

Monte Carlo methods

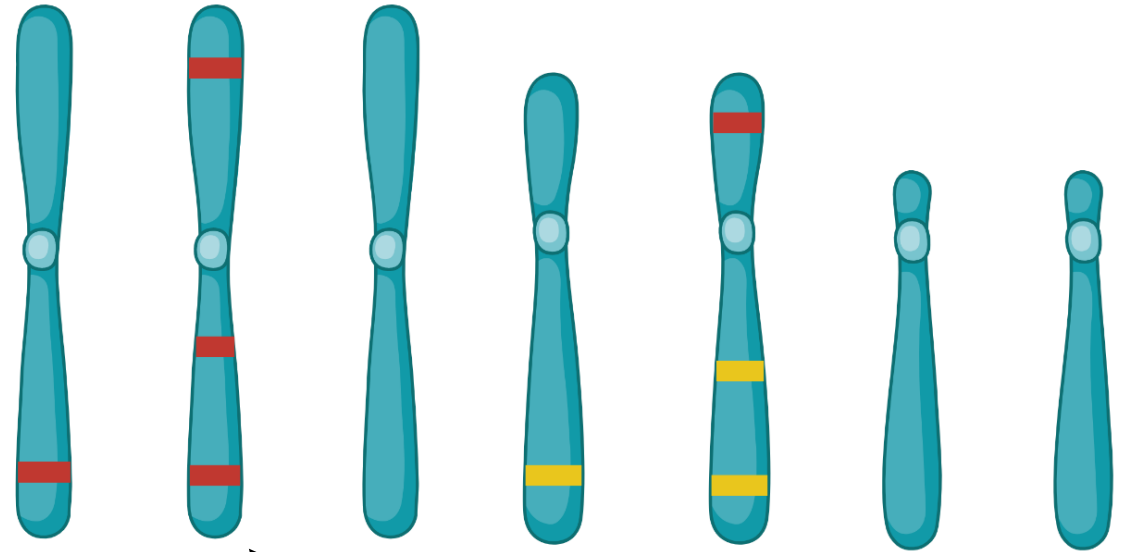
Named after the resort in Monaco well known for its casinos



You will eventually be faced with questions where you can't find a clear statistical test that will be appropriate to your data.



Are these beetles randomly distributed, aggregating or avoiding each other?



Does chromosome 2 have more “red” genes than I would expect by chance?

