

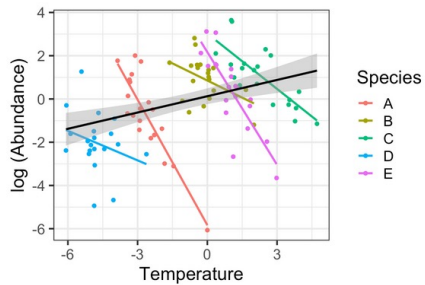
Job trends in statistics and data analysis

With so much hinging on statistics - national and international policy, funding, corporate decision making, government commitment and research - **demand for statisticians and data analysts is expected to grow around 34% between 2014 and 2024**. This vastly outstrips the average across all jobs.

source - <https://www.environmentalscience.org/career/environmental-data-analyst>

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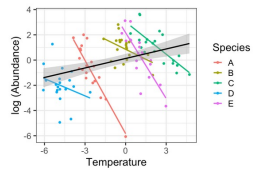
One common feature here is that species may differ in the way they are structured by environmental and/or trait variation. This can be well described by the Simpson's paradox (Simpson 1951), which is defined "as a phenomenon in probability and statistics in which a trend appears in several groups of data but disappears or reverses when the groups are combined."



Important enough to have its own Wikipedia page:
https://en.wikipedia.org/wiki/Simpson%27s_paradox

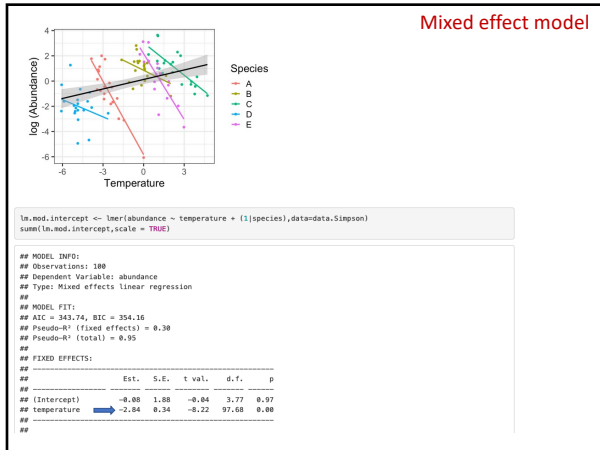
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Fixed effect model



```
## MODEL INFO:
## Observations: 100
## Dependent Variable: abundance
## Type: OLS linear regression
##
## MODEL FIT:
## F(1,98) = 15.13, p = 0.00
## R^2 = 0.13
## Adj. R^2 = 0.12
##
## Standard errors: OLS
##
##               Est.   S.E.   t val.   p
## [Intercept]   -0.08  0.18   -0.48   0.63
## scale(temperature)  0.09  0.10    3.09   0.00
##
## Continuous predictors are mean-centered and scaled by 1 s.d.
```

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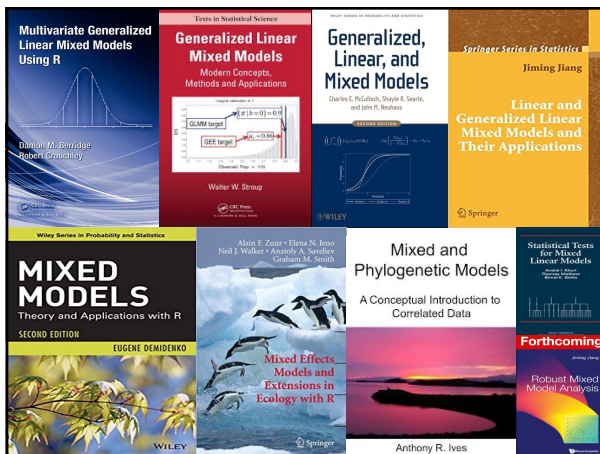
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General linear models (not Generalized linear model)

