Host activity before and during infection influences resulting parasite intensities

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```
# Clear the working environment
rm(list = ls())
# Visualization
library(ggplot2)
library(visreg)
library(gridExtra)
source("http://highstat.com/Books/BGS/GAMM/RCodeP2/HighstatLibV6.R")
# (generalized) Linear mixed modeling
library(lme4)
library(glmmTMB)
library(lmodel2)
# Statistcal analysis reporting and model validation
library(performance)
library(car)
library(lmtest)
library(DHARMa)
# Data wrangling
library(dplyr)
library(plyr)
library(tidyverse)
library(tidylog)
library(splancs)
```

Visualize some patterns in the raw data

```
IndBehav7 <- IndBehav7 %>%
   rename(Wormaf = Wormbf, AUC = AUC2)
## rename: renamed 2 variables (AUC, Wormaf)
# Subsetting down to Infected individuals
IndBehav7I <- IndBehav7 %>%
   filter(Infection == 1 & wormJump >= 0)
## filter: removed 172 rows (50%), 175 rows remaining
# getting the mean and SD of worms jumped
summary(IndBehav7I$wormJump)
##
      Min. 1st Qu. Median
                              Mean 3rd Qu.
                                               Max.
     0.000 2.000 2.000 1.909 2.000
                                              3.000
sqrt(var(IndBehav7I$wormJump))
## [1] 0.4189282
# Setting some of the factors back to factors
IndBehav7$fishID <- as.factor(IndBehav7$fishID)</pre>
IndBehav7$TrialTime <- as.factor(IndBehav7$TrialTime)</pre>
IndBehav7$Sex <- as.factor(IndBehav7$Sex)</pre>
IndBehav7$Infection <- as.factor(IndBehav7$Infection)</pre>
IndBehav7$ContInf <- as.factor(IndBehav7$ContInf)</pre>
IndBehav7$Died <- as.factor(IndBehav7$Died)</pre>
IndBehav7$Treatment <- as.factor(IndBehav7$Treatment)</pre>
# Calculating tissue tolerance for each individual.
IndBehav7 <- IndBehav7 %>%
   mutate(ChSMI = LateSMI - PreSMI) %>%
   mutate(TisTol = ChSMI/Totworm)
## mutate: new variable 'ChSMI' (double) with 85 unique values and 0% NA
## mutate: new variable 'TisTol' (double) with 47 unique values and 46% NA
# Subsetting down to female and males only to scale their SMI Males
IndBehavM <- IndBehav7 %>%
   filter(Sex == "M") %>%
   mutate(ScPSMI = scale(PreSMI), ScLSMI = scale(LateSMI))
## filter: removed 180 rows (52%), 167 rows remaining
## mutate: new variable 'ScPSMI' (double) with 41 unique values and 0% NA
           new variable 'ScLSMI' (double) with 41 unique values and 0% NA
```

```
# Females
IndBehavF <- IndBehav7 %>%
   filter(Sex == "F") %>%
   mutate(ScPSMI = scale(PreSMI), ScLSMI = scale(LateSMI))
## filter: removed 167 rows (48%), 180 rows remaining
## mutate: new variable 'ScPSMI' (double) with 45 unique values and 0% NA
          new variable 'ScLSMI' (double) with 45 unique values and 0% NA
# Combine the two separate dataframes together
IndBehav8 <- rbind(IndBehavF, IndBehavM)</pre>
# Scaling some variables to make them biologically comparable and better for
# model fitting
IndBehav8 <- IndBehav8 %>%
   mutate(ScVarvelBef = c(scale(VarvelBef)), ScNRatebf = c(scale(NRatebf)), ScBehavTol = c(scale(BehavTol))
        ScTotworm = c(scale(Totworm)), ScAUC = c(scale(AUC)), ScBehavVig = c(scale(BehavVig)),
        ScChSMI = c(scale(ChSMI)), ScTisTol = c(scale(TisTol)))
## mutate: new variable 'ScVarvelBef' (double) with 86 unique values and 2% NA
           new variable 'ScNRatebf' (double) with 85 unique values and 37% NA
           new variable 'ScBehavTol' (double) with 49 unique values and 1% NA
##
##
           new variable 'ScTotworm' (double) with 47 unique values and 46% NA
           new variable 'ScAUC' (double) with 49 unique values and 43% NA
           new variable 'ScBehavVig' (double) with 87 unique values and 1\% NA
##
           new variable 'ScChSMI' (double) with 85 unique values and 0% NA
           new variable 'ScTisTol' (double) with 47 unique values and 46% NA
##
# Filtering down to only infected and only individuals who have VIE injections
# for sample size questions
IndBehavind <- IndBehav8 %>%
   distinct(fishID, .keep_all = TRUE) %>%
   filter(Treatment == "VIE") %>%
   filter(Infection == "1")
## distinct: removed 260 rows (75%), 87 rows remaining
## filter: removed 52 rows (60%), 35 rows remaining
## filter: removed 18 rows (51%), 17 rows remaining
IndBehavCh <- IndBehav8 %>%
   filter(TrialTime != "Before") %>%
   mutate(TrialTime = recode(TrialTime, Early = "1", Late = "2", Later = "3"))
## filter: removed 86 rows (25%), 261 rows remaining
## mutate: changed 261 values (100%) of 'TrialTime' (0 new NA)
IndBehavCh$TrialTime <- as.factor(IndBehavCh$TrialTime)</pre>
IndBehavCh$Infection <- as.factor(IndBehavCh$Infection)</pre>
IndBehavChI <- IndBehavCh %>%
   filter(Infection == "1")
```

filter: removed 117 rows (45%), 144 rows remaining

