

stir — Report incidence-rate comparison[Syntax](#)[Remarks and examples](#)[Also see](#)[Menu](#)[Stored results](#)[Description](#)[Methods and formulas](#)[Options](#)[Reference](#)

Syntax

stir *exposedvar* [*if*] [*in*] [, *options*]

<i>options</i>	Description
<hr/>	
Main	
<u>strata</u> (<i>varname</i>)	stratify on <i>varname</i>
<u>noshow</u>	do not show st setting information
<hr/>	
Options	
<u>ird</u>	report incidence-rate difference rather than ratio
<u>estandard</u>	combine external weights with within-stratum statistics
<u>istandard</u>	combine internal weights with within-stratum statistics
<u>standard</u> (<i>varname</i>)	combine user-specified weights with within-stratum statistics
<u>pool</u>	display pooled estimate
<u>noc crude</u>	do not display crude estimate
<u>nohom</u>	do not display homogeneity test
<u>tb</u>	calculate test-based confidence intervals
<u>level</u> (#)	set confidence level; default is <code>level(95)</code>

Options except `noshow`, `tb`, and `level(#)` are relevant only if `strata()` is specified.

You must `stset` your data before using `stir`; see [ST] `stset`.

`by` is allowed; see [D] `by`.

`fweights` and `iweights` may be specified using `stset`; see [ST] `stset`. `stir` may not be used with `pweighted` data.

Menu

Statistics > Survival analysis > Summary statistics, tests, and tables > Report incidence-rate comparison

Description

`stir` reports point estimates and confidence intervals for the incidence-rate ratio and difference. `stir` is an interface to the `ir` command; see [ST] `epitab`.

By the logic of `ir`, `exposedvar` should be a 0/1 variable, with 0 meaning unexposed and 1 meaning exposed. `stir`, however, allows any two-valued coding and even allows `exposedvar` to be a string variable.

`stir` may not be used with `pweighted` data.

`stir` can be used with single- or multiple-record or single- or multiple-failure st data.

Options

Main

strata(*varname*) specifies that the calculation be stratified on *varname*, which may be a numeric or string variable. Within-stratum statistics are shown and then combined with Mantel–Haenszel weights.

noshow prevents **stir** from showing the key st variables. This option is seldom used because most people type **stset**, **show** or **stset**, **noshow** to set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] **stset**.

Options

ird, **estandard**, **istandard**, **standard(*varname*)**, **pool**, **nocrude**, and **nohom** are relevant only if **strata()** is specified; see [ST] **epitab**.

tb and **level(#)** are relevant in all cases; see [ST] **epitab**.

Remarks and examples

[stata.com](http://www.stata.com)

stir examines the incidence rate and time at risk.

```
. use http://www.stata-press.com/data/r13/page2
. stir group, noshow
```

note: Exposed <-> group==2 and Unexposed <-> group==1

	group		Total
	Exposed	Unexposed	
Failure Time	19 5023	17 4095	36 9118
Incidence rate	.0037826	.0041514	.0039482
	Point estimate	[95% Conf. Interval]	
Inc. rate diff.	-.0003688	-.002974	.0022364
Inc. rate ratio	.9111616	.4484366	1.866047 (exact)
Prev. frac. ex.	.0888384	-.8660469	.5515634 (exact)
Prev. frac. pop	.04894		
(midp) Pr(k<=19) =		0.3900 (exact)	
(midp) 2*Pr(k<=19) =		0.7799 (exact)	

Stored results

stir stores the following in r():

Scalars

r(p)	one-sided <i>p</i> -value
r(ird)	incidence-rate difference
r(lb_ird)	lower bound of CI for ird
r(ub_ird)	upper bound of CI for ird
r(irr)	incidence-rate ratio
r(lb_irr)	lower bound of CI for irr
r(ub_irr)	upper bound of CI for irr
r(afe)	attributable (prev.) fraction among exposed
r(lb_afe)	lower bound of CI for afe
r(ub_afe)	upper bound of CI for afe
r(afp)	attributable fraction for the population
r(chi2_mh)	Mantel–Haenszel homogeneity χ^2
r(chi2_p)	pooled homogeneity χ^2
r(df)	degrees of freedom

Methods and formulas

stir simply accumulates numbers of failures and time at risk by exposed and unexposed (by strata, if necessary) and passes the calculation to ir; see [ST] **epitab**.

Reference

Dupont, W. D. 2009. *Statistical Modeling for Biomedical Researchers: A Simple Introduction to the Analysis of Complex Data*. 2nd ed. Cambridge: Cambridge University Press.

Also see

[ST] **epitab** — Tables for epidemiologists

[ST] **stset** — Declare data to be survival-time data

[ST] **stsum** — Summarize survival-time data