## The allometry of field metabolic rates in tropical vs. temperate hummingbirds

Anusha Shankar, Joseph R. Canepa, Catherine H. Graham, Donald R. Powers Thursday, March 17, 2016

## Abstract

Allometric scaling relationships describe how the scaling of one trait (e.g. body size) affects the scaling of another trait (e.g. field metabolic rate, brain size). These scaling relationships have intrigued scientists for decades because there seem to be rules that govern how metabolic processes scale up with body size. In birds, field metabolic rate (FMR) scales with body mass (M) in the form FMR = aM0.67. This relationship between metabolic rate and mass has been found to vary depending on the taxonomic level being studied. Thus, a scaling exponent on the scale of all birds could mask localized taxon-specific patterns. Hummingbirds have among the highest mass-specific rates of all vertebrates, as well as a much higher wing aspect ratio than predicted for their size. We collected FMR and mass data for hummingbirds, and examined the scaling of FMR with body mass among hummingbird. We found that hummingbirds have an FMR to mass scaling exponent >0.9, much higher than any group previously studied. Further, since temperate hummingbirds feed on densely clumped resources more than tropical hummingbirds do, we expect tropical birds to have a higher field metabolic rate to body size exponent than temperate birds to support increased foraging effort. We find that tropical hummingbirds have a higher FMR to mass ratio on average than temperate birds.

Tropical birds have been found to have significantly lower BMR than temperate birds, though within the tropics (Peru) elevation had no effect on BMR (Londono et al. 2015). 'Pace of life' is thus slower in tropical than temperate regions.

## **Introduction:**

**Hypotheses**: 1. Whole animal daily energy expenditure will increase as mass increases 2. Daily energy expenditure will increase if daily temperatures vary more widely + Higher thermoregulatory costs (avg. daily temperatures are low) would contribute to increased DEE (both directly because of increased thermoregulatory costs, and indirectly because they would need to be more active to gain energy to deal with these increased costs) 3. Tropical resident species have a higher DEE than temperate long distance migrants 4. Territorial species

A log-log plot of hummingbird daily energy expenditure (kJ) vs. mass (g), including values from this study as well as from the literature. Colors represent species.

Mass-corrected daily energy expenditure vs. Mass, colored by species

Plot of total daily energy expenditure per bird, colored by species (not mass-corrected)

```
##
## lm(formula = log(kJ_day) ~ log(Initial_mass), data = dlw)
##
## Residuals:
                       Median
        Min
                  1Q
                                             Max
  -0.65537 -0.25841 0.03412 0.25133
                                         0.94224
##
##
## Coefficients:
                     Estimate Std. Error t value Pr(>|t|)
                                 0.13276
                                            16.32
## (Intercept)
                      2.16647
                                                    <2e-16 ***
```

```
## log(Initial_mass) 0.93116
                                0.07984
                                         11.66 <2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.342 on 88 degrees of freedom
## Multiple R-squared: 0.6072, Adjusted R-squared: 0.6027
## F-statistic: 136 on 1 and 88 DF, p-value: < 2.2e-16
##
## Call:
## lm(formula = log(kJ_day) ~ log(Initial_mass), data = dlw[dlw$Big_site ==
##
      "EC", ])
##
## Residuals:
       Min
                 1Q
                     Median
## -0.66219 -0.15843 0.02263 0.12887 0.57555
##
## Coefficients:
##
                    Estimate Std. Error t value Pr(>|t|)
                                 0.6824
                                          2.802 0.01413 *
## (Intercept)
                      1.9120
                      1.2073
                                 0.3191
                                          3.784 0.00201 **
## log(Initial_mass)
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.2866 on 14 degrees of freedom
## Multiple R-squared: 0.5056, Adjusted R-squared: 0.4703
## F-statistic: 14.32 on 1 and 14 DF, p-value: 0.002014
##
## Call:
## lm(formula = log(kJ_day) ~ log(Mass_g), data = dlw_mean)
##
## Residuals:
##
       Min
                 1Q
                     Median
                                   30
## -0.46776 -0.09714 0.00693 0.13382 0.40697
##
## Coefficients:
              Estimate Std. Error t value Pr(>|t|)
##
                2.1642
                           0.3626
                                    5.969 0.00021 ***
## (Intercept)
## log(Mass_g)
                0.9577
                           0.2015
                                    4.753 0.00104 **
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.2477 on 9 degrees of freedom
## Multiple R-squared: 0.7151, Adjusted R-squared: 0.6834
## F-statistic: 22.59 on 1 and 9 DF, p-value: 0.00104
##
## Call:
## lm(formula = log(kJ_day) ~ log(Mass_g) + Region, data = dlw_mean)
## Residuals:
##
       Min
                 1Q
                      Median
                                   3Q
                                           Max
```

```
## -0.21710 -0.09355 0.00264 0.06574 0.24674
##
## Coefficients:
##
            Estimate Std. Error t value Pr(>|t|)
## (Intercept) 2.3103
                      0.2285 10.112 1.99e-05 ***
                      0.1341 5.702 0.000734 ***
## log(Mass_g) 0.7646
## RegionCR
               ## RegionEC
               0.4460
                        0.1105 4.035 0.004965 **
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
\mbox{\tt \#\#} Residual standard error: 0.1538 on 7 degrees of freedom
## Multiple R-squared: 0.9145, Adjusted R-squared: 0.8779
## F-statistic: 24.97 on 3 and 7 DF, p-value: 0.0004105
```