

The likelihood ratio for continuous data

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Summary

This document introduces the likelihood ratio for continuous data and models, and explains its connection with discrete models.

Prerequisites

Be familiar with the [likelihood ratio for discrete data](#).

Definition

Recall that if M_0 and M_1 are fully-specified models for discrete data $X = x$, with probability mass functions $p(x | M_0)$ and $p(x | M_1)$, then the likelihood ratio for M_1 vs. M_0 is

$$\text{LR}(M_1, M_0) := \frac{p(x | M_1)}{p(x | M_0)}.$$

Now suppose that the data and models are continuous. So instead of a probability *mass* function, each model has a probability *density* function. Then the likelihood ratio for M_1 vs. M_0 is usually defined as the ratio of the probability density functions. That is, we have exactly the same expression for the LR,

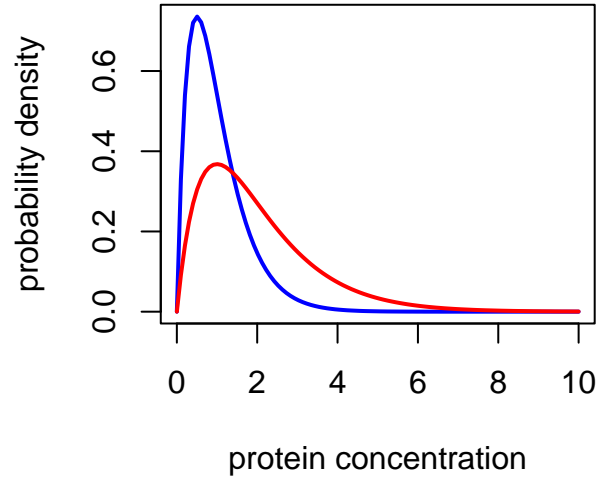
$$\text{LR}(M_1, M_0) := \frac{p(x | M_1)}{p(x | M_0)},$$

but now $p(\cdot | M_1)$ and $p(\cdot | M_0)$ are probability density functions instead of probability mass functions.

Example

A medical screening test for a disease involves measuring the concentration (X) of a protein in the blood. In normal individuals, X has a Gamma distribution with mean 1 and shape 2. In diseased individuals, the protein becomes elevated, and X has a Gamma distribution with mean 2 and shape 2. Plotting the probability density functions of these distributions yields:

```
x <- seq(0,10,length.out = 100)
plot(x,dgamma(x,scale = 0.5,shape = 2),type = "l",col = "blue",lwd = 2,
      xlab = "protein concentration",ylab = "probability density")
lines(x,dgamma(x,scale = 1,shape = 2),type = "l",col = "red",lwd = 2)
```



Suppose that for a particular patient we observe $X = 4.02$. Then the likelihood ratio for the model that this patient is from the normal group (M_n) vs. the model that the patient is from the diseased group (M_d) is $\text{dgamma}(4.02, \text{scale} = 0.5, \text{shape} = 2) / \text{dgamma}(4.02, \text{scale} = 1, \text{shape} = 2)$ which is 0.0718. That is, the data favour this individual being diseased by a factor of approximately 14.

Connection with discrete models

Often the likelihood ratio for continuous models is simply *defined* as the ratio of the densities, as above. However, an alternative approach, which can yield greater insight, is instead to *derive* this result as an approximation, from the definition of likelihood ratio for discrete models, as follows.

The first step is to recognize that in practice all observations are actually discrete, because of finite precision. Sometimes the measurement precision is made explicit, but often it is implicit in the number of decimal places used to report an observation. For example, in the example above, where we were told that we observed a protein concentration of $X = 4.02$, it would be reasonable to think that the measurement precision is 2 decimal places, and that this observation actually corresponds to “ X lies in the interval $[4.015, 4.025]$ ”. The probability of this observation, under a continuous model for X , is the integral of the probability density function from 4.015 to 4.025. In other words, it is $F_X(4.025) - F_X(4.015)$, where F_X denotes the cumulative distribution function for X .

With this view, the likelihood for the “observation” $X = 4.02$ under M_n is actually $\text{pgamma}(4.025, \text{scale} = 0.5, \text{shape} = 2) - \text{pgamma}(4.015, \text{scale} = 0.5, \text{shape} = 2)$, which is 0.00005183. Similarly, the likelihood under M_d is 0.0007217, and therefore the likelihood ratio is 0.0718.

As you can see, this approach yields a LR that is numerically very close to that obtained using the ratio of the densities, as above. This is not a coincidence! Here is why we should expect this to happen more generally. Suppose we assume that the measurement precision is ϵ . So the “observation” $X = x$ really means $X \in [x - \epsilon, x + \epsilon]$. Then the likelihood for a model M , given this observation, is $\Pr(X \in [x - \epsilon, x + \epsilon] \mid M)$. Provided that the density $p(x \mid M)$ is approximately constant in the region within radius ϵ around x , then this probability is approximately $2\epsilon p(x \mid M)$.

Thus, the LR for two models M_1 vs. M_0 is

$$LR = \frac{\Pr(X \in [x - \epsilon, x + \epsilon] \mid M_1)}{\Pr(X \in [x - \epsilon, x + \epsilon] \mid M_0)} \approx \frac{2\epsilon p(x \mid M_1)}{2\epsilon p(x \mid M_0)} = \frac{p(x \mid M_1)}{p(x \mid M_0)}.$$

An example where the approximation breaks down

The approximation usually works well, but here is a simple example to illustrate how the approximation could break down in principle.

Consider observing a single data point X and we compare the models that $M_0 : X \sim N(0, \sigma_0)$ vs. $M_1 : X \sim N(0, \sigma_1)$. Suppose that we observe $X = 0.00$, assumed to be correct to the nearest 0.01. So the “true” LR is given by

```
trueLR <- function (s0, s1) {
  L0 <- pnorm(0.005, sd = s0) - pnorm(-0.005, sd = s0)
  L1 <- pnorm(0.005, sd = s1) - pnorm(-0.005, sd = s1)
  return(L0/L1)
}
```

and the approximation is

```
approxLR <- function (s0, s1)
  dnorm(0, sd = s0)/dnorm(0, sd = s1)
```

Now, if σ_0 and σ_1 are both not too small, the approximation works fine. For example, for $\sigma_0 = 0.5, \sigma_1 = 1$, the truth and approximation are 2 and 2.

But σ_0 or σ_1 is small, we have the problem that the density is not approximately constant within the region $[-0.005, 0.005]$. For example, at $\sigma_0 = 0.001, \sigma_1 = 1$ we have the truth and approximation as 250.6637 and 1000.

Summary

In most cases, the likelihood ratio for model M_1 vs. model M_0 for a continuous random variable X , given an observation $X = x$, can be well approximated by the ratio of the model densities of X , evaluated at x . This approximation comes from assuming that the model density functions are approximately constant within the neighborhood of x that has radius equal to the measurement precision.