Linear Model	Common name
$Y = \mu + X$	Simple linear regression
$Y = \mu + A_1$	One-factorial (one-way) ANOVA
$Y = \mu + A_1 + A_2 + A_1 \times A_2$	Two-factorial (two-way) ANOVA
$Y = \mu + A_1 + X (+A_1 \times X)$	Analysis of Covariance (ANCOVA)
$Y = \mu + X_1 + X_2 + X_3$	Multiple regression
$Y = \mu + A_1 + g + A_1 \times g$	Mixed model ANOVA
$Y_1 + Y_2 = \mu + A_1 + A_2 + A_1 \times A_2$	Multivariate ANOVA (MANOVA)

Y (response) is a continuous variable
X (predictor) is a continuous variable
A represents categorical predictors (factors)
g represents groups of data (more on this later)
(+A₁ × X) - step 1 on an ANCOVA, but not in the final analysis
Multiple factors A₁ + A₂ + etc (and their interactions)

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**Fixed effects are often crossed in relation to other fixed effects
(e.g., typical two-way ANOVA)**

Fixed effect factor: Data have been gathered from all the levels of the factor that are of interest.

Example: Contrasting the effects of three specific dosages of a drug on the response. "Dosage" is the factor; the three specific dosages in the experiment are the levels; there is no intent to say anything about other dosages.

(source: <https://www.ma.utexas.edu/users/mks/statmistakes/fixedsrandom.html>)

Drug
(fixed factor)

Dose
(fixed factor)

Drug A

Drug B

D₁

D₂

D₃

D₄

D₁

D₂

D₃

D₄

each different dose is tested for (across) each drug (i.e., factors are crossed)

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Random versus fixed effects – a hierarchical view

Random effect factor (sometimes referred to as a variance component model): The factor has many possible levels. Although there is interest in all possible levels, only a random sample of levels can be included in the data (either due to lack of knowledge all all possible levels, or costs, or both, or other issues).

Example: In an animal breeding experiment conducted to estimate the breeding value of sires (father) from a certain breed, several sires were **randomly selected from a population** and each sire was mated with several dams (mother). The weights of all the newborn animals were recorded.

Breed
(fixed factor)

Sire
(random factor)

Breed A

Breed B

S₁

S₂

S₃

S₄

S₅

S₆

S₇

S₈

↑ ↑ ↑ ↑ ↑ ↑ ↑ ↑

different sires are used across different breeds

↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓

Newborns (offsprings)

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Random versus fixed effects – a hierarchical view

The factor is "sire" (male gamete). The analysis will not estimate the effect of each of the sires in the sample; instead, it will **estimate the variability attributable to the factor "sire"**.

It is a type of hierarchical linear model, which assumes that the data being analysed are drawn from a hierarchy of different populations whose differences relate to that hierarchy.

Breed
(fixed factor)

Sire
(random factor)

Breed A

Breed B

S₁

S₂

S₃

S₄

S₅

S₆

S₇

S₈

↑ ↑ ↑ ↑ ↑ ↑ ↑ ↑

different sires are used across different breeds
and sires are more related within than between breeds (pedigree is a hierarchical structure)

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Random effects are often **hierarchical** in relation to fixed effects

Here, sires are a random variable (factor) that will change from study to study. In contrast (based on another example), drug levels are of true interest that won't change from study to study.

Breed (fixed factor)

Siren (random factor)

Breed A	Breed B						
S ₁	S ₂	S ₃	S ₄	S ₅	S ₆	S ₇	S ₈

different sires are used across different breeds

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Random effects – let's focus on a single breed

In a one-way (factor) random effect ANOVA, the goal is to estimate the variance of a breed (variation among sires). The sires are merely a sample from which inferences are to be made concerning the single population (here Breed A).

weights of all the newborn animals (1 per dam) were recorded for each sire

Do sires vary in their newborn weights? This can be answered by assessing whether there is more variation between sires than within sires.