```
IndBehavChU <- IndBehavCh %>%
  filter(Infection == "0")
```

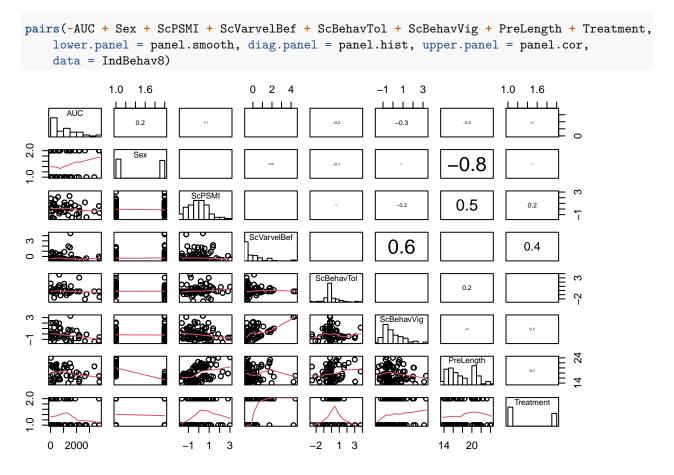
filter: removed 144 rows (55%), 117 rows remaining

What hypotheses we want to test with these data and what data we can use to test them?

- Does host behavioral vigor trade-off with their ability to resist parasite infection?
- Does higher host activity during infection lead to higher parasite infection?

What factors are important for host infection intensity over the course of infection and do is there any sexual variation in host infection intensity?

Visually inspection of the explanatory variables that will be used in the analyses



Description, development, and fitting of linear model for the analysis

We will use a linear model to analyze how infection intensity differs by sexual variation and other important host traits. Given each host only has one infection intensity measure, we do not need the fishID random effect used in previous models.

- Deterministic
- $AUC_{det} = a + b_1 Sex + b_2 ScBehavVig + b_3 ScVarVelBef + b_4 ScResidPLSMI + b_5 ScRPLength + b_6 Sex: ScBehavVig + b_7 Sex: ScResidPLSMI$
- Stochastic

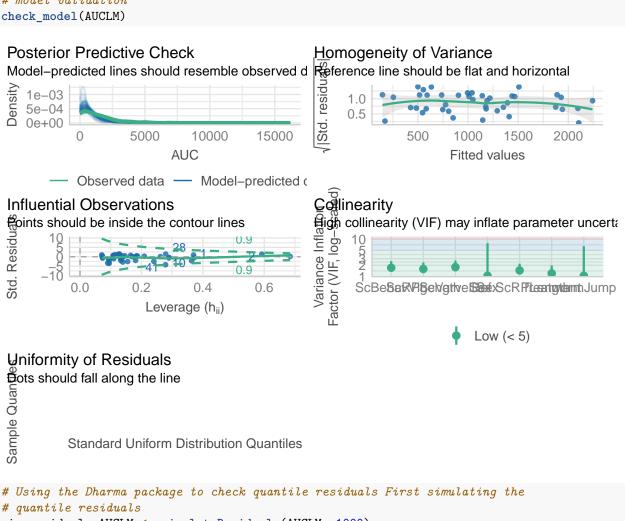
```
- AUC ~ N(AUC_{det}, \sigma^2)
```

- Fixed
 - Sex
 - Scaled behavioral vigor
 - Scaled Pre-infection SMI
 - Scaled variance in velocity before infection
 - Scaled Residuals of Length and Sex
 - VIE treatment
 - Interaction between Sex and behavioral vigor
 - Interaction between sex and body condition

