848, p-value = 0.07297
```

```
> ncv.test(lm2)
```

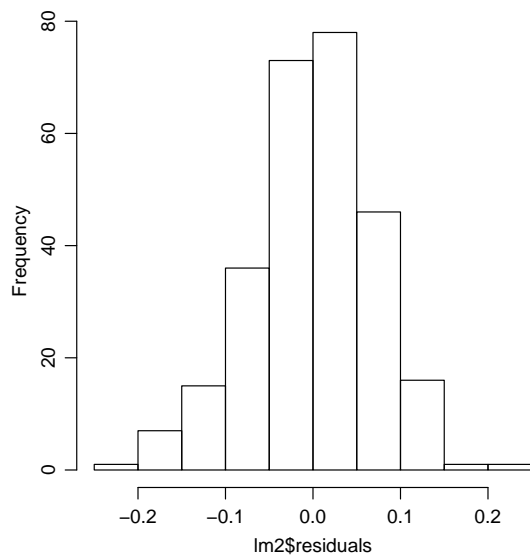
```
Non-constant Variance Score Test
Variance formula: ~ fitted.values
Chisquare = 13.57363    Df = 1    p = 0.0002293850
```



```
> residual.plot(lm2, main = "")
```



```
> hist(lm2$residuals, main = "")
```



Results

The second linear model fit to all three life-stages and the fresh preserved and 8% sugar-buffered formalin treatments showed that there was a significantly steeper slope for the 8% sugar-buffered formalin treatment ($p=0.012$) but no difference in slope among the three life-stages (0.4124).

```
> Anova(lm2)
```

```
Anova Table (Type II tests)
```

```
Response: logwt
```

	Sum Sq	Df	F value	Pr(>F)
loglen	11.8507	1	2326.8062	< 2.2e-16 ***
tx	2.1941	1	430.7889	< 2.2e-16 ***
stage	0.0605	2	5.9392	0.003002 **
loglen:tx	0.0326	1	6.4052	0.011965 *
loglen:stage	0.0091	2	0.8888	0.412395
tx:stage	0.0103	2	1.0138	0.364265
loglen:tx:stage	0.0108	2	1.0634	0.346778
Residuals	1.3344	262		

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> comp.slopes(lm2)
```

```
Multiple comparison control procedures used: fdr
```

```
Multiple Slope Comparisons
```

comparison	diff	lwr	upr	raw.p	adj.p
1 Fresh-8SBF	-0.25238	-0.47839	-0.02638	0.02876	0.02876

```
Slope Information
level slopes lwr upr raw.p adj.p
2 Fresh 2.84961 2.69511 3.00411 0 0
1 8SBF 3.10199 2.93705 3.26694 0 0
```

Within a preservation treatment there was a significant difference in mean log(weight) adjusted for log(length) (0.0241) with gravid females being significantly heavier than male or non-gravid females for all values of log(length) ($p < 0.0336$).

```
> lm2a <- lm(logwt ~ loglen + stage)
> anova(lm2a)
```

Analysis of Variance Table

```
Response: logwt
      Df Sum Sq Mean Sq  F value    Pr(>F)
loglen   1 14.4866  14.4866 1085.5607 < 2e-16 ***
stage    2  0.1009   0.0504   3.7795 0.02405 *
Residuals 270  3.6031   0.0133
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> comp.intercepts(lm2a)
```

```
Tukey HSD on adjusted means assuming parallel lines.
comparison      diff      lwr      upr      p.adj
1 Grav-Fem  0.049036498  0.003022977  0.095050020  0.03361352
2 Male-Fem -0.007539737 -0.043875446  0.028795972  0.87659483
3 Male-Grav -0.056576235 -0.103661253 -0.009491217  0.01375084
```

```
Mean adjusted values at a covariate value of 1.12492564403163
      Fem      Grav      Male
0.7755453 0.8245818 0.7680056
```

```
> detach(mysis3.no8BF)
```

