# Lecture Five: Comparing Multiple Samples: Non-Parametric tests (Cont.)

#### 1. Weighted log rank Tests

• For each time interval (t<sub>(j-1)</sub>, t<sub>(j)</sub>], in which there is only one distinct failure time (allow ties), we have a 2 by 2 table

| Group | # of deaths at $t_{(j)}$ | # of surviving beyond t <sub>(j)</sub> | # at risk just before t <sub>(j)</sub> |
|-------|--------------------------|--|--|
| Ι     | d <sub>1j</sub>          | $n_{1j}$ - $d_{1j}$                    | n <sub>1j</sub>                        |
| II    | d <sub>2j</sub>          | $n_{2j}$ - $d_{2j}$                    | n <sub>2j</sub>                        |
| Total | dj                       | nj-dj                                  | nj                                     |

The expected events:

$$e_{1j} = d_j * n_{1j}/n_j$$
  
 $e_{2j} = d_j * n_{2j}/n_j$ 

 $d_{1j}|d_j$  has hypergeometric distribution with

$$E(d_{1j}|d_j) = e_{1j}$$
  
Var(d\_{1j}|d\_j) =  $v_{1j} = \frac{n_{1j}n_{2j}d_j(n_j - d_j)}{n_j^2(n_j - 1)}$ 

• A family of weighted log rank statistics

$$U_{WT} = \sum_{j=1}^{r} w_j (d_{1j} - e_{1j})$$

• A general weighting scheme (the Peto-Peto statistic when  $\rho = 1$  and  $\gamma = 0$ ).

$$w_j = S(t_j)^{\rho} (1 - S(t_j))^{\gamma}$$

Here,  $\rho \ge 0$ ,  $\gamma \ge 0$ ,  $S(t_j)$  is the KM estimate pooled from both groups.

- o Effects of weights
  - $\triangleright \rho = 0$  and  $\gamma = 0$ : equal weight
  - >  $\rho > 0$  and  $\gamma > 0$ : more weight on difference in the middle
  - >  $\rho > 0$  and  $\gamma = 0$ : more weight on earlier difference
  - $\triangleright \rho = 0$  and  $\gamma > 0$ : more weight on later difference



- Splus implementation :  $\gamma = 0$  and  $w(t) = S(t)^{\rho}$  (Ref.: *Biometrika* vol. 69, pp. 553-566 (1982) by Harrington and Fleming)
  - ✤ Splus function: survdiff()
  - $\rho = 0$ : w(t) = 1, log-rank/Mental-Haenszel
  - $\rho = 1$ : w(t) = S(t), Peto-Peto/Prentice (generalized Wilcoxon)
  - ✤  $\rho > 0$ : more weight on earlier difference (S(t) is non-deceasing function)
  - ♦ ρ < 0: more weight on later difference (interpretation less natural)
- SAS implementation: strata statement (test option)

# • Other weighted rank based Tests

 $w_j = n_j$  (The Gehan (1965) statistic)  $w_j = S(t_j)$  (The Peto-Peto (1972) statistic)  $w_j = n_j^{1/2}$  (One of Tarone and Ware (1977) test statistics)

• The Wilcoxon Test

$$U_W = \sum_{j=1}^r n_j (d_{1j} - e_{1j})$$

The variance of the Wilcoxon statistic above is

$$V_w = \sum_{j=1}^r n_j^2 v_{1j}$$

and the Wilcoxon test statistic is

$$W_W = U_W^2 / V_W \sim \chi^2(1),$$

when the null hypothesis is true (why?).

- SAS implementation: see options of *strata statement* of PROC LIFETEST.
- Example 2.13: Wilcoxon test (see output for example 2.12)
- Comparison of the log rank, Wilcoxon and Peto-Peto tests
  - Equal weight (detect difference that is consistent over time) for log rank test, more weight on the earlier difference for Wilcoxon test.
  - Log rank: more suitable when assumption of proportional hazards is satisfied  $(h_1(t) = \varphi h_2(t))$
  - Necessary (not a sufficient) condition for proportional hazards: The true survivor functions do not cross  $(S_1(t) = [S_2(t)]^{\varphi})$
  - Example 2.14: KM plot

### 2. Comparison of more than two samples

- Same idea as in two group case: measuring discrepancy
- Kruskal-Wallis tests (more general than Wilcoxon tests)
- log-rank tests based on sequence of 2 by g tables (g > 2)

$$U_{Lk} = \sum_{j=1}^{r} (d_{kj} - \frac{n_{kj}d_j}{n_j}), \text{ [Wilcoxon test: } U_{Wk} = \sum_{j=1}^{r} n_j (d_{kj} - \frac{n_{kj}d_j}{n_j}) \text{]}$$

for k = 1, 2, ..., g-1. The variance matrix for log-rank test is

$$V_L = (V_{Lkk'}),$$

where

$$V_{Ikk} = \sum_{j=1}^{r} \frac{n_{kj} d_j (n_j - d_j)}{n_j (n_j - 1)} (\delta_{kk} - \frac{n_{kj}}{n_j}).$$

• The test statistic:  $U_L V_L^{-1} U_L \sim \chi^2(g-1)$  (why?)