```
##
## Call:
  glm(formula = AUC ~ Sex + ScBehavVig + ScVarvelBef + ScRPLength +
      Treatment + wormJump + Sex:ScRPLength, family = Gamma(link = "log"),
##
##
      data = IndBehavI)
##
## Coefficients:
                  Estimate Std. Error t value Pr(>|t|)
##
## (Intercept)
                              0.61592 10.062 9.95e-12 ***
                  6.19754
## SexM
                   0.53484
                              0.26311
                                       2.033 0.04995 *
## ScBehavVig
                  -0.55887
                              0.15749 -3.549 0.00116 **
## ScVarvelBef
                   0.45869
                              0.18094
                                       2.535 0.01601 *
## ScRPLength
                  -0.55197
                              0.16684 -3.308 0.00222 **
## TreatmentVIE
                              0.29171 -0.178 0.85975
                  -0.05194
## wormJump
                   0.20270
                              0.30905
                                       0.656 0.51631
## SexM:ScRPLength 0.51726
                              0.29635
                                      1.745 0.08994
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
##
## (Dispersion parameter for Gamma family taken to be 0.689384)
##
##
      Null deviance: 69.030 on 41 degrees of freedom
## Residual deviance: 55.847 on 34 degrees of freedom
    (6 observations deleted due to missingness)
## AIC: 675.03
```

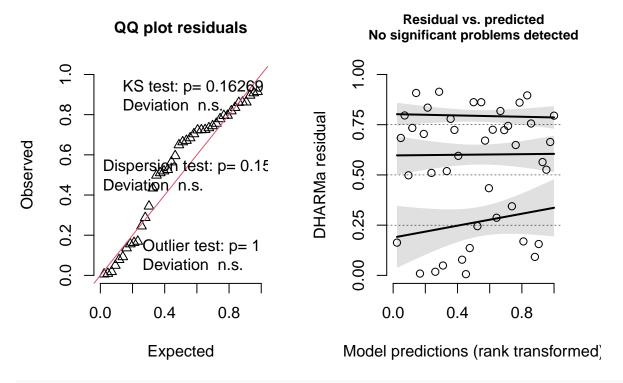
Validate that the model fits well and there are no problems

Using the check_model function from the perforamnce package to check the
model validation
check_model(AUCLM)



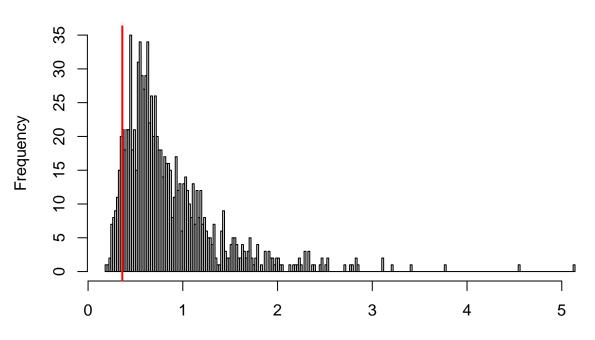
```
# Using the Dharma package to check quantile residuals First simulating the
# quantile residuals
sim_residuals_AUCLM <- simulateResiduals(AUCLM, 1000)
# Plotting the quantile residuals to test how quantile residuals look
plot(sim_residuals_AUCLM)</pre>
```

DHARMa residual



Testing for dispersion
testDispersion(sim_residuals_AUCLM)

DHARMa nonparametric dispersion test via sd of residuals fitted vs. simulated



Simulated values, red line = fitted model. p-value (two.sided) = 0.154

```
##
## DHARMa nonparametric dispersion test via sd of residuals fitted vs.
## simulated
##
## data: simulationOutput
## dispersion = 0.42402, p-value = 0.154
## alternative hypothesis: two.sided
```

Testing the significance of factors in our model

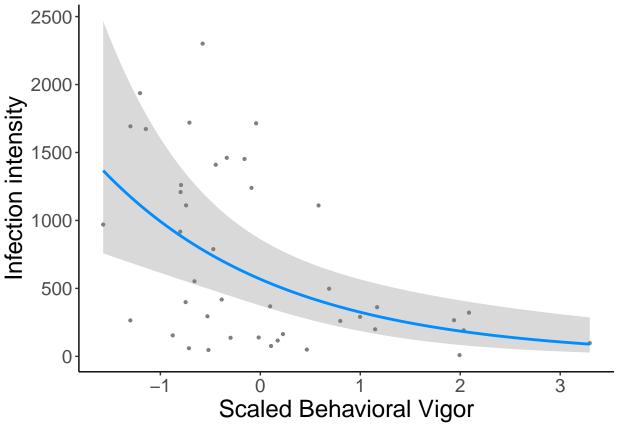
```
Anova(AUCLM, type = 3)
## Analysis of Deviance Table (Type III tests)
##
## Response: AUC
##
               LR Chisq Df Pr(>Chisq)
                3.9769 1 0.046127 *
## Sex
               10.4695 1 0.001214 **
## ScBehavVig
               7.2344 1 0.007152 **
## ScVarvelBef
## ScRPLength
                6.8738 1 0.008747 **
## Treatment
                0.0325 1 0.856908
## wormJump 0.5585 1 0.454867
## Sex:ScRPLength 2.5644 1 0.109294
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
```

Visualize the important explanatory factors for infection intensity

```
# Behavioral Vigor graph
InfIntbyVig = visreg(AUCLM, scale = "response", "ScBehavVig", partial = T, gg = TRUE) +
    theme_classic() + theme(legend.position = "none") + ylab("Infection intensity") +
    xlab("Scaled Behavioral Vigor") + theme(text = element_text(size = 18))

