

# BIOST537\_Project

Dante Ramirez, Nada Ali, Jiayu Sun, Machi Kaneko

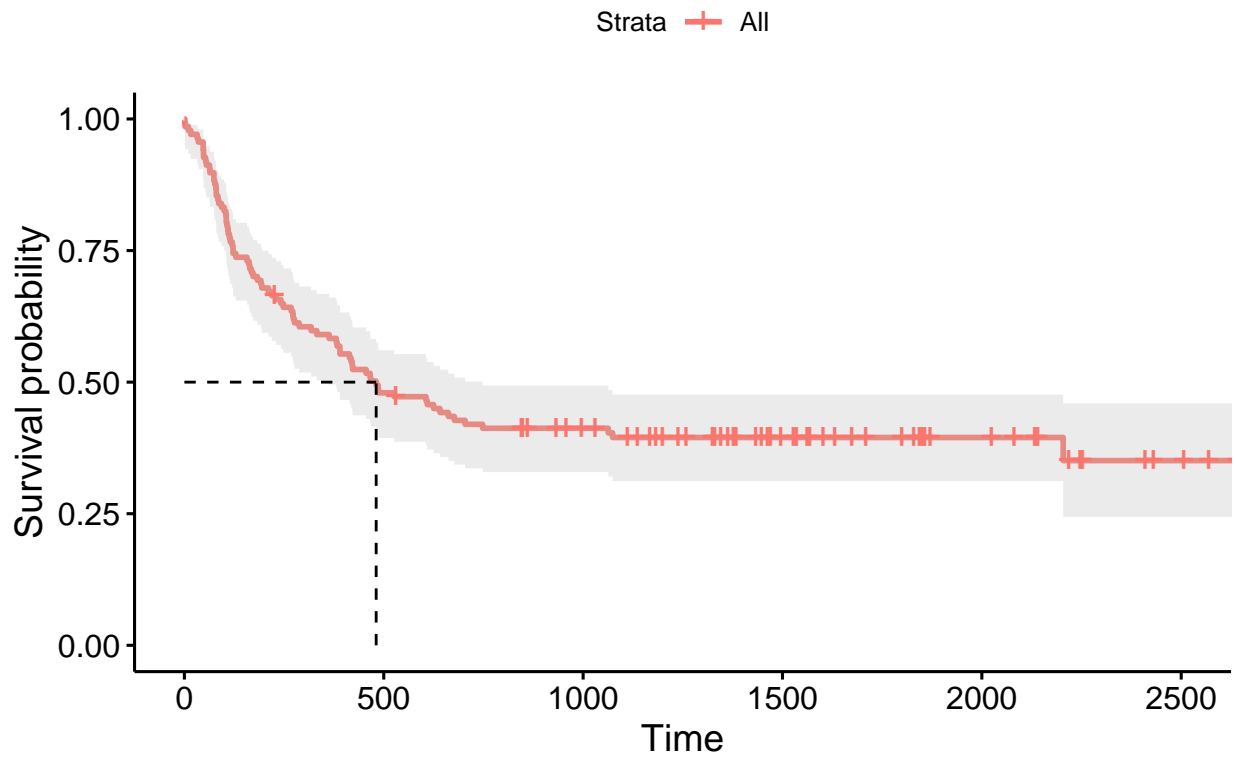
2023-03-13

Directive 1

```
## [1] 782.0292
```

```
## [1] 0.3941606
```

## Kaplan–Meier survival estimate



```
## records      n.max    n.start    events    rmean se(rmean)    median    0.95LCL
## 137.0000  137.0000  137.0000    83.0000 1186.1053  100.5981   481.0000   363.0000
## 0.95UCL
## 748.0000
```