#### 3. Further Generalizations

# • Stratification within a treatment group is necessary when subjects are not homogenous: Section 2.8

- Handle additional covariates (confounding variables).
- Example: Multi-center clinical trial (stratified by center); stratified by sex or other potential risk factors.
- Stratified log-rank/Wilcoxon test: Basically, Calculating the values of U- and V-statistics for each stratum, then combine them (see following test statistic).
- Test statistic

$$W_{S} = \frac{\left(\sum_{k=1}^{s} U_{lk}\right)^{2}}{\sum_{k=1}^{s} V_{lk}} \sim \chi^{2}(1)$$

• Example 2.15: Two vaccines after surgery for melanoma patients Summarized output from following SAS program:

| Age group             | $U_L$          | $V_L$                  | $W_L(\frac{U_L^2}{V_L})$      |
|-----------------------|----------------|------------------------|-------------------------------|
| 21-40                 | -0.2571        | 1.1921                 | 0.055                         |
| 41-60                 | 0.4778         | 0.3828                 | 0.596                         |
| 61-                   | 1.0167         | 0.6497                 | 1.591                         |
| Total                 | 1.2374         | 2.2246                 |                               |
| $W_S = 1.2374^2/2.22$ | 246=0.688. Tes | t statistic $W_S \sim$ | $\chi^2(1)$ . P-value = 0.41. |

```
/* SAS program: melanoma.sas (SAS Version 8) */
```

```
options pagesize=60 linesize=79 nodate nonumber;
libname fu '../../sdata';
data fu.melanoma;
infile '../../data/melanoma.dat';
input age tx survt censor;
data w1;
     set fu.melanoma;
if age = 1;
proc lifetest notable;
     time survt*censor(0);
     strata tx;
data w2;
     set fu.melanoma;
if age = 2;
proc lifetest notable;
     time survt*censor(0);
     strata tx;
data w3;
```

```
set fu.melanoma;
if age = 3;
proc lifetest notable;
    time survt*censor(0);
    strata tx;
run;
/* SAS program: melanoma.sas (SAS Version 9) */
options pagesize=60 linesize=79 nodate nonumber;
libname fu '../../sdata';
data w;
    set fu.melanoma;
proc lifetest notable;
    time survt*censor(0);
strata age / group = tx;
run;
```

• It's not flexible as Cox model (proportional hazards model).

### • When treatment groups are ordered in some way: Log-rank test for trend

- Examples: groups correspond to increasing doses of a treatment; the stage of a disease, or age group.
- Log-rank test may not lead to a significant difference among groups even though the hazard of death increase or decrease across the groups
- Mathematically, the alternative hypothesis is

$$H_A : S_1(t) < S_2(t) < \dots < S_g(t)$$

• Log-rank test for trend statistic:

$$U_T = \sum_{k=1}^{g} w_k (d_{k.} - e_{k.}),$$

where  $w_k$  is a code assigned to the k'th group, k = 1, 2, ..., g and

$$d_{k.} = \sum_{j=1}^{r_k} d_{kj}, \quad e_{k.} = \sum_{j=1}^{r_k} e_{kj}$$

are the observed and expected numbers of deaths in the k'th group. The variance of  $U_T$  is given by

$$V_T = \sum_{k=1}^{g} (w_k - \overline{w})^2 e_{k.},$$

where

$$\overline{w} = \frac{\sum_{k=1}^{g} w_k e_{k.}}{\sum_{k=1}^{g} e_{k.}},$$

Then, the statistic  $W_T = U_T^2 / V_T \sim \chi^2(1)$  under  $H_0: S_1(t) = S_2(t) = \dots = S_g(t)$ 

Example 2.16: Melanoma patients (BCG arm only: trend over age?) (page 46)

SAS output:

### Trend Tests

| Test     | Test<br>Statistic | Standard<br>Error | z-Score | Pr >  z |
|----------|-------------------|-------------------|---------|---------|
| Log-Rank | 2.5692            | 1.5465            | 1.6613  | 0.0967  |
| Wilcoxon | 25.0000 1         | 4.4568            | 1.7293  | 0.0838  |

# SAS program:

options pagesize=60 linesize=79 nodate nonumber; libname fu '../../sdata'; data w; set fu.melanoma; if tx = 1; proc lifetest notable; time survt\*censor(0); strata age / trend;

• More flexible approach: Cox model (next chapter)

Renyi type of test (Similar to Kolmogorov-Smirnov test, but with censored data)

See pages 223-224 of Klein & Moeschberger's book (reference #1 in the syllabus).

**Reading assignment: Read section 2.10**