

Using Multiple Imputation, Survival Analysis, And Propensity Score Analysis In Cancer Data With Missingness

Master's Thesis

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1 Introduction

- The Problem
- Missing data
- Survival Analysis
- Causal Analysis

2 Methods

- Imputation
- Survival
- Causal Analysis

3 Application

- Breast Brain Mets Example

4 Conclusion

- Discussion

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In an ideal world

- We would have a large dataset
 - That was obtained from a randomized controlled trial (RCT)
 - That would help answer a clearly defined question
 - That had all the covariates of scientific interest
 - That contained no missing data

- RCT's are expensive and often unethical
 - We often get retrospective observational data
 - Pulled from a database or historical records
- The questions we have may not be answerable from the data on hand
 - The data obtained often doesn't support the original question in mind
- The covariates collected are out of our control
 - Since often no control of experiment, no control over what is collected
- Lots of missing data
 - Since no control over how the data is collected, we can't guarantee that everything is collected
 - This issue is seemingly omnipresent in all types of data collection

YES!

- Without an RCT, we can't be sure if differences in outcome is due to the treatment or something else
- Omitting important factors may bias our results
- With missing data, we will be throwing away data and biasing our results

The Solution

This thesis aims to fix some of these problems

- Fill in missing data via multiple imputation
- Create meaningful analytical models via survival analysis
- Get a causal interpretation from observational data

Goal: To be able to apply methods to cancer data

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What is missing data

- Missing data happens when we intend to collect a piece of data but don't actually get it
- Historical approaches
 - Complete Case (CC) analysis: Throw away any record that is not complete
 - Available Case (AC) analysis: Use records so long as they are complete for the specific analysis in question

Definition

The English verb “to impute” comes from the Latin *imputo*, which means to reckon, attribute, make account of, charge, ascribe. [1]

- In the 1930's, Allan, Wishart, and Yates laid framework for missing data
 - Idea: Fill in the missing value, deduct degrees of freedom to account for it
 - Issue: Dogmatic, and variance can't be estimates correctly

Multiple Imputation

Throughout the 70's and 80's Donald Rubin worked to improve on this

- Instead of imputing one value, lets impute it $m \geq 2$ times
- Draw the values from the missing data's posterior distribution given the observed data and the process that generated the missing data

This idea is called Multiple Imputation (MI) and was formalized in 1987 [2]. It is the gold standard method for missing data currently.

How does MI work?

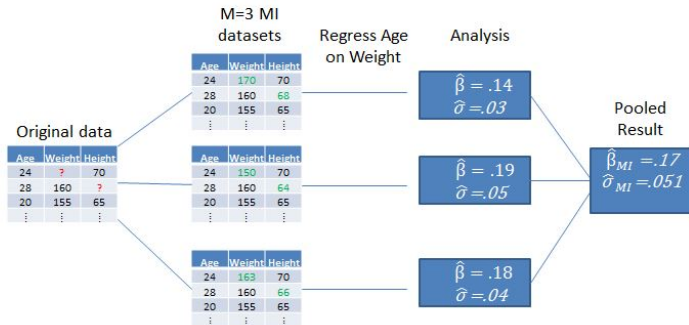


Figure 1: Visualization of MI data

Missingness is displayed by ?'s and the imputed data is shown as #'s. We then regress age on weight, get the results from the individual datasets, and then pool them together.

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Survival Analysis

Survival analysis is a field of statistics concerned with analyzing time to event data, often in the face of censoring or truncation.

Example:

- The survival of patients after a liver transplant in a hospital
- Censoring/Truncation:
 - study ending and no death,
 - subject dies before study starts
 - subject moves away,
 - exact death time only known in an interval

Kaplan-Meier Estimator

- The survival function $S(t) = P(T > t) = \int_t^\infty f(u)du$ is estimated by the nonparametric Kaplan-Meier Estimator

$$\hat{S}(t) = \prod_{t_i < t} \frac{n_i - d_i}{n_i}$$

- n_i is the number of subject in the risk set at time t_i
- d_i is the number of deaths at time t_i

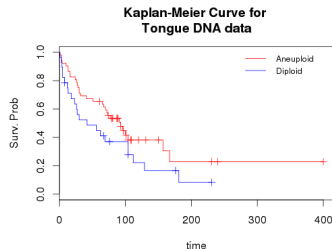


Figure 2: An example of a Kaplan-Meier curve

Log rank test

H_0 : No difference between the survival curves of the two populations

$$\frac{\sum_{j=1}^J (O_{1j} - E_{1j})}{\sqrt{\sum_{j=1}^J V_j}} \sim N(0, 1)$$

- $N_j = N_{1j} + N_{2j}$ is the number at risk at time j (composed from deaths in each group)
- $O_j = O_{1j} + O_{2j}$ is the observed number of deaths at time j (composed from the observed deaths in each group)
- $E_{1j} = \frac{O_j N_{1j}}{N_j}$
- $V_j = \frac{O_j(N_{1j}/N_j)(1-N_{1j}/N_j)(N_j-O_j)}{N_j-1}$