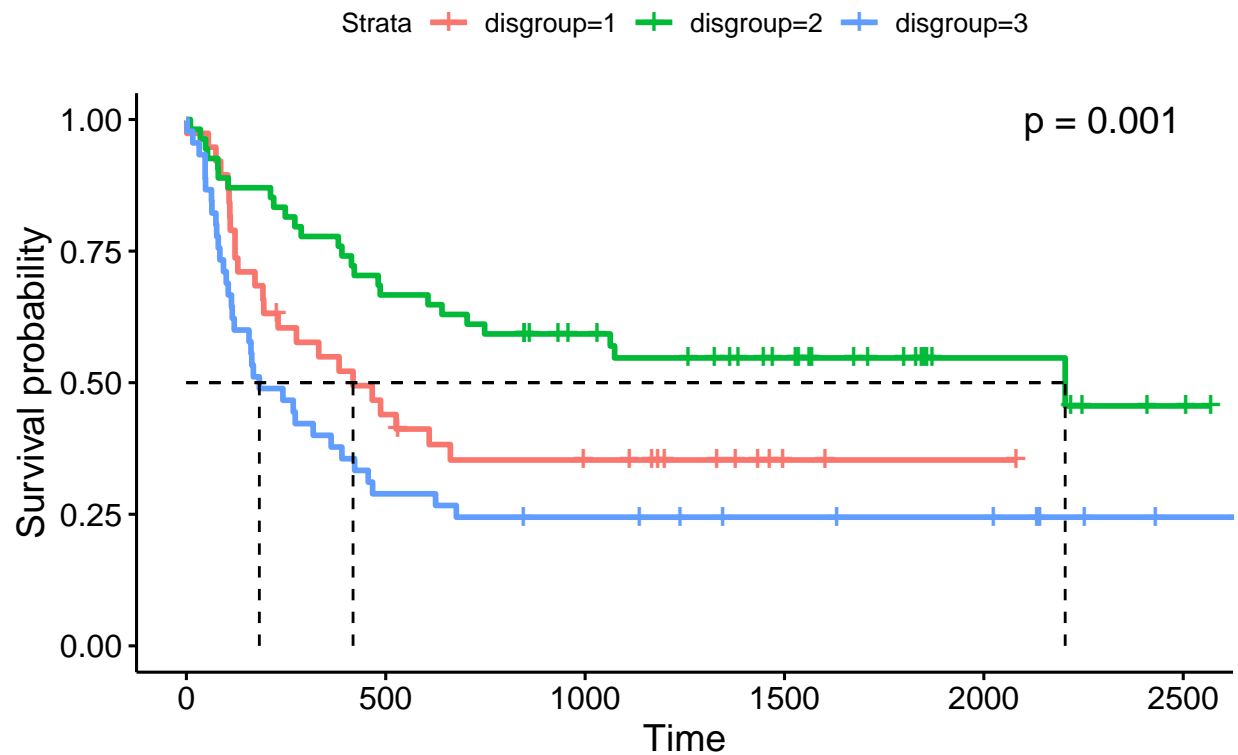
Directive 2

# Disease Subgrouping

Table 1: Call: `s_bmt ~ disgroup` Chisq = 13.803722 on 2 degrees of freedom,  $p = 0.001006$

	N	Observed	Expected	(O-E)^2/E	(O-E)^2/V
<b>disgroup=1</b>	38	24	21.85	0.2112	0.2893
<b>disgroup=2</b>	54	25	39.97	5.604	11.01
<b>disgroup=3</b>	45	34	21.18	7.756	10.53

## Kaplan–Meier survival estimate, by Disease Group



	Disease Group 1	Disease Group 2	Disease Group 3
mean_age	24.421	29.407	30.444
sd_age	7.295	8.764	11.220
count_males	26.000	30.000	24.000
prop_males	0.684	0.556	0.533
count_females	12.000	24.000	21.000
prop_females	0.316	0.444	0.467
count_cmv	15.000	26.000	27.000
prop_cmv	0.395	0.481	0.600
count_mtx	17.000	12.000	11.000
prop_mtx	0.447	0.222	0.244
count_hospital	64.000	118.000	81.000
mean_donor_age	26.789	28.074	29.933
sd_donor_age	8.933	9.245	12.057
count_donor_males	26.000	34.000	28.000
prop_donor_males	0.684	0.630	0.622
count_donor_cmv	17.000	22.000	19.000
prop_donor_cmv	0.447	0.407	0.422

```
##          chiSq df pChisq
## 1          13.8037  2      5
## n          16.2407  2      1
## sqrtN      15.6529  2      4
## S1         15.7260  2      3
## S2         15.7781  2      2
## FH_p=1_q=1  9.9331  2      6
## $tft
##          Q          Var          Z pNorm
## 1         -10.6695      42.7801 -1.63127      5
## n        -1294.0000  439987.8847 -1.95081      1
## sqrtN     -118.1769   4202.2583 -1.82302      4
## S1         -9.2667     23.2023 -1.92379      3
## S2         -9.1996     22.7588 -1.92839      2
## FH_p=1_q=1  -1.0948      1.4957 -0.89516      6
##
## $scores
## [1] 1 2 3

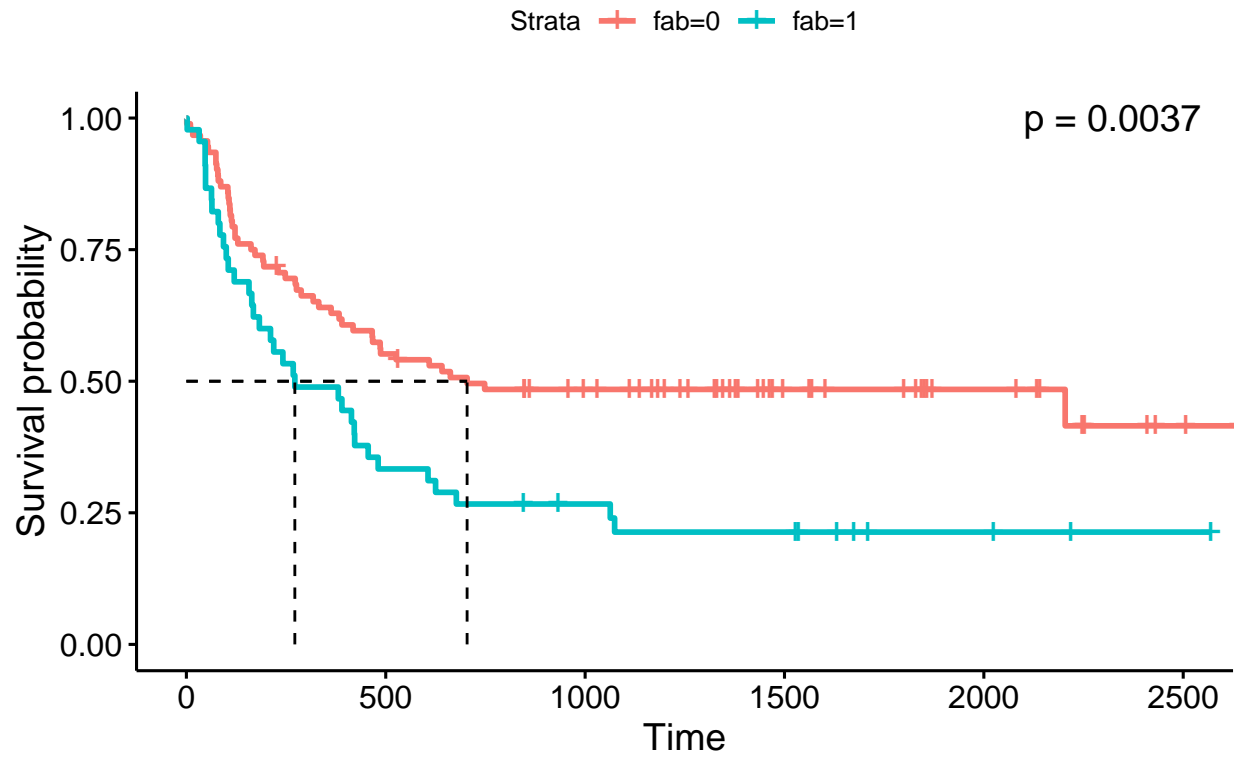
## [1] 0.05107965
```