2.5 Length-Weight Regressions for Different Groups

2.5.1 Males and Females in Fresh Group

This isolates just the fresh treatment and creates a stage variable that has gravid females and males/females combined. It then fits a model with a common slope and separate intercepts by the new stage variable.

```
> mysis3.fresh <- mysis3[mysi3$tx == "Fresh", ]
> mysis3.fresh$stage1 <- factor(ifelse(mysis3.fresh$stage == "Grav", "Grav",
+   "MF"))
> mysis3.fresh$tx <- factor(mysis3.fresh$tx)
> attach(mysis3.fresh)
> lm3.fresh <- lm(logwt ~ loglen + stage1)
> summary(lm3.fresh)
```



```

Call:
lm(formula = logwt ~ loglen + stage1)

Residuals:
    Min       1Q   Median       3Q      Max
-0.223649 -0.059177  0.007352  0.063210  0.209712

Coefficients:
              Estimate Std. Error t value Pr(>|t|)
(Intercept) -2.37190     0.11558 -20.522  < 2e-16 ***
loglen       2.76585     0.09684  28.562  < 2e-16 ***
stage1MF    -0.05854     0.02087  -2.806  0.00577 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.0853 on 134 degrees of freedom
Multiple R-Squared:  0.878,    Adjusted R-squared:  0.8762
F-statistic: 482.4 on 2 and 134 DF,  p-value: < 2.2e-16

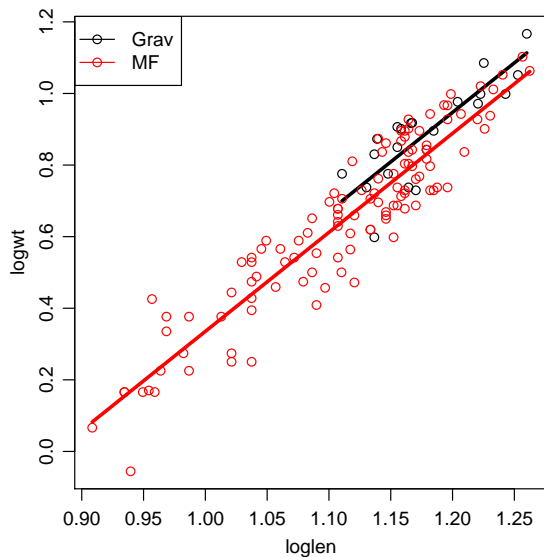
```

Thus, the equations of the lines are (these are in the manuscript)...

- Fresh,MF: $\log_{10}(W) = -2.4304 + 2.7659\log_{10}(L)$
- Fresh,Gravid: $\log_{10}(W) = -2.3719 + 2.7659\log_{10}(L)$

The following is not in the manuscript – just a visual check

```
> fit.plot(lm3.fresh, main = "", leg = "topleft")
```



```
> detach(mysis3.fresh)
```

2.5.2 Males and Females in Fresh Group

This isolates the 8BF and 8SBF groups (excludes fresh) and combines males and females as above. It then fits the linear model with a common slope and separate intercepts.

```
> mysis3.nfresh <- mysis3[mysis3$tx != "Fresh", ]
> mysis3.nfresh$stage1 <- factor(ifelse(mysis3.nfresh$stage == "Grav",
+   "Grav", "MF"))
> mysis3.fresh$tx <- factor(mysis3.fresh$tx)
> mysis3.nfresh$treat1 <- factor(mysis3.nfresh$stage1:mysis3.nfresh$tx)
> attach(mysis3.nfresh)
> lm3.nfresh <- lm(logwt ~ loglen + treat1)
> summary(lm3.nfresh)
```

Call:

```
lm(formula = logwt ~ loglen + treat1)
```

Residuals:

```
      Min       1Q   Median       3Q      Max
-0.217285 -0.029277  0.003547  0.043603  0.112868
```

Coefficients:

```
              Estimate Std. Error t value Pr(>|t|)
(Intercept)  -2.55867    0.07330  -34.906  <2e-16 ***
loglen         3.06650    0.06124   50.071  <2e-16 ***
treat1MF:8BF  -0.15985    0.01536  -10.406  <2e-16 ***
treat1MF:8SBF -0.02362    0.01310   -1.804    0.073 .
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Residual standard error: 0.05762 on 175 degrees of freedom
Multiple R-Squared:  0.9528,    Adjusted R-squared:  0.952
F-statistic: 1178 on 3 and 175 DF,  p-value: < 2.2e-16
```

Thus, the equations of the lines are (these are in the manuscript)...

