Omit two sequencing runs and luminal samples and perform combat seq

Then filter on sequencing depth of 10k or less, and prevalence filter to 10%

Genotype* Site_General interaction

```
Permutation: free
Number of permutations: 10000
Terms added sequentially (first to last)
                    Df SumsOfSqs MeanSqs F.Model R2
                                                        Pr(>F)
                         21.98 21.975 18.305 0.03730 9.999e-05 ***
Sequencing_Run
                    1
                          2.78 2.782 2.317 0.00472 0.09689 .
                     1
Sex
Genotype
                     2
                         38.68 19.340 16.110 0.06566 9.999e-05 ***
                    1 192.28 192.285 160.170 0.32640 9.999e-05 ***
Site_General
                         2.06 1.028 0.856 0.00349 0.49525
Genotype:Site_General 2
                         331.34 1.201
                   276
Residuals
                                              0.56243
Total
                   283
                         589.11
                                              1.00000
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Genotype* Site Interaction

```
Permutation: free
Number of permutations: 10000
Terms added sequentially (first to last)
               Df SumsOfSqs MeanSqs F.Model R2 Pr(>F)
                      21.98 21.975 21.635 0.03730 9.999e-05 ***
Sequencing_Run 1
                      2.78 2.782 2.739 0.00472 0.06259 .
               1
Sex
              2 38.68 19.340 19.040 0.06566 9.999e-05 ***
Genotype
Site 5 249.42 49.884 49.113 0.42339 9.999e-05 ***
Genotype:Site 10 8.11 0.811 0.798 0.01376 0.72883
Residuals 264 268.15 1.016
                                           0.45517
             283 589.11
Total
                                           1.00000
Signif. codes: 0 '*** '0.001 '** '0.01 '* '0.05 '.' 0.1 ' '1
```