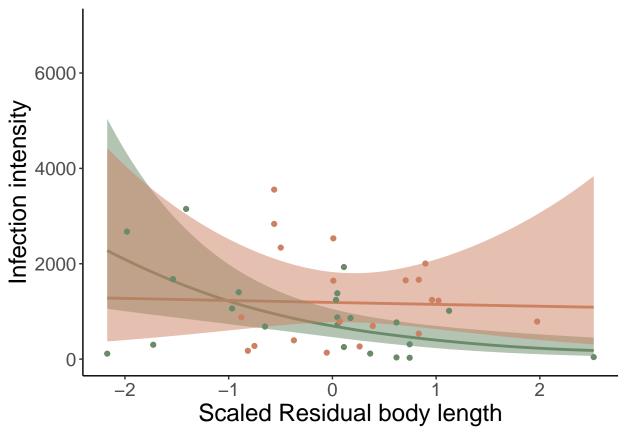
## Conditions used in construction of plot
## Sex: F
## ScVarvelBef: -0.5119451
## ScRPLength: 0.04746859
## Treatment: UNTOUCHED
## wormJump: 2

# Print the graph
print(InfIntbyVig)
```



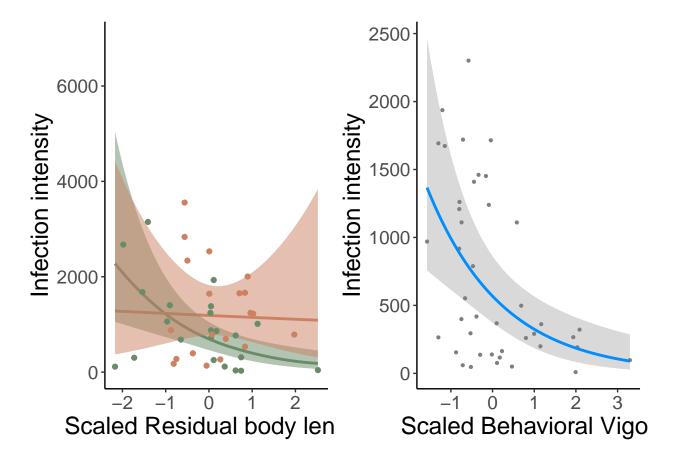
```
# Behavioral Vigor graph Extracting fit and residuals from model
InfIntbyLen = visreg(AUCLM, scale = "response", "ScRPLength", "Sex", partial = T,
   plot = FALSE, overlay = T)
# Extracting fit
InfIntbyLenfit <- InfIntbyLen$fit</pre>
# Extracting residuals
InfIntbyLenres <- InfIntbyLen$res</pre>
cpsex = c("darkseagreen4", "lightsalmon3")
InfIntbyLengraph <- ggplot() + geom_smooth(data = InfIntbyLenfit, aes(x = ScRPLength,</pre>
    y = visregFit, group = Sex, color = Sex)) + geom_ribbon(data = InfIntbyLenfit,
    aes(x = ScRPLength, y = visregFit, ymin = visregLwr, ymax = visregUpr, fill = Sex),
    alpha = 0.5) + geom_point(data = InfIntbyLenres, aes(x = ScRPLength, y = visregRes,
    group = Sex, color = Sex)) + theme_classic() + theme(legend.position = "none") +
   ylab("Infection intensity") + xlab("Scaled Residual body length") + theme(text = element_text(size
    scale_color_manual(values = cpsex) + scale_fill_manual(values = cpsex) + ylim(0,
    7000)
# Print the graph
print(InfIntbyLengraph)
```

'geom_smooth()' using method = 'loess' and formula = 'y ~ x'



```
# Sex by SMI graph Extracting the fit and residuals
InfIntbySex = visreg(AUCLM, scale = "response", "Sex", partial = T, plot = FALSE)
# Extract fit
InfIntbySexFit <- InfIntbySex$fit</pre>
# Extract residuals
InfIntbySexRes <- InfIntbySex$res</pre>
# Add colorscheme by sex
cpsex = c("darkseagreen4", "lightsalmon3")
# Plot the graph
InfSexgraph <- ggplot(data = InfIntbySexRes, aes(x = Sex, y = visregRes)) + geom_boxplot(aes(fill = Sex</pre>
    geom_jitter(alpha = 0.5) + theme_classic() + theme(legend.position = "none") +
    ylab("Infection intensity") + xlab(" ") + theme(text = element_text(size = 18)) +
    scale_fill_manual(values = cpsex) + scale_x_discrete(breaks = c("F", "M"), labels = c("Females",
    "Males"))
# Loading in gridExtra for multiple graphs in one image
library(gridExtra)
InfIntgraphs <- grid.arrange(InfIntbyLengraph, InfIntbyVig, nrow = 1)</pre>
```

'geom_smooth()' using method = 'loess' and formula = 'y ~ x'



Fitting posthoc model to test for pattern within sex

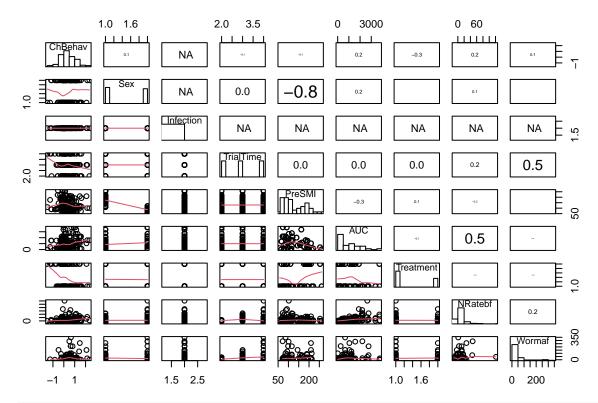
```
# Subsetting down to females only
IndBehavIF <- IndBehavI %>%
    filter(Sex == "F")
## filter: removed 23 rows (48%), 25 rows remaining
# Subsetting down to males only
IndBehavIM <- IndBehavI %>%
    filter(Sex == "M")
## filter: removed 25 rows (52%), 23 rows remaining
# Fit a linear model for infection intensity by sex Female model
AUCLMF <- glm(AUC ~ ScBehavVig + ScVarvelBef + ScResidPLSMI + ScRPLength + Treatment,
    family = Gamma(link = "log"), IndBehavIF)
# Male model
AUCLMM <- glm(AUC ~ ScBehavVig + ScVarvelBef + ScResidPLSMI + ScRPLength + Treatment,
    family = Gamma(link = "log"), IndBehavIM)
# Summary to see the relationship of the variables. Females
summary(AUCLMF)
```