- 8BF,MF: $\log_{10}(W) = -2.7185 + 3.0665\log_{10}(L)$
- 8SBF,MF: $\log_{10}(W) = -2.5823 + 3.0665\log_{10}(L)$
- 8SBF,Gravid: $\log_{10}(W) = -2.5587 + 3.0665\log_{10}(L)$

The following is not in the manuscript, it is just a visual and computational check.

```
> comp.intercepts(lm3.nfresh, 0)
```

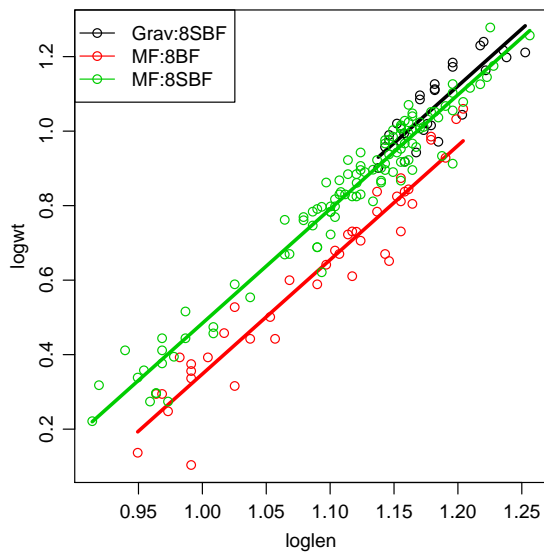
Tukey HSD on adjusted means assuming parallel lines.

```
      comparison      diff      lwr      upr      p.adj
1  MF:8BF-Grav:8SBF -0.15984574 -0.1933487 -0.126342799 6.061818e-14
2  MF:8SBF-Grav:8SBF -0.02362329 -0.0527940  0.005547414 1.376686e-01
3    MF:8SBF-MF:8BF  0.13622245  0.1115867  0.160858192 4.918288e-14
```

Mean adjusted values at a covariate value of 0

```
Grav:8SBF  MF:8BF  MF:8SBF
-2.558665 -2.718511 -2.582288
```

```
> fit.plot(lm3.nfresh, main = "", leg = "topleft")
```



```
> detach(mysis3.nfresh)
```

2.5.3 R-square For Above Equations

The R^2 for the above equations can be computed easily by exploiting the fact that the R^2 for an equation can be computed with the R^2 between the observed response values and the predicted response values from the equation.

Begin by isolating the data for each of the five different groups.

```
> mysis4 <- mysis3
> mysis4$stage1 <- factor(ifelse(mysis4$stage == "Grav", "Grav", "MF"))
> data.Fmf <- subset(mysis4, stage1 == "MF" & tx == "Fresh")
> data.Fg <- subset(mysis4, stage1 == "Grav" & tx == "Fresh")
> data.8Bmf <- subset(mysis4, stage1 == "MF" & tx == "8BF")
> data.8Smf <- subset(mysis4, stage1 == "MF" & tx == "8SBF")
> data.8Sg <- subset(mysis4, stage1 == "Grav" & tx == "8SBF")
```

Then find predicted $\log_{10}(W)$ for each group

```
> pred.Fmf <- 2.7659 * data.Fmf$loglen - 2.4304
> pred.Fg <- 2.7659 * data.Fg$loglen - 2.3719
> pred.8Bmf <- 3.0665 * data.8Bmf$loglen - 2.7185
> pred.8Smf <- 3.0665 * data.8Smf$loglen - 2.5823
> pred.8Sg <- 3.0665 * data.8Sg$loglen - 2.5587
```