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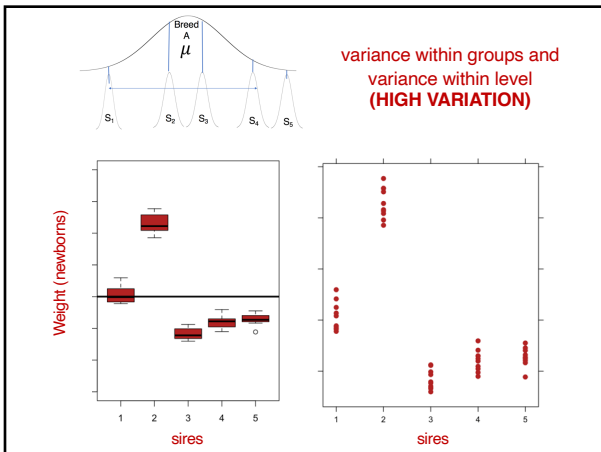
variance within groups and variance within level
(LOW VARIATION)

Weight (newborns)

sires

sires

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Fixed versus random effects may depend on the question and not always the data

STUDY: Suppose five cuts of meat are taken from each of three pigs, all from the same breed, and the fat content is measured in each cut.

FIXED EFFECT QUESTION – Do the different cuts differ in their fat content? One-way (fixed) ANOVA with five treatment levels (cuts) and three replicates per cut (pigs).

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```

cut.weights  cut.factor  pig
1 1.1 1 1
2 2.1 2 1
3 3.1 3 1
4 4.1 4 1
5 5.1 5 1
6 1.2 1 2
7 2.2 2 2
8 3.2 3 2
9 4.2 4 2
10 5.1 5 2
11 0.9 1 3
12 2.3 2 3
13 3.1 3 3
14 4.2 4 3
15 5.2 5 3
    
```

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```

pig1 <- c(1.1,2.1,3.1,4.1,5.1)
pig2 <- c(1.2,2.2,3.2,4.2,5.1)
pig3 <- c(0.9,2.3,3.1,4.2,5.2)

cut.weights <- c(pig1,pig2,pig3)
cut.factor <- as.factor(rep(1:5,3))
pig <- as.factor(c(rep(1,5),rep(2,5),rep(3,5)))
View(cbind(cut.weights,cut.factor,pig))
anova(lm(cut.weights~cut.factor))

```

	cut.weights	cut.factor	pig
1	1.1	1	1
2	2.1	2	1
3	3.1	3	1
4	4.1	4	1
5	5.1	5	1
6	1.2	1	2
7	2.2	2	2
8	3.2	3	2
9	4.2	4	2
10	5.1	5	2
11	0.9	1	3
12	2.3	2	3
13	3.1	3	3
14	4.2	4	3
15	5.2	5	3

```

> anova(lm(cut.weights~cut.factor))
Analysis of Variance Table

Response: cut.weights
          Df Sum Sq Mean Sq F value    Pr(>F)
cut.factor  4 30.6923   7.6573   863.54 1.07e-12 ***
Residuals 10   0.0867   0.0087
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

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Fixed versus random effects may depend on the question and not always the data

STUDY: Suppose five cuts of meat are taken from each of three pigs, all from the same breed, and the fat content is measured in each cut.

FIXED EFFECT QUESTION – Do the different cuts differ in their fat content? One-way (fixed) ANOVA with five treatment levels (cuts) and three replicates per cut (pigs).

RANDOM EFFECT QUESTION - Is there more variation in fat content among or within pigs (i.e., animal-to-animal and within-animal variation)? A fat pig could have their cuts fatter (i.e., hierarchical variation).

In this case, the three pigs selected are not of interest. This would be a one-way random effects ANOVA.

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Note that there is more variation within pigs (i.e., among cuts within pigs) than between pigs