### *FAB Subgrouping*

Table 2: Call: `s_bmt ~ fab` Chisq = 8.435337 on 1 degrees of freedom, p = 0.003680

	N	Observed	Expected	(O-E)^2/E	(O-E)^2/V
<b>fab=0</b>	92	48	59.83	2.337	8.435
<b>fab=1</b>	45	35	23.17	6.034	8.435

## Kaplan–Meier survival estimate, by FAB Group



	FAB Classification 1	FAB Classification 2
mean_age	28.598	27.889
sd_age	9.478	9.810
count_males	56.000	24.000
prop_males	0.609	0.533
count_females	36.000	21.000
prop_females	0.391	0.467
count_cmv	44.000	24.000
prop_cmv	0.478	0.533
count_mtx	32.000	8.000
prop_mtx	0.348	0.178
count_hospital	178.000	85.000
mean_donor_age	29.000	26.956
sd_donor_age	9.669	11.133
count_donor_males	58.000	30.000
prop_donor_males	0.630	0.667
count_donor_cmv	44.000	14.000
prop_donor_cmv	0.478	0.311

##	Q	Var	Z	pNorm
## 1	1.1825e+01	1.6590e+01	2.9033	1
## n	1.0830e+03	1.6628e+05	2.6559	6
## sqrtN	1.1217e+02	1.6047e+03	2.8001	2
## S1	7.9035e+00	8.7832e+00	2.6668	4
## S2	7.8227e+00	8.6118e+00	2.6657	5
## FH_p=1_q=1	2.1652e+00	6.0024e-01	2.7948	3

```
##          maxAbsZ      Var      Q pSupBr
## 1      1.2047e+01 1.6590e+01 2.9578      1
## n      1.0850e+03 1.6628e+05 2.6608      6
## sqrtN    1.1283e+02 1.6047e+03 2.8167      3
## S1      7.9834e+00 8.7832e+00 2.6938      4
## S2      7.8946e+00 8.6118e+00 2.6902      5
## FH_p=1_q=1 2.2184e+00 6.0024e-01 2.8633      2

## [1] 0.007909707
```

