## Odds Ratio and Relative Risk

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Last week we considered the transmission disequilibrium test and made note that it controlled for effects of *population stratification* 

As this will come up again in future lectures, it's worth briefly covering now. We'll start with a working definition

Population stratification is the presence of systematic differences in allele frequencies between subpopulations in a population as a results of *non-random mating* between individuals. As such, it is an important confounding variable in genome association studies

Here, we will consider two populations X and Y that together make up a larger population, Z

Ρορι		
AA		
BB	10	
AB	20	

Population Y		
AA	30	
BB	40	
AB	30	

Population Z		
AA	100	
BB	50	
AB	50	

Because of Population X, A has a much higher allele frequency than B in the general population

The primary consequence of the higher allele frequency lies in the relative portions of homozygous parents, either AA or BB

Population Z		
AA	100	
BB	50	
AB	50	

	Non-transmitted		
Transmitted	А	В	Total
A	100	b	
В	с	50	
Total			200

In disease association studies, we control for this confounding by only considering the offspring of those parents who are heterozygous

If there is no association between the disease and allele, we should expect that of the heterozygous parents, 50% would transmit A while 50% would transmit B

Population Z		
AA 100		
BB	50	
AB	50	

	Non-transmitted		
Transmitted	A	В	Total
А	100	25	
В	25	50	
Total			200

The *p*-value for this table using McNemar's test is p = 0.887

If, on the other hand, there is an association between disease status and, say, allele A, we would expect over-representation of A passed from heterozygous parents

Population Z		
AA 100		
BB	50	
AB	50	

	Non-transmitted		
Transmitted	А	В	Total
А	100	40	
В	10	50	
Total			200

The *p*-value for this table using McNemar's test is p = 0.00004

- Last week, we considered the  $\chi^2$  test, Fisher's Exact Test, and the exact binomial test for analyzing association of groups in 2  $\times$  2 tables
- Critically, we understand that these were *binary* tests of association, telling us nothing about the *magnitude* of association (or lack of association) between groups
- Today, we will focus on methods for determining this magnitude

Odds are expressed as a proportion of successes to failures. For example, using a fair six sided die, we would say that the odds of rolling a six are 1 to 5 (one six to five not-sixes), or 0.2.

We could equally say the odds of not rolling a six to rolling a six are 5 to 1, or 5.0. In other words, depending on what is classified as success as failure in a binary outcome gives us two different odds that are inverses of each other

What we may be interested in comparing is the odds of some outcome between two populations

	Side Effects		
Treatment Status	Yes	No	Total
Control	10	40	50
Treatment	30	20	50
Total	40	60	100

For the control group, the odds of having side effects are 10 to 40, or 0.25. Similarly, the odds of having side effects for the treatment group is 30 to 20, or 1.5

From here, two equivalent statements can be made:

	Side Effects			
Treatment Status	Yes	No	Total	Odds
Control	10	40	50	0.25
Treatment	30	20	50	1.5
Total	40	60	100	

We might consider how the odds between Control and Treatment groups are related, and from here, two equivalent statements can be made:

Comparing the odds of side effects from treatment to control, we have an *odds ratio* of 1.5/.25 = 6. That is, the odds of having side effects are 6 times higher in the treatment group, compared to control.

Equivalently, we also could have described the odds from control to treatment as  $.25/1.5\,=\,0.1667$ 

Typically, we compute the odds in a standard way, which divides the odds of the first row by the odds of the second. Of course, which group is on which line is arbitrary

When  $1 < \theta < \infty$ , the odds of the first response are  $\theta$  times higher in the first row than the second. When  $0 < \theta < 1$ , the odds are  $(1/\theta)$  higher in the second row than the first.  $\theta = 1$  if and only if the two groups are independent

The odds ratio remains the same if the columns and rows of the table are switched: in other words,  $\theta$  is not determined by which margins (if any) are assumed to be fixed

$$OR = \hat{\theta} = \frac{a/b}{c/d} = \frac{ad}{bc}$$

## Odds Ratio Test Statistic

Having an estimate for the odds ratio is nice, but otherwise meaningless without having a sense of the variability in this estimate. In other words, it would be convenient if we could compute a confidence interval as well.

Under the null  $H_0: \theta = 1$ , we see that the odds ratio is just as likely to fall in the interval (0, 1] as it is  $[1, \infty)$ . Consequently, the distribution of  $\hat{\theta}$  is hardly normal. It can be shown that this is remedied with a log transformation:

$$\log \hat{ heta} \sim N( heta, \hat{\sigma}(\log \hat{ heta}))$$

where

$$\hat{\sigma}(\log \hat{\theta}) = \left(\frac{1}{a} + \frac{1}{b} + \frac{1}{c} + \frac{1}{d}\right)^{1/2}$$

## Odds Ratio and CLT

Given that  $\log \hat{\theta}$  is approximately normally distributed, we can compute the same style of confidence intervals that we used in the CLT. With  $z_{\alpha/2}$  being our critical value, we can construct a  $(1 - \alpha)$ % confidence interval as

$$\log \hat{ heta} \pm z_{lpha/2}(\hat{\sigma}(\log \hat{ heta}))$$

Taking the exponential of the endpoints of this interval, we end up with a confidence interval for the odds ratio

$$egin{aligned} &\exp\left(\log\hat{ heta}\pm z_{lpha/2}(\hat{\sigma}(\log\hat{ heta}))
ight) = \exp\left(\log\hat{ heta}
ight)\exp\left(\pm z_{lpha/2}(\hat{\sigma}(\log\hat{ heta}))
ight) \ &= \hat{ heta}\exp\left(\pm z_{lpha/2}(\hat{\sigma}(\log\hat{ heta}))
ight) \end{aligned}$$