```

pig1 <- c(1.1,2.1,3.1,4.1,5.1)
pig2 <- c(1.2,2.2,3.2,4.2,5.1)
pig3 <- c(0.9,2.3,3.1,4.2,5.2)

cut.weights <- c(pig1,pig2,pig3)
cut.factor <- as.factor(rep(1:5,3))
pig <- as.factor(c(rep(1,5),rep(2,5),rep(3,5)))
View(cbind(cut.weights,cut.factor,pig))
anova(lm(cut.weights~cut.factor))

```

```

library(nlme)
gls(cut.weights ~ 1, correlation=corCompSymm(form=~ 1 | pig))

```

```

Correlation Structure: Compound symmetry
Formula: ~1 | pig
Parameter estimate(s):
  Rho
-0.2490236
Degrees of freedom: 15 total; 14 residual
Residual standard error: 1.431177

```

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Now, these data have more variation between pigs (i.e., among cuts within pigs) than within pigs

```

pig1 <- c(1,1,2,1,3,1,4,1,5,1)
pig2 <- c(1,2,2,3,2,4,2,5,1) + 10
pig3 <- c(0,9,2,3,3,1,4,2,5,2) + 100
cut.weights <- c(pig1,pig2,pig3)
gls(cut.weights ~ 1, correlation=corCompSymm(forme ~ 1 | pig))
    
```

	cut.weights	cut.factor	pig
1	1.1	1	1
2	2.1	2	1
3	3.1	3	1
4	4.1	4	1
5	5.1	5	1
6	1.2	1	2
7	2.2	2	2
8	3.2	3	2
9	4.2	4	2
10	5.1	5	2
11	0.9	1	3
12	2.3	2	3
13	3.1	3	3
14	4.2	4	3
15	5.2	5	3

```

Correlation Structure: Compound symmetry
Formula: ~1 | pig
Parameter estimate(s):
  Rho
0.9991573
Degrees of freedom: 15 total; 14 residual
Residual standard error: 55.00793
    