```
##
## Call:
## glm(formula = AUC ~ ScBehavVig + ScVarvelBef + ScResidPLSMI +
      ScRPLength + Treatment, family = Gamma(link = "log"), data = IndBehavIF)
## Coefficients:
               Estimate Std. Error t value Pr(>|t|)
                            0.2365 27.004 5.12e-16 ***
## (Intercept)
                 6.3853
## ScBehavVig
                -0.2783
                            0.1928 -1.444 0.166036
## ScVarvelBef
                 0.3585
                            0.3033
                                    1.182 0.252570
## ScResidPLSMI -0.4320
                            0.2557 -1.689 0.108417
## ScRPLength
                -0.6701
                            0.1582 -4.237 0.000496 ***
                                    0.970 0.345034
## TreatmentVIE 0.3790
                            0.3909
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for Gamma family taken to be 0.6339407)
##
      Null deviance: 44.276 on 23 degrees of freedom
## Residual deviance: 35.674 on 18 degrees of freedom
    (1 observation deleted due to missingness)
## AIC: 376.28
##
## Number of Fisher Scoring iterations: 22
# Males
summary(AUCLMM)
##
## Call:
## glm(formula = AUC ~ ScBehavVig + ScVarvelBef + ScResidPLSMI +
      ScRPLength + Treatment, family = Gamma(link = "log"), data = IndBehavIM)
##
## Coefficients:
               Estimate Std. Error t value Pr(>|t|)
##
                           0.25048 28.785 3.28e-15 ***
## (Intercept)
                7.21005
                           0.26406 -2.635
## ScBehavVig
               -0.69585
                                             0.0180 *
## ScVarvelBef
                0.51982
                           0.25857
                                    2.010
                                             0.0616 .
## ScResidPLSMI -0.27102
                           0.19305 - 1.404
                                             0.1795
## ScRPLength
                0.03954
                                    0.143
                                             0.8884
                           0.27736
## TreatmentVIE -0.30223
                           0.44366 -0.681
                                             0.5055
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for Gamma family taken to be 0.701784)
##
      Null deviance: 23.688 on 21 degrees of freedom
## Residual deviance: 18.147 on 16 degrees of freedom
    (1 observation deleted due to missingness)
## AIC: 368.34
## Number of Fisher Scoring iterations: 9
```

```
# Testing for significance for females
Anova(AUCLMF, type = 3, test = "LR")
## Analysis of Deviance Table (Type III tests)
##
## Response: AUC
##
               LR Chisq Df Pr(>Chisq)
## ScBehavVig
                 1.5676 1
                              0.210552
## ScVarvelBef
                 1.5190 1
                              0.217771
## ScResidPLSMI 3.1025 1
                              0.078170 .
## ScRPLength
                 8.3693 1
                              0.003816 **
## Treatment
                 0.7356 1
                              0.391066
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
# Testing for significance for males
Anova(AUCLMM, type = 3, test = "LR")
## Analysis of Deviance Table (Type III tests)
## Response: AUC
##
               LR Chisq Df Pr(>Chisq)
## ScBehavVig
                 6.3953 1
                               0.01144 *
## ScVarvelBef
                 3.9527 1
                               0.04680 *
## ScResidPLSMI 1.3761 1
                              0.24077
                               0.90005
## ScRPLength
                 0.0158 1
## Treatment
                 0.3521 1
                               0.55291
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
Examining change in activity for only infected individuals
# Setting some variables as factors to confirm they are this way for the
# analysis
IndBehavChI$Sex <- as.factor(IndBehavChI$Sex)</pre>
IndBehavChI$Infection <- as.factor(IndBehavChI$Infection)</pre>
IndBehavChI$Treatment <- as.factor(IndBehavChI$Treatment)</pre>
# Subsetting our dataframe to relevant data and making some
IndBehavChI <- IndBehavChI %>%
   mutate(ScWormaf = scale(Wormaf)) %>%
   mutate(ScChBehav = scale(ChBehav))
## mutate: new variable 'ScWormaf' (double) with 43 unique values and 53% NA
## mutate: new variable 'ScChBehav' (double) with 135 unique values and 7% NA
pairs(~ChBehav + Sex + Infection + TrialTime + PreSMI + AUC + Treatment + NRatebf +
   Wormaf, lower.panel = panel.smooth, diag.panel = panel.hist, upper.panel = panel.cor,
   data = IndBehavChI)
```