Then compute regressions

```
> lm.Fmf <- lm(pred.Fmf ~ data.Fmf$logwt)
> lm.Fg <- lm(pred.Fg ~ data.Fg$logwt)
> lm.8Bmf <- lm(pred.8Bmf ~ data.8Bmf$logwt)
> lm.8Smf <- lm(pred.8Smf ~ data.8Smf$logwt)
> lm.8Sg <- lm(pred.8Sg ~ data.8Sg$logwt)
```

And then extract R^2 values

```
> summary(lm.Fmf)$r.squared
```

```
[1] 0.8688818
```

```
> summary(lm.Fg)$r.squared
```

```
[1] 0.6865749
```

```
> summary(lm.8Bmf)$r.squared
```

```
[1] 0.9206869
```

```
> summary(lm.8Smf)$r.squared
```

```
[1] 0.9501737
```

```
> summary(lm.8Sg)$r.squared
```

```
[1] 0.7329088
```

For comparison purposes, the following is the R^2 from the fit of the linear model just to the Fresh, MF group. This is on the raw data, as if the model was fit to just this group, and is not used in the manuscript.

```
> lm.Fmf1 <- lm(data.Fmf$logwt ~ data.Fmf$loglen)
> coef(lm.Fmf1)
```

```
(Intercept) data.Fmf$loglen
-2.440190      2.774618
```

```
> summary(lm.Fmf1)$r.squared
```

```
[1] 0.8688818
```

2.5.4 The Plot for the Manuscript

```
> plot(mysis4$c.wt ~ mysis4$len, log = "xy", col = "white", axes = FALSE,
+      xlab = xlbl1, ylab = ylbl1, xlim = c(8, 20), ylim = c(0.8, 20))
> axis(2, at = c(0.8, 0.9, seq(1, 10, by = 1), 20), labels = F)
> axis(2, at = c(1, 5, 10, 20), labels = T)
> axis(1, at = c(8:20), labels = F)
> axis(1, at = c(8, 10, 15, 20), labels = T)
> points(data.Fmf$c.wt ~ data.Fmf$len, col = "black", pch = 19)
> points(data.Fg$c.wt ~ data.Fg$len, col = "black", pch = 17, cex = 1.25)
> points(data.8Bmf$c.wt ~ data.8Bmf$len, col = "black", pch = 0)
> points(data.8Smf$c.wt ~ data.8Smf$len, col = "black", pch = 1)
> points(data.8Sg$c.wt ~ data.8Sg$len, col = "black", pch = 2)
> lines(c(8.1, 19.5), 10^c(2.7659 * log10(8.1) - 2.4304, 2.7659 * log10(19.5) -
+   2.4304), lwd = 3, col = "black", lty = 1)
> lines(c(12.9, 19.5), 10^c(2.7659 * log10(12.9) - 2.3719, 2.7659 * log10(19.5) -
+   2.3719), lwd = 3, col = "black", lty = 3)
> lines(c(8.9, 19.5), 10^c(3.0665 * log10(8.9) - 2.7185, 3.0665 * log10(19.5) -
+   2.7185), lwd = 3, col = "gray", lty = 2)
> lines(c(8.2, 19.5), 10^c(3.0665 * log10(8.2) - 2.5823, 3.0665 * log10(19.5) -
+   2.5823), lwd = 3, col = "gray", lty = 1)
> lines(c(13.7, 19.5), 10^c(3.0665 * log10(13.7) - 2.5587, 3.0665 * log10(19.5) -
+   2.5587), lwd = 3, col = "gray", lty = 3)
> legend(x = 14, y = 1.5, legend = c("Fresh,MF", "Fresh,Grav"), lty = c(1,
+   3), lwd = 3, inset = 0.02)
> legend(x = 14, y = 1.5, legend = c("", ""), pch = c(19, 17), pt.cex = 1.25,
+   inset = 0.02, bty = "n")
> legend("topleft", legend = c("8BF,MF", "8SBF,MF", "8SBF,Grav"), lty = c(2,
+   1, 3), lwd = 3, col = "gray", inset = 0.02)
> legend("topleft", legend = c("", "", ""), pch = c(0, 1, 2), pt.cex = 1.25,
+   inset = 0.02, bty = "n")
```