```

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More on the hierarchical nature of data



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The advantages of mixed models - increase statistical power and estimation accuracy through dependent replication and design convenience (particularly in observational studies).

Do we need a random effect here?

Effects of temperature on fish growth (difference in growth begin/end of study)

Low temperature			
Intermediate temperature			
High temperature			

22

The advantages of mixed models - increase statistical power and estimation accuracy through dependent replication and design convenience (particularly in observational studies).

Do we need a random effect here?

Effects of temperature on fish growth
(difference in growth begin/end of study)

Low temperature			
Intermediate temperature			
High temperature			

23

The advantages of mixed models - increase statistical power and estimation accuracy through dependent replication and design convenience (particularly in observational studies).

Do we need a random effect here?

Effects of temperature on fish growth
(difference in growth begin/end of study)

Low temperature			
-----------------	--	--	--

24

The advantages of mixed models - increase statistical power and estimation accuracy through dependent replication and design convenience (particularly in observational studies).

Do we need a random effect here?

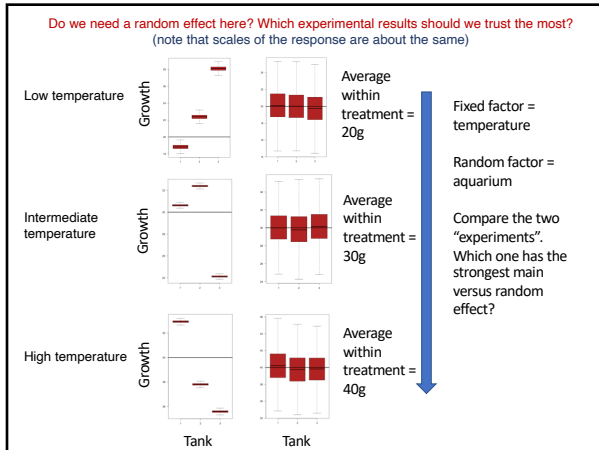
Effects of temperature on fish growth
(difference in growth begin/end of study)

Low temperature			
-----------------	--	--	--

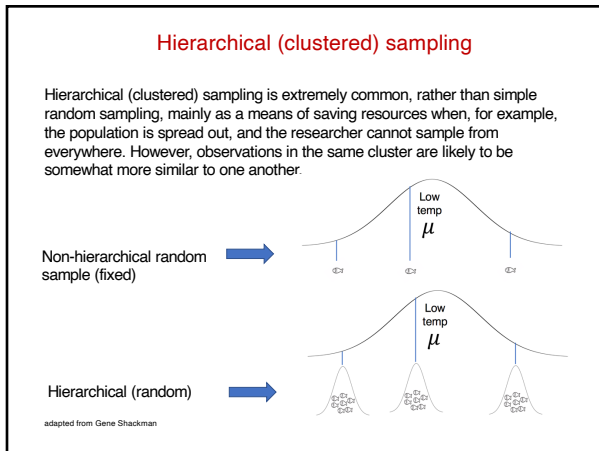
versus

Note that treatment (low temp.) means considering all individuals are the same in both situation

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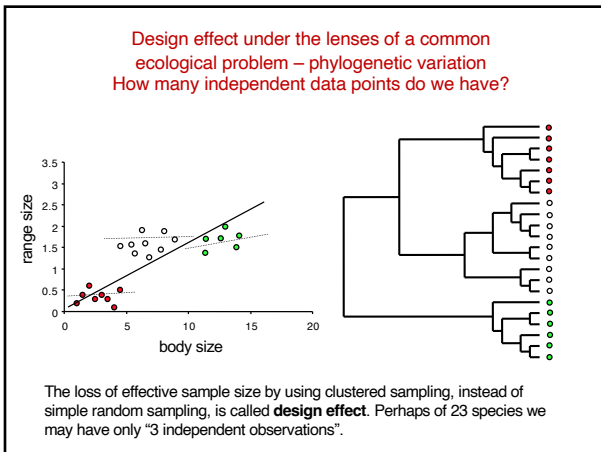


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Hierarchical (clustered) sampling

As a result, in a clustered sample selecting, an additional observation from the same cluster adds less new information than would a completely independent selection. As such, the sample is not as variable a random sample would be, so that the effective sample size is reduced. The loss of effective sample size by using clustered sampling, instead of simple random sampling, is called **design effect**.

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The American Naturalist

JOURNAL ARTICLE
Phylogenies and the Comparative Method

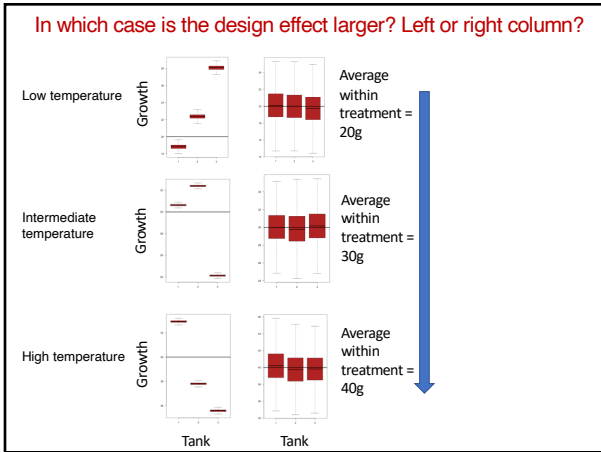
Joseph Felsenstein
The American Naturalist
Vol. 125, No. 1 (Jan., 1985), pp. 1-15

Published by: [The University of Chicago Press](http://www.jstor.org/stable/2461605) for [The American Society of Naturalists](http://www.jstor.org/stable/2461605)