```
## Warning in par(usr): argument 1 does not name a graphical parameter
## Warning in par(usr): argument 1 does not name a graphical parameter
## Warning in par(usr): argument 1 does not name a graphical parameter
## Warning in cor(x, y, use = "pairwise.complete.obs"): the standard deviation is
## Warning in cor(x, y, use = "pairwise.complete.obs"): the standard deviation is
## zero
## Warning in par(usr): argument 1 does not name a graphical parameter
## Warning in cor(x, y, use = "pairwise.complete.obs"): the standard deviation is
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```

```
## Warning in cor(x, y, use = "pairwise.complete.obs"): the standard deviation is
## zero
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## Warning in par(usr): argument 1 does not name a graphical parameter
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## zero
## Warning in par(usr): argument 1 does not name a graphical parameter
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## zero
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## Warning in par(usr): argument 1 does not name a graphical parameter
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## Warning in par(usr): argument 1 does not name a graphical parameter
## Warning in par(usr): argument 1 does not name a graphical parameter
```



cpsex = c("darkseagreen4", "lightsalmon3")

We will use a linear mixed model to analyze how Change in activity differs by sexual variation and how infected an individual fish was. FishID is included as a random term to allow for non-independence of individuals due to multiple measurements per individual across time.

- Deterministic
- $Log_{10}(InfInt_{det}+1)=a+b_1TrialTime+b_2ChBehav+b_3Sex+b_4ScPSMI+b_5Treatment+b_6wormJump+b_7ScChSMI+b_8BehavVig+a_i$
- Stochastic

- ChVel ~
$$N(Log_{10}(InfInt_{det} + 1), \sigma^2)$$

- $a_i \sim N(0, \sigma_{fishID}^2)$

- Fixed
 - TrialTime
 - Change in behavior between time points
 - Sex
 - Scaled pre-infection scaled mass index
 - VIE treatment of the fish
 - number of worms that initially started the infection
 - Scaled change in scaled mass index
 - Behavior vigor of each fish
- Random
 - fishID

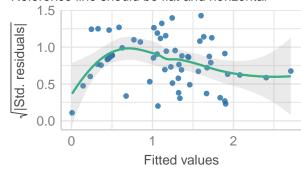
```
# Fit a linear model for checking what explanatory factors are important for
# Variance in activity Note this is a linear mixed model because we have
# multiple measures per fish and therefore, need to account for
# non-independence between measures.
InfIntLMInf <- lmer(log10(Wormaf + 1) ~ Sex + ScPSMI + ScChSMI + TrialTime + ScBehavVig +</pre>
   Treatment + ChBehav + wormJump + (1 | fishID), IndBehavChI)
# Summary to see the relationship of the variables.
summary(InfIntLMInf)
## Linear mixed model fit by REML ['lmerMod']
## Formula: log10(Wormaf + 1) ~ Sex + ScPSMI + ScChSMI + TrialTime + ScBehavVig +
##
      Treatment + ChBehav + wormJump + (1 | fishID)
##
     Data: IndBehavChI
##
## REML criterion at convergence: 129.3
##
## Scaled residuals:
                     Median
       Min
                1Q
                                   30
                                           Max
## -2.03509 -0.69817 -0.04025 0.72865 1.60854
##
## Random effects:
           Name
## Groups
                        Variance Std.Dev.
           (Intercept) 0.005027 0.0709
## fishID
                        0.406932 0.6379
## Residual
## Number of obs: 61, groups: fishID, 21
## Fixed effects:
##
               Estimate Std. Error t value
## (Intercept) 0.55319 0.41074
                                   1.347
## SexM
               -0.11606
                           0.18735 -0.619
## ScPSMI
               -0.06753
                           0.14011 -0.482
                          0.13309 -0.567
## ScChSMI
               -0.07540
## TrialTime2
              0.77036 0.19797
                                   3.891
## TrialTime3
              1.35524 0.20857
                                   6.498
## ScBehavVig
                           0.13822
                                   0.526
                0.07273
## TreatmentVIE 0.20701
                           0.22032
                                    0.940
                                   2.339
## ChBehav
                           0.19621
                0.45893
## wormJump
               -0.18333
                           0.17373 - 1.055
## Correlation of Fixed Effects:
##
            (Intr) SexM ScPSMI ScCSMI TrlTm2 TrlTm3 ScBhvV TrtVIE ChBehv
## SexM
              0.015
              0.228 -0.281
## ScPSMI
## ScChSMI
              0.193 -0.100 0.559
## TrialTime2 -0.197 0.018 -0.008 -0.020
## TrialTime3 -0.334 -0.011 0.007 0.016 0.444
              0.149 -0.018  0.523  0.334 -0.049
## ScBehavVig
## TreatmntVIE -0.377   0.004 -0.440 -0.412   0.007 -0.023 -0.541
## ChBehav
           -0.388 -0.168  0.072  0.190 -0.105  0.240  0.468 -0.068
## wormJump
              -0.860 -0.256 -0.073 -0.051 -0.029 0.074 -0.118 0.175 0.277
```