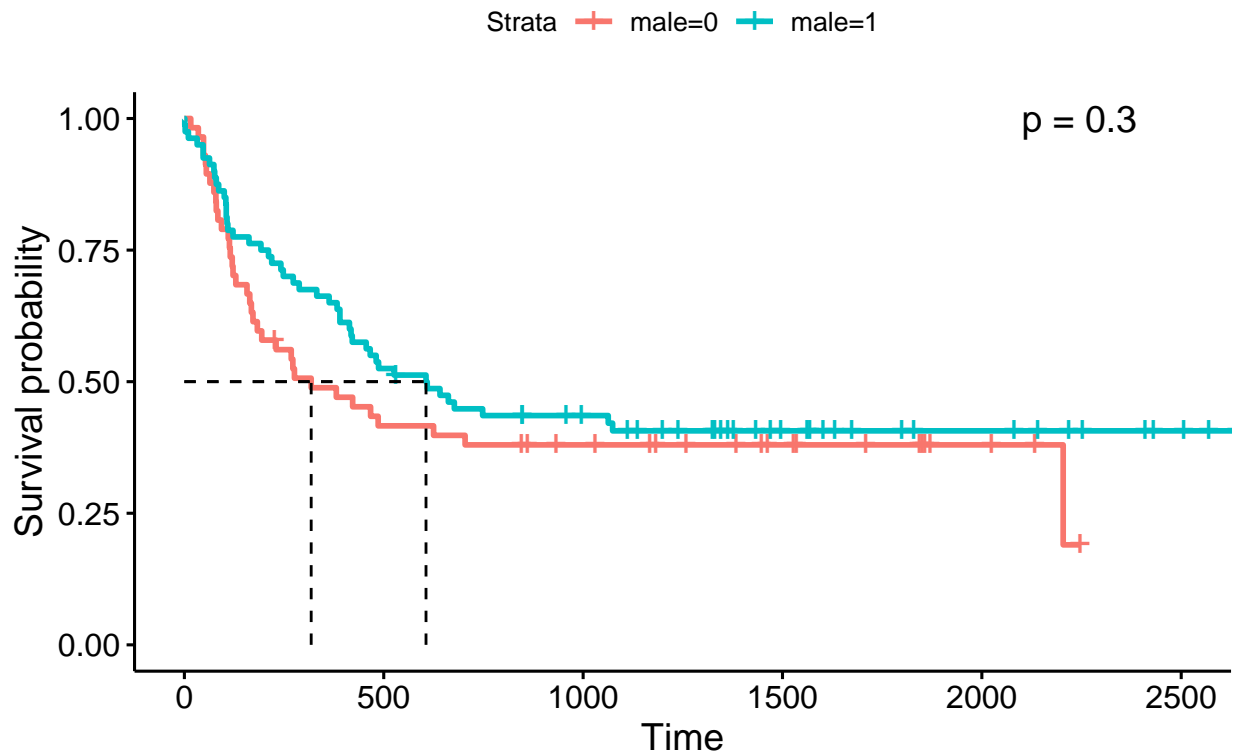
### Directive 3

*Sex Subgrouping*

Table 3: Call: `s_bmt ~ male` Chisq = 1.078766 on 1 degrees of freedom,  $p = 0.298974$

	N	Observed	Expected	(O-E)^2/E	(O-E)^2/V
male=0	57	36	31.42	0.6662	1.079
male=1	80	47	51.58	0.4059	1.079