<https://www.jstor.org/stable/2461605>
Page Count: 15

The loss of effective sample size by using clustered sampling, instead of simple random sampling, is called **design effect**. Perhaps of 23 species we may have only "3 independent observations".

INCREASED TYPE I ERROR IF KEPT AT 23 species (i.e., wrongly considering too many independent observations) AND LOSS OF POWER if only "3 species" is used.

30



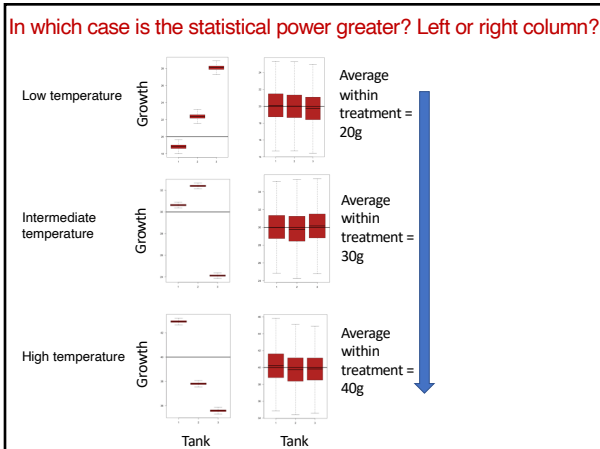
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Hierarchical (clustered) sampling

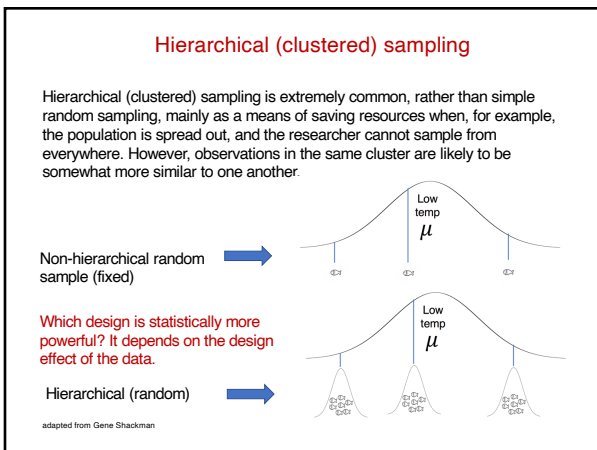
The **design effect** is a correction factor that is used to adjust the sample size based on clustered sampling. This accounts for the loss of information inherent in the clustered design and is used when estimating random effects.

Once the design effect is calculated, the sample size calculated for a standard design can be adjusted accordingly (i.e., degrees of freedom are corrected). As such, the statistical power may change according to the design effect.

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Intraclass correlation as a way to understand why to use a random effect and how much “random” has an “effect”



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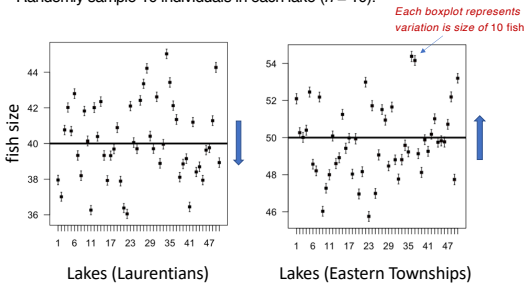
Hierarchical (clustered) sampling – intraclass correlation & design effect (one extreme example)

- Consider a study that wants to estimate fish size between two regions in Quebec (Laurentians and Eastern Townships).
- Draw a random sample of 50 lakes in each region.
- Randomly sample 10 individuals in each lake ($n = 10$).

One fixed factor = region (these two regions cannot change).
 One random factor = lakes (because they are more likely to be similar within regions and they are not crossed between regions, i.e., lakes differ).

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- Consider a study that wants to estimate fish size between two regions in Quebec (Laurentians and Eastern Townships).
- Draw a random sample of 50 lakes in each region.
- Randomly sample 10 individuals in each lake ($n = 10$).



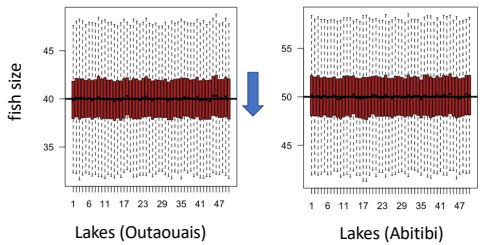
Very high (~1) intraclass correlation here because individuals are very similar within lakes than among lakes; so, a random effect would be important.

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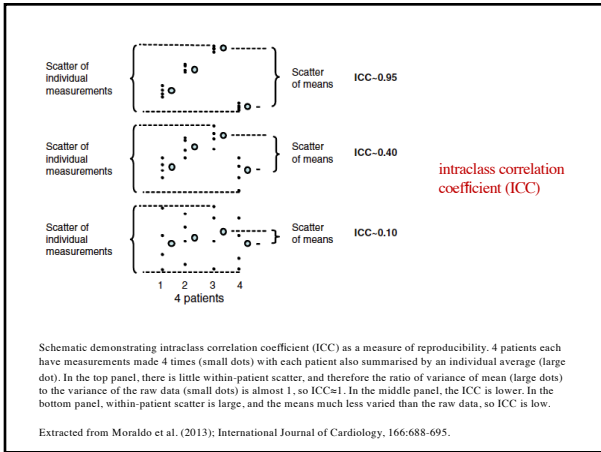
- Consider a study that wishes to estimate fish size between two regions in Quebec (Laurentians and Eastern Townships).
 - Draw a random sample of 50 lakes in each region.
 - Randomly sample 10 individuals in each lake ($n = 10$).