This estimate tends to be slightly conservative, leading to the possibility of incorrectly rejecting a null hypothesis

## 95 % CI for Odds Ratio

	Side Effects			
Treatment Status	Yes	No	Total	Odds
Control	10	40	50	0.25
Treatment	30	20	50	1.5
Total	40	60	100	

$$\hat{ heta} = 6, \quad \log \hat{ heta} = 1.792, \quad z_{lpha/2} = 1.96$$
 $\hat{\sigma}(\log \hat{ heta}) = \left(rac{1}{10} + rac{1}{40} + rac{1}{30} + rac{1}{20}
ight)^{1/2} = 0.456$ 

Then

$$\log \hat{ heta} \pm z_{lpha/2}(\hat{\sigma}(\log \hat{ heta})) = (0.8972, 2.6864)$$

$$\hat{ heta} \exp\left(\pm z_{lpha/2}(\hat{\sigma}(\log \hat{ heta}))
ight) = (2.4526, 14.6783)$$

A slightly more sophisticated version of this test is returned by the <code>fisher.test</code> in R

```
> fisher.test(mm)
```

Fisher's Exact Test for Count Data

```
data: mm
p-value = 0.000083
alternative hypothesis: true odds ratio is not equal to 1
95 percent confidence interval:
    2.2658 16.3909
sample estimates:
odds ratio
    5.8813
```

A more meaningful statistic for clinicians is often the *relative risk*, which expresses a ratio of probabilities of outcomes between populations based on exposure. For example, we might ask, "What is the proportional increase in heart disease for smokers compared to non-smokers?"

Caution needs to be used here as we are asking a question about conditional probabilities. Computing this directly requires knowing the probability of an outcome *in general*, which we may or may not have, depending on the study

	Variable		
Population	+	_	Total
A	а	b	a+b
В	с	d	c+d
Total	a + c	b+d	N

 $\pi_{1|1} = \{ \text{Probability of '+' outcome, given Population A} \} = \frac{a}{a+b}$ 

 $\pi_{1|2} = \{ \text{Probability of '+' outcome, given Population B} \} = \frac{c}{c+d}$ 

$$RR = \frac{\pi_{1|1}}{\pi_{1|2}} = \frac{a(c+d)}{c(a+b)}$$

A report published in 1988 summarizes results of a Harvard Medical School clinical trial determining effectiveness of aspiring in preventing heart attacks in middle-aged male physicians

	Myocardial Infarction		
Treatment Status	Attack	No Attack	
Placebo	189	10,845	
Asprin	104	10,933	

Here, the row totals were fixed by study design. As the column values were not determined a priori, these observations reflect the probabilities of an outcome in the study

	Myocardial Infarction		
Treatment Status	Attack	No Attack	
Placebo	189	10,845	
Asprin	104	10,933	

 $\pi_{1|1} = \{\text{Probability of '+' outcome, given Population A}\} = \frac{189}{11034}$  $\pi_{1|2} = \{\text{Probability of '+' outcome, given Population B}\} = \frac{104}{11037}$ 

$$RR = \frac{\pi_{1|1}}{\pi_{1|2}} = \frac{a(c+d)}{c(a+b)} = \frac{189 \cdot 11037}{104 \cdot 11039} = 1.812$$

A report published in 1950 summarizes the results of a case-control study between 20 London hospitals investigating relationship between cigarette smoking and cancer. The study involved 1,418 patients, matching 709 cases with lung cancer against 709 controls without lung cancer on the basis of gender and age

	Lung Cancer	
Smoking Status	Present	Not Present
Smoker	688	650
Non-Smoker	21	59
Total	709	709

Now it is the totals for the *outcome* that we determined a priori. From this data, it would be impossible to determine the actual prevelance of lung cancer within the population. Without these probabilities, relative risk cannot be computed

	Variable		
Population	+	_	Total
A	а	b	a+b
В	с	d	c+d
Total	a + c	b+d	N

Fortunately, the odds ratio can be used as an approximation to the relative risk when incidence of the disease is close to zero (i.e., rare diseases). In such cases, relative to b and d, a and c will be small:

$$RR = rac{a(c+d)}{c(a+b)} pprox rac{ad}{bc} = OR$$

Consider this in the Myocardial Infarction study, where the incidence of a heart attack is relatively small

	Myocardial Infarction		
Treatment Status	Attack	No Attack	
Placebo	189	10,845	
Asprin	104	10,933	

$$RR = rac{a(c+d)}{c(a+b)} = 1.812$$

$$OR = \frac{ad}{bc} = 1.832$$

Returning to the smoking example, as the true incidence of lung cancer is known to be relatively small, the odds ratio can still provide a rough indication of relative risk

	Lung Cancer	
Smoking Status	Present	Not Present
Smoker	688	650
Non-Smoker	21	59
Total	709	709

$$OR = \frac{688 \cdot 59}{21 \cdot 650} = 2.974$$