### Kaplan–Meier survival estimate, by Sex

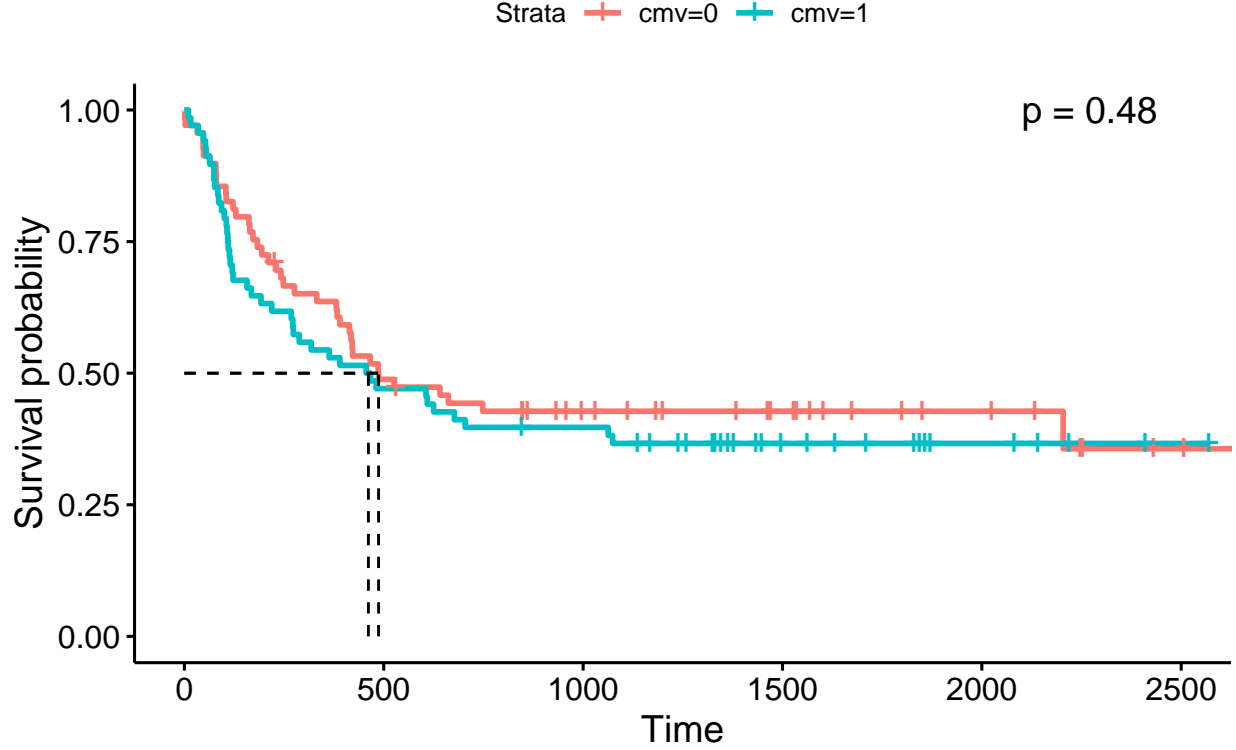


*CMV Subgrouping*

Table 4: Call: `s_bmt ~ cmv` Chisq = 0.497423 on 1 degrees of freedom,  $p = 0.480635$

	N	Observed	Expected	$(O-E)^2/E$	$(O-E)^2/V$
<b>cmv=0</b>	69	40	43.2	0.2375	0.4974
<b>cmv=1</b>	68	43	39.8	0.2579	0.4974

## Kaplan–Meier survival estimate, by CMV

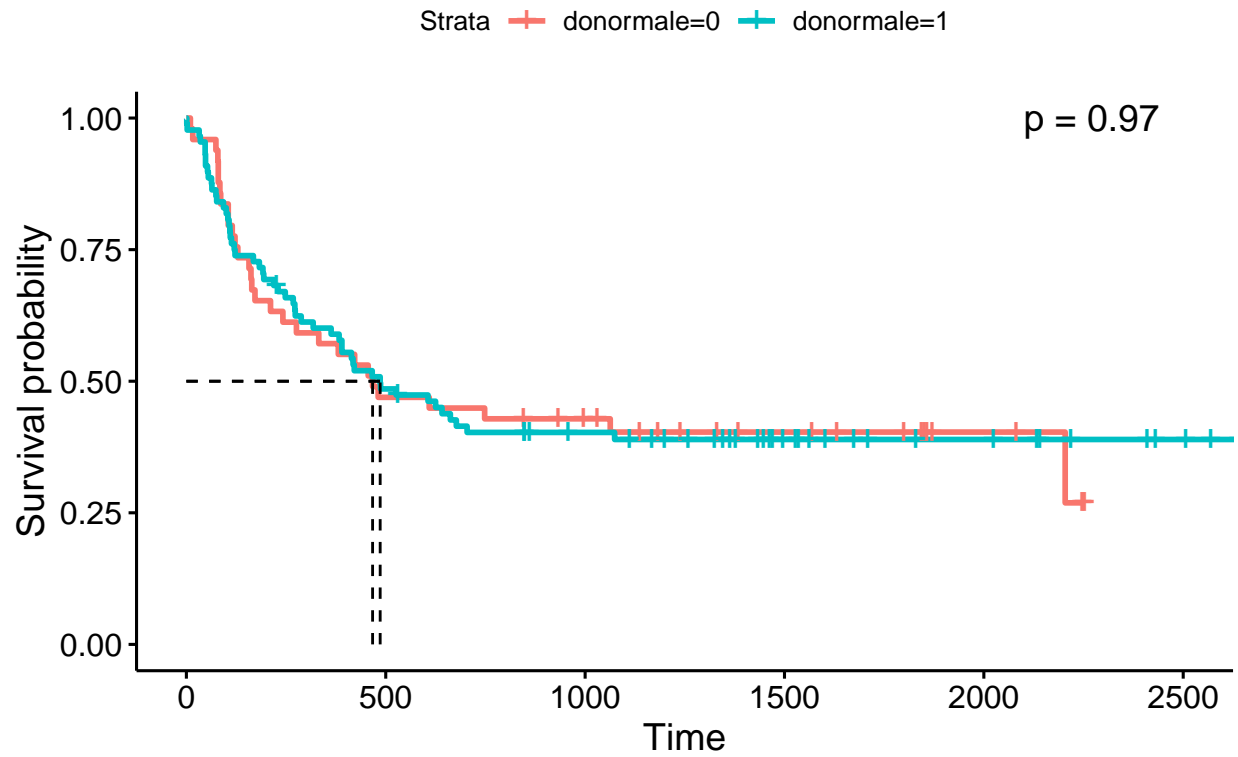


### Donor Sex Subgrouping

Table 5: Call: `s_bmt ~ donormale` Chisq = 0.001359 on 1 degrees of freedom,  $p = 0.970591$