Validate that the model fits well and there are no problems

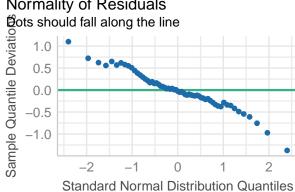
```
# Using the check_model function from the perforamnce package to check the
# model validation
check_model(InfIntLMInf, check = c("qq", "normality", "homogeneity"))
```

Homogeneity of Variance

Reference line should be flat and horizontal

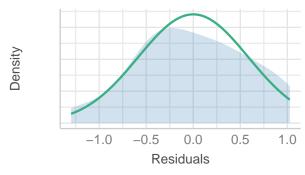


Normality of Residuals



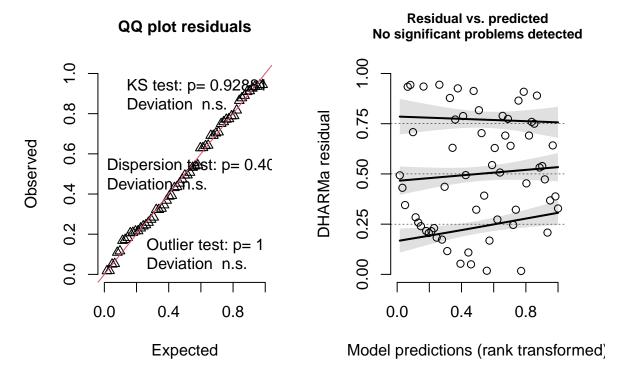
Normality of Residuals

Distribution should be close to the normal curve



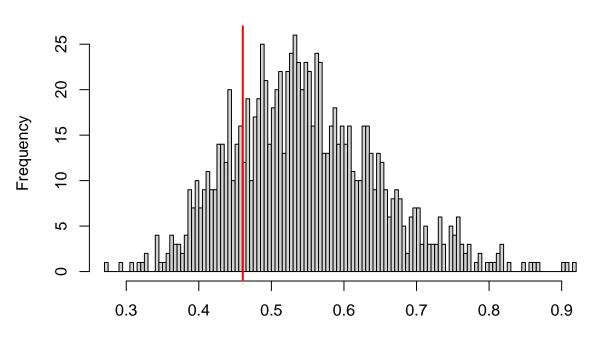
```
# Using the Dharma package to check quantile residuals First simulating the
# quantile residuals
sim_residuals_InfIntLMInf <- simulateResiduals(InfIntLMInf, 1000)</pre>
# Plotting the quantile residuals to test how quantile residuals look
plot(sim_residuals_InfIntLMInf)
```

DHARMa residual



Testing for dispersion
testDispersion(sim_residuals_InfIntLMInf)

DHARMa nonparametric dispersion test via sd of residuals fitted vs. simulated



Simulated values, red line = fitted model. p-value (two.sided) = 0.408

```
##
## DHARMa nonparametric dispersion test via sd of residuals fitted vs.
## simulated
##
## data: simulationOutput
## dispersion = 0.84272, p-value = 0.408
## alternative hypothesis: two.sided

# There are some problems with this model validation. It doesnt look model
# breaking but definitely should look at other model error structures to
# resolve the issues.
```

Testing the significance of factors in our model using a Kenward-Rodgers F test for infected only infection intensity model

```
# F test to test for signficance of slope of variables
Anova(InfIntLMInf, test = "F", type = 3)
## Analysis of Deviance Table (Type III Wald F tests with Kenward-Roger df)
## Response: log10(Wormaf + 1)
                   F Df Df.res
                               Pr(>F)
## (Intercept) 1.7844 1 17.834 0.19841
## Sex
             0.3819 1 13.514 0.54686
## ScPSMI
             0.2320 1 12.774 0.63816
## ScChSMI
             0.3180 1 15.068 0.58110
## TrialTime 21.5646 2 40.186 4.341e-07 ***
## ScBehavVig 0.2706 1 16.492 0.60982
## Treatment 0.8818 1 12.696 0.36524
## ChBehav
             4.9482 1 42.669 0.03146 *
             1.1043 1 13.713 0.31148
## wormJump
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

Visualize the important explanatory factors for infection intensity for infected individuals

```
## 'geom_smooth()' using formula = 'y ~ x'
Log (number of parasites + 1)
                                                                                     2
                                        0
                                   Change in activity
# Subset to only infected females
IndBehavChIF <- IndBehavChI %>%
    filter(Sex == "F" & Infection == "1") %>%
    mutate(ScWormaf = scale(Wormaf))
## filter: removed 69 rows (48%), 75 rows remaining
## mutate: changed 32 values (43%) of 'ScWormaf' (0 new NA)
# Subset to only infected males
IndBehavChIM <- IndBehavChI %>%
    filter(Sex == "M" & Infection == "1") %>%
    mutate(ScWormaf = scale(Wormaf))
## filter: removed 75 rows (52%), 69 rows remaining
```

mutate: changed 36 values (52%) of 'ScWormaf' (0 new NA)