- Assume for the sake of discussion that all individuals within each lake had the exact same size but size differed between each of the lakes, then
- intraclass correlation = 1 and a design effect = $1 + 1(50 - 1) = 50$.*
- In this case, we have started our sample with 500 individuals across lakes, but the "design" can only "use" 50 values ("observations") in the analysis because all individuals were too similar within lakes! So, to increase the degrees of freedom, we would need to sample now 500 lakes per region instead! So, this design effect reduced the statistical power of the ANOVA.

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- Consider now these data (two other regions): If variation in size *within* each lake is the same for all lakes, then
- intraclass correlation = 0 and what we call design effect = $1 + 0(50 - 1) = 1$.*
- In this extreme case, each additional lake adds no new information about the fish size in each region.
 - Only surveying one lake would give us the same information (with the same standard error) about fish size as we get from surveying 50 lakes. So, to get the same degrees of freedom we only need to sample now 500 fish in a single lake per region instead of 10 individuals across 50 lakes!



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Mixed models mix random and fixed effects and allows estimating conducting statistical testing (inference) via proper estimation of design effects for hierarchical (clustered) sampling! It also affects parameter estimation (e.g., Simpson's paradox)

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Why consider a mixed-model?
 Some factors you may be able to control (fixed) and others you won't (random)

- (i) Models using random effects are important for inference when analyzing data that exhibit non-independence (hierarchical structure).
- (ii) Random effects provide a unifying statistical framework for models that might otherwise seem unrelated, for example, time-series models for populations, spatial models, genetics models, and models for variation among individuals;
- (iii) Models that include random effects are increasingly easy to build and customize for specific problems using publicly available modelling tools and software.

adapted from Thorson and Minto

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Time for reading



A brief introduction to mixed effects modelling and multi-model inference in ecology

Xavier A. Harrison¹, Lynda Donaldson^{2,3}, Maria Eugenia Correa-Cano⁵, Julian Evans^{4,5}, David N. Fisher^{4,6}, Cecily E.D. Goodwin⁷, Beth S. Robinson^{2,7}, David J. Hodgson⁸ and Richard Inger^{3,4}

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Mixed models mixes random and
fixed effects!

(next lecture)