	N	Observed	Expected	$(O-E)^2/E$	$(O-E)^2/V$
<b>donormale=0</b>	49	30	29.84	0.0008686	0.001359
<b>donormale=1</b>	88	53	53.16	0.0004875	0.001359

## Kaplan–Meier survival estimate, by Donor Sex

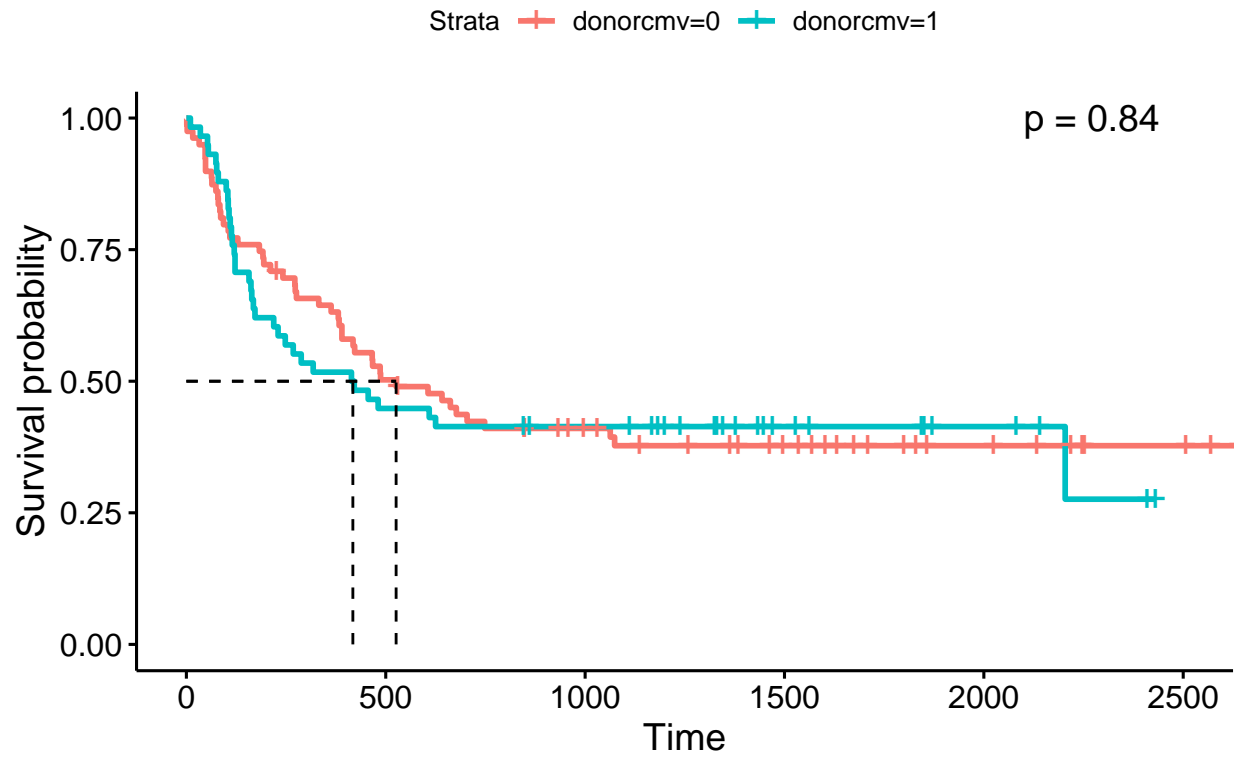


### Donor CMV Subgrouping

Table 6: Call: `s_bmt ~ donorcmv` Chisq = 0.043347 on 1 degrees of freedom,  $p = 0.835073$

	N	Observed	Expected	$(O-E)^2/E$	$(O-E)^2/V$
<b>donorcmv=0</b>	79	48	48.93	0.01772	0.04335
<b>donorcmv=1</b>	58	35	34.07	0.02544	0.04335