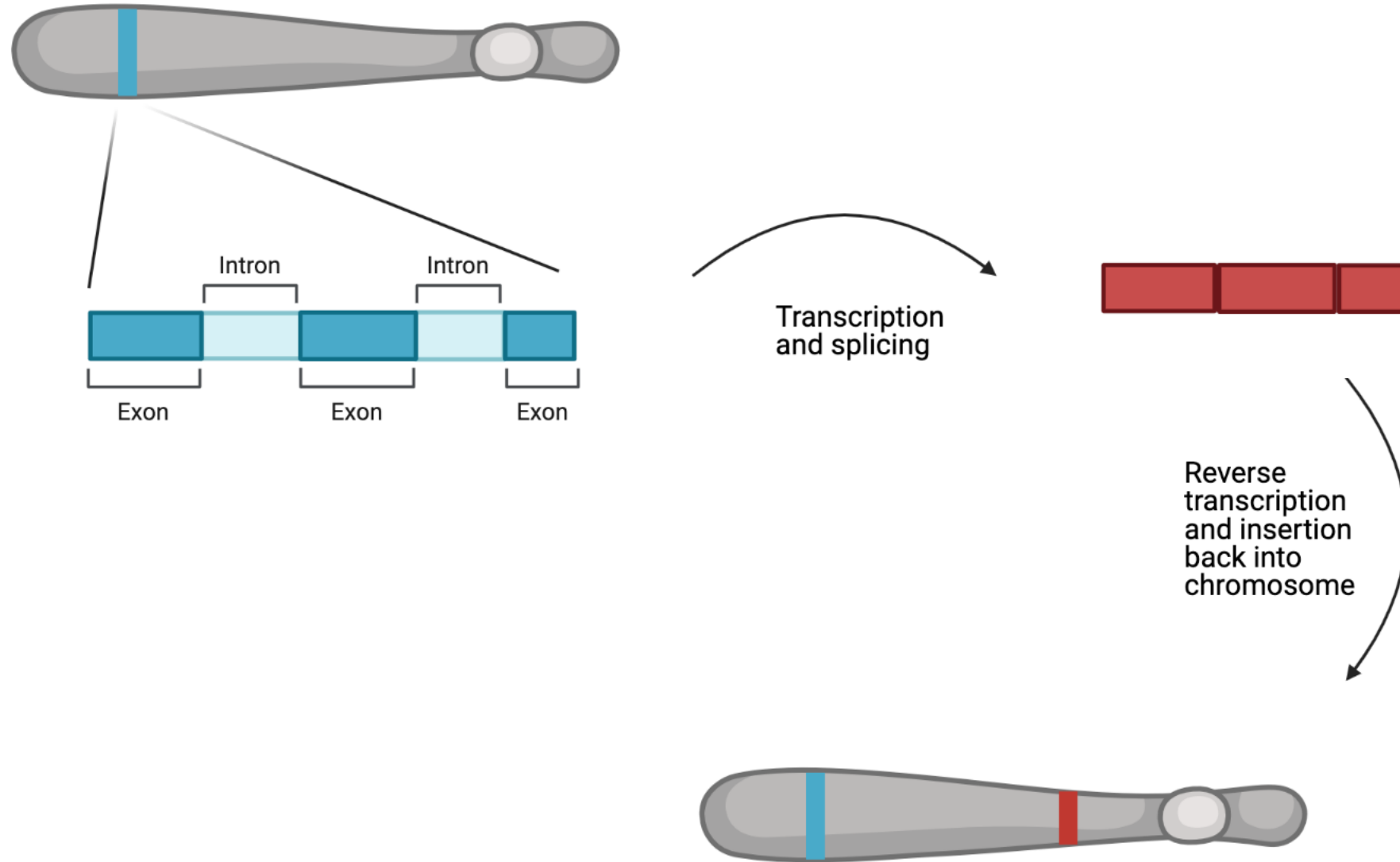
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Monte Carlo methods are most similar to permutation methods that we have used previously. The distinction between the two is that with permutations we simply randomized data. In contrast, with Monte Carlo we will use a simulation process that represents the biology of a null hypothesis.

Monte Carlo methods offer an approach to answer difficult often complex questions.

Lets look at an example with retrogenes.

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Sexual antagonism hypothesis of retrogene distribution:

If some genes have variation that is beneficial to one sex but harmful to the other fitness could be increased in two ways.

- 1) make a duplicate of a gene and express one version in males and one in females
- 2) make a duplicate of a gene and move it from an autosome to an X or Y chromosome

Additionally in many species the sex chromosomes do not get expressed during meiosis if you have a gene that can benefit males during spermatogenesis moving this type of gene from a sex chromosome to an autosome will be beneficial.

From this sexual antagonism hypothesis we might expect to see more retrogenes on sex chromosomes and possibly more parental genes on sex chromosomes than we would expect by chance.

We identified retrogenes in a Tribolium beetle that has a recent fusion of chromosome 2 (lg2) to the sex chromosome. This means that the X in this species now include everything that is in lg2 as well as lgX.

	lg1	lg2	lgX	lg4	lg5	lg6	lg7	lg8	lg9	lg10	Total
GeneNumber	3254	2941	3705	1507	1543	1287	994	684	325	212	16512
PhysicalSize	45	41	38	30	20	15	14	11	12	9	235
Parents	22	45	39	9	8	7	8	3	1	0	142
Daughters	24	40	35	15	7	6	5	4	5	1	142

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What is the probability that we would observe this many parents and daughters assuming that chromosome 2 is no different than any other chromosomes?

From this we might hypothesize that we should see more retrogenes on sex chromosomes and possibly more parental genes on sex chromosomes than we would expect by chance.

- 1) Size of the chromosome (determines probability of a daughter gene being on a chromosome)
- 2) Number of genes on each chromosome (determines the probability of a parent gene being on a a chromosomes)

1. First decide what we want to test
 - a) the number of parents on $\lg 2$
 - b) the number of daughters on $\lg 2$
2. Create an object to store a null distribution of these values
3. Simulate an expected number of parents and an expected number of daughters on $\lg 2$.
 - a) when simulating account for chromosome size and gene number
4. Repeat step 3 thousands of times.
5. Compare our observation to the null distribution to calculate a p-value for our observation.

R example