```
# Fit linear mixed models to post hoc check for significance for male and
# female slopes
ChVelLMInfM <- lmer(ChBehav ~ ScPSMI + ScChSMI + TrialTime + ScBehavVig + Treatment +
   ScNRatebf + (1 | fishID), IndBehavChIM)
# Summary to see the relationship of the variables for female model.
summary(ChVelLMInfF)
## Linear mixed model fit by REML ['lmerMod']
## Formula: ChBehav ~ ScPSMI + ScChSMI + TrialTime + ScBehavVig + Treatment +
##
      ScNRatebf + (1 | fishID)
##
     Data: IndBehavChIF
##
## REML criterion at convergence: 115.2
## Scaled residuals:
       Min
                    Median
                1Q
                                  3Q
## -2.40025 -0.59721 0.01767 0.39831 2.49141
##
## Random effects:
## Groups Name
                       Variance Std.Dev.
## fishID (Intercept) 0.1516 0.3893
## Residual
                        0.1465
                                0.3827
## Number of obs: 74, groups: fishID, 25
##
## Fixed effects:
##
              Estimate Std. Error t value
## (Intercept) 0.35892 0.13500 2.659
## ScPSMI
               0.03171 0.19273 0.165
## ScChSMI
               -0.12455 0.20790 -0.599
## TrialTime2 -0.12062 0.12073 -0.999
## TrialTime3 -0.08185 0.11226 -0.729
## ScBehavVig
             -0.29924
                          0.09676 -3.093
## TreatmentVIE -0.14657
                          0.20917 -0.701
## ScNRatebf
             0.20238
                          0.07599 2.663
##
## Correlation of Fixed Effects:
             (Intr) ScPSMI ScCSMI TrlTm2 TrlTm3 ScBhvV TrtVIE
## ScPSMI
              0.239
## ScChSMI
             -0.110 -0.849
## TrialTime2 -0.419 -0.040 0.004
## TrialTime3 -0.423 -0.024 0.008 0.525
## ScBehavVig 0.035 -0.134 0.264 -0.057 -0.030
## TreatmntVIE -0.553 -0.435  0.344 -0.012  0.010  0.015
## ScNRatebf 0.134 0.090 -0.008 -0.443 -0.210 0.128 0.026
# Summary to see the relationship of the variables for male model.
summary(ChVelLMInfM)
## Linear mixed model fit by REML ['lmerMod']
## Formula: ChBehav ~ ScPSMI + ScChSMI + TrialTime + ScBehavVig + Treatment +
##
      ScNRatebf + (1 | fishID)
```

```
##
     Data: IndBehavChIM
##
## REML criterion at convergence: 87.2
##
## Scaled residuals:
##
       Min
              1Q
                     Median
                                   3Q
                                           Max
## -1.77079 -0.56125 -0.08803 0.54617 2.27980
##
## Random effects:
## Groups
            Name
                        Variance Std.Dev.
## fishID
             (Intercept) 0.1907
                                0.4366
                        0.1050
                                 0.3240
## Residual
## Number of obs: 60, groups: fishID, 22
##
## Fixed effects:
##
               Estimate Std. Error t value
                           0.14960
## (Intercept)
              0.55518
                                    3.711
## ScPSMI
               -0.13224
                           0.18568 -0.712
                           0.17887 -0.596
## ScChSMI
               -0.10664
## TrialTime2
                0.13984
                           0.10301
                                    1.358
## TrialTime3
              -0.28284
                           0.12219 -2.315
## ScBehavVig -0.42170
                           0.10922 -3.861
                           0.24076 -0.790
## TreatmentVIE -0.19015
## ScNRatebf
             -0.02104
                           0.03640 -0.578
##
## Correlation of Fixed Effects:
##
              (Intr) ScPSMI ScCSMI TrlTm2 TrlTm3 ScBhvV TrtVIE
## ScPSMI
              -0.175
## ScChSMI
              -0.120 0.769
## TrialTime2 -0.335 -0.007 -0.010
## TrialTime3 -0.302 -0.027 -0.029 0.518
## ScBehavVig
              0.243 0.240 -0.007 -0.040 -0.094
## TreatmntVIE -0.603 0.134 0.221 0.003 0.013 -0.400
## ScNRatebf
               0.078 0.021 0.032 -0.317 -0.439 0.127 -0.010
# Anova to test for significance for females
Anova(ChVelLMInfF, test = "F", type = 2)
## Analysis of Deviance Table (Type II Wald F tests with Kenward-Roger df)
##
## Response: ChBehav
##
                  F Df Df.res
                                Pr(>F)
## ScPSMI
             0.0271 1 19.801 0.870991
## ScChSMI
             0.3589 1 19.547 0.555988
## TrialTime 0.5239 2 47.777 0.595551
## ScBehavVig 9.5583 1 20.050 0.005741 **
## Treatment 0.4909 1 19.840 0.491657
## ScNRatebf 6.8075 1 61.242 0.011393 *
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
# Anova to test for significance for males
Anova(ChVelLMInfM, test = "F", type = 2)
```