## Kaplan–Meier survival estimate, by Donor CMV



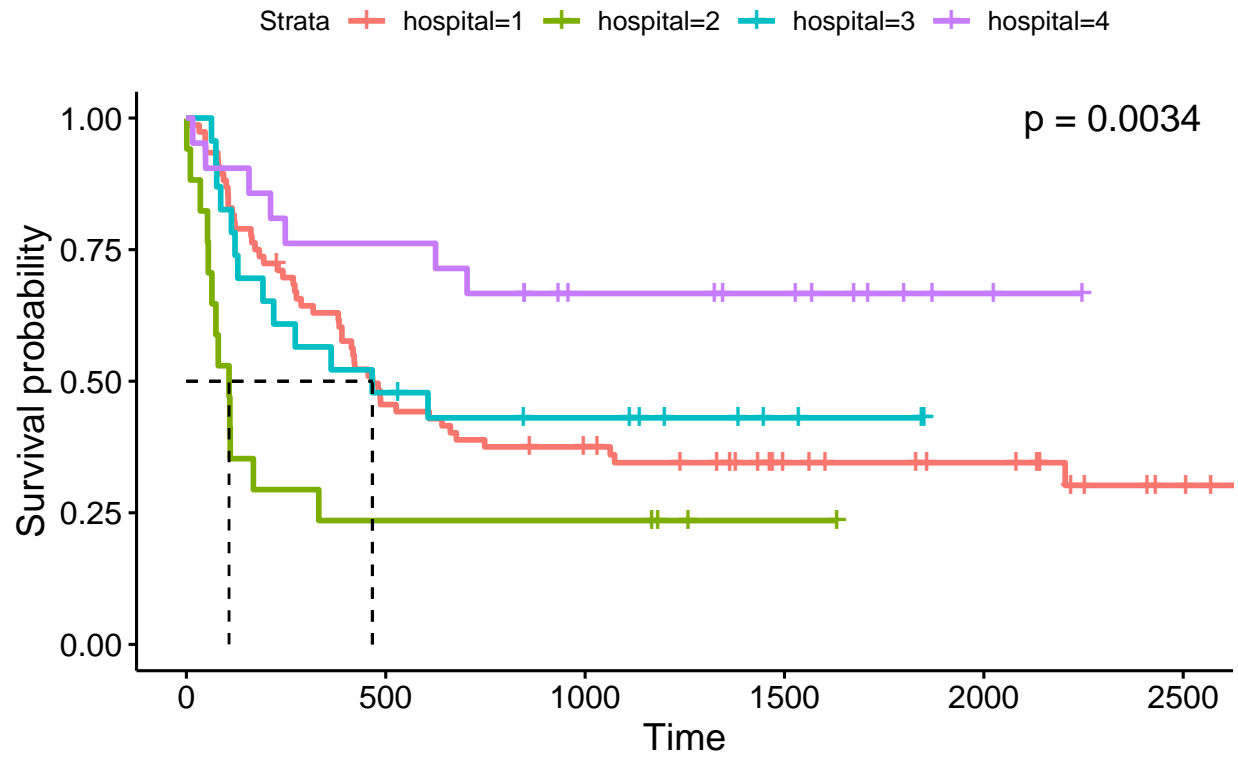
### *Hospital Subgrouping*

Table 7: Call: `s_bmt ~ hospital` Chisq = 13.680494 on 3 degrees of freedom,  $p = 0.003374$

	N	Observed	Expected	(O-E) <sup>2</sup> /E	(O-E) <sup>2</sup> /V
<b>hospital=1</b>	76	50	47.71	0.1101	0.2613
<b>hospital=2</b>	17	13	5.905	8.524	9.258
<b>hospital=3</b>	23	13	13.62	0.02779	0.03339
<b>hospital=4</b>	21	7	15.77	4.879	6.076



## Kaplan–Meier survival estimate, by Hospital

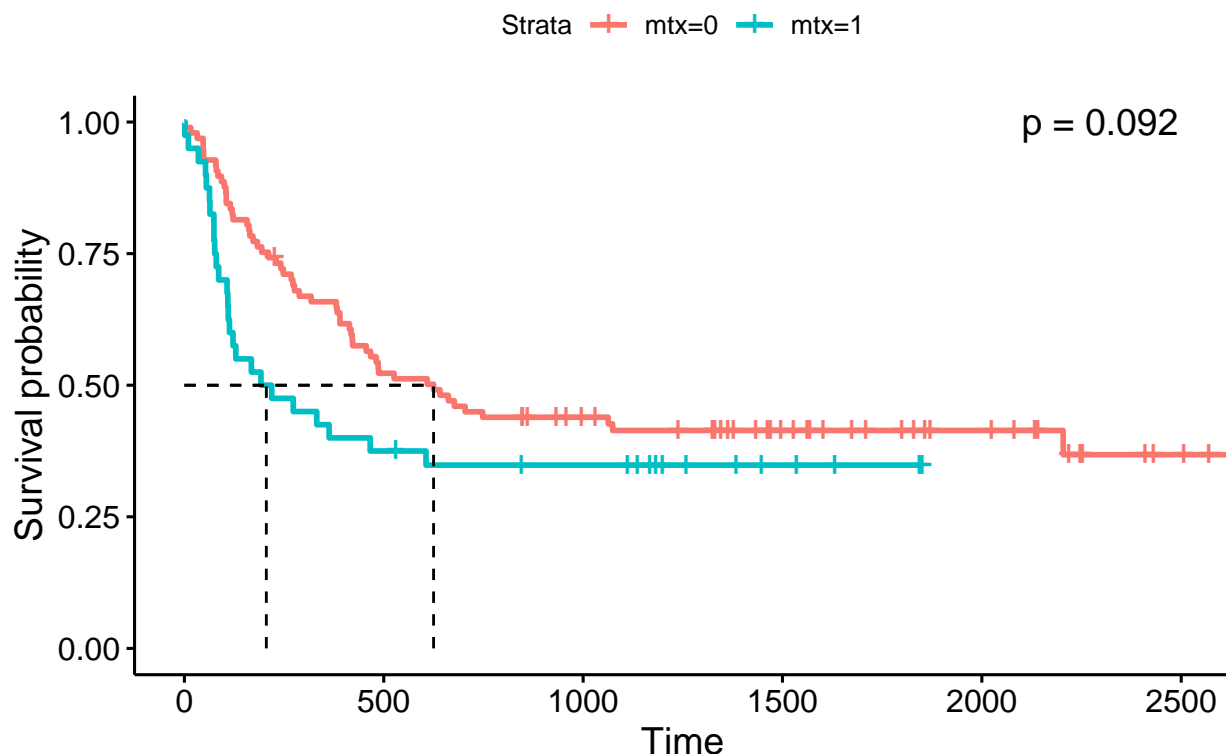


### MTX Subgrouping

Table 8: Call: `s_bmt ~ mtb` Chisq = 2.838053 on 1 degrees of freedom,  $p = 0.092056$

	N	Observed	Expected	$(O-E)^2/E$	$(O-E)^2/V$
<b>mtx=0</b>	97	57	63.48	0.6614	2.838
<b>mtx=1</b>	40	26	19.52	2.151	2.838

## Kaplan–Meier survival estimate, by MTX



Two of the hospitals appear to be significant, but since each has little data, the power might be low, the SE might be high, and the CI may cross.

```
#nonparametric survival function survfit.bmt <- survfit(s.bmt~1, data=bmt, conf.type="log-log")#1.
estimate median disease free survival time print(survfit.bmt)#2How do patients in different disease
groups or in different FAB classifications compare to each other with respect to other available baseline
measurements? #table 1: columns = disease groups, rows= baseline characteristics #table 2: columns =
FAB classifications, rows = baseline characteristics#3 Are any of the measured baseline variables associated
with differences in disease-free survival?#4 Is occurrence of aGVHD after transplantation associated with
improved disease-free survival? summary(coxph(s.bmt ~deltaaa + age + cmv + donorcmv + strata(hospital),
data=bmt))#Is it associated with a decreased risk of relapse? summary(coxph(s.relapse~deltaaa + age
+ cmv + donorcmv, data=bmt))#5 Among the patients who develop aGVHD, are any of the measured
baseline factors associated with differences in disease-free survival? gvhd <- survfit(s.gvhd ~ mtx, data =
bmt, conf.type = "log-log") plot(gvhd, conf.int = F, main="Kaplan-Meier GVHD survival estimate, by
MTX", xlab="Time (in days)", ylab="Survival probability", col="black", lty="solid", lwd=2)gvhdcmv <-
survfit(s.gvhd ~ cmv, data = bmt, conf.type = "log-log") plot(gvhdcmv, conf.int = F, main="Kaplan-Meier
GVHD survival estimate, by recipient CMV status", xlab="Time (in days)", ylab="Survival probability",
col="black", lty="solid", lwd=2)gvhdhospital <- survfit(s.gvhd ~ hospital, data = bmt, conf.type =
"log-log") plot(gvhdhospital, conf.int = F, main="Kaplan-Meier GVHD survival estimate, by hospital",
xlab="Time (in days)", ylab="Survival probability", col="black", lty="solid", lwd=2)gvhddonorcmv
<- survfit(s.gvhd ~ donorcmv, data = bmt, conf.type = "log-log") plot(gvhddonorcmv, conf.int =
F, main="Kaplan-Meier GVHD survival estimate, by donor CMV status", xlab="Time (in days)",
ylab="Survival probability", col="black", lty="solid", lwd=2)#6 Is prophylactic use of methotrexate asso-
ciated with an increased or decreased risk of developing aGVHD? **incude confounders s.gvhd <- with(bmt,
Surv(ta, deltaa==1)) summary(coxph(s.gvhd~mtx + donorcmv + strata(hospital), data=bmt))#Provide
an estimate of the survival function of time from transplant until onset of aGVHD separatefor patients
either administered methotrexate or not. In doing so, consider the importance of accounting for relevant
```

```

confounding factors. s.gvhdm <- with(bmt, Surv(ta, deltaa==1)) survfit.gvhdmx <- survfit(s.gvhdm~mtx,
data=bmt, conf.type="log-log" ) plot(survfit.gvhdmx) summary(survfit.gvhdmx, times=c(7, 14, 21, 28,
35, 42, 49, 56))#7 Is recovery of normal platelet levels associated with improved disease-free survival? -
yes summary(coxph(s.bmt ~deltap + age + donorcmv + strata(hospital), data=bmt))#Is it associated
with a decreased risk of relapse? - no s.relapse <- with(bmt, Surv(agediagnosis, ageevent, deltar==1))
summary(coxph(s.relapse~deltap + age + donorcmv + strata(hospital), data=bmt)

```