Hazard Function

- Hazard is the instantaneous rate of event given that you have survived until time t , given by

$$h(t) = \lim_{\Delta t \rightarrow 0+} \frac{P[t \leq T < t + \Delta t | T \geq t]}{\Delta t}$$

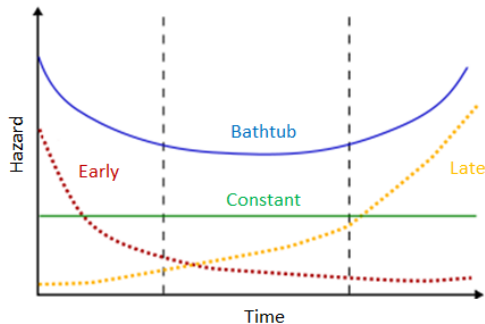


Figure 3: A few different hazard function shapes [3]

Cox Regression

- Cox regression models hazard by

$$h(t|Z) = \underbrace{h_0(t)}_{\text{time}} * \underbrace{\exp(\sum_{k=1}^p \beta_k Z_k)}_{\text{covariates}}$$

- Where $h_0(t)$ is the baseline hazard
- Z_k is the k^{th} covariate
- β_k 's are found by maximizing the partial likelihood function
- The covariates act to multiply the hazard function.

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Example: Examining weight loss between new weight loss drug or placebo

- We would like to be able to say “The drug leads to more weight loss”
 - But need an RCT to say this
 - Randomization minimizes differences between groups at baseline
 - We only have observational data
 - Thus differences could be attributed to the drug or confounding
 - e.g. healthier people were much more likely to take the drug at baseline

Idea: Try to balance the covariates to reduce the effects of confounding so the two groups seem identical at baseline

- Suppose that for or every person, there are two potential outcomes
 - $Y_i(0)$ - The outcome if they had taken the control, $T = 0$
 - $Y_i(1)$ - The outcome if they had taken the treatment, $T = 1$
- The observed value for subject i : $Y_i = Y_i(1)T + Y_i(0)(1 - T)$

Counterfactual Example

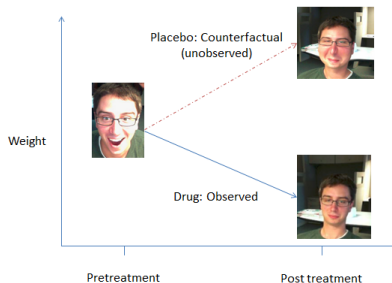


Figure 4: Example of a counterfactual

- Obviously we only observe one. *The fundamental problem of causal inference*
- If we could observe both, then we could observe the causal effects for each person

Rubin's Causal Model

- Stable unit treatment value assumption (SUTVA): Treatment status of another subject does not affect outcome of other units. Single version of each treatment
- Ignorability/No Unmeasured Confounders: $(Y(0), Y(1)) \perp T|X$, [4]

Estimands of Interest

- Individual Treatment Effect: $Y_i(1) - Y_i(0)$
- Average Treatment Effect (ATE): $E[Y(1) - Y(0)]$. The effect of moving entire population from treated to untreated
- Average treatment effect for the treated (ATT): $E[Y(1) - Y(0) | T = 1]$. The average treatment effect for those actually treated
- Note: $E[Y(1) | T = 1] \neq E[Y(1)]$, because $E[Y | T = 1] = E[Y_1 T + Y_0(1 - T) | T = 1] = E[Y_1 | T = 1] \neq E[Y(1)]$

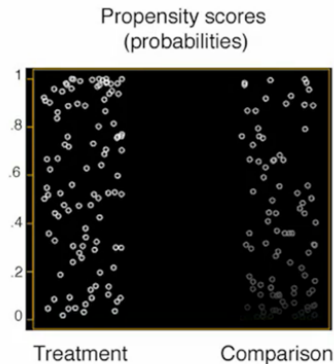
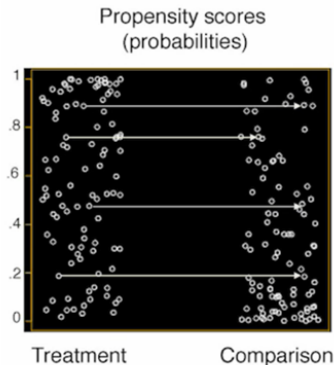
Definition

The propensity score is the probability that the subject received the treatment given the subjects *pretreatment* covariates. It is computed using the patient's baseline (pretreatment) information [4]

- Defined as $e_i(x) = P(T_i = 1|X_i)$
- Assume that the covariates play a role in how the subject chose treatment
- If we assume that $(Y(0), Y(1)) \perp T|X \implies (Y(0), Y(1)) \perp T|e(X)$, [4]
- Controlling for propensity score will make groups seem indistinguishable
- Thus, we may treat it as if it were an RCT

Common Propensity Score Methods

- Matching: Match treatment and controls on their propensity score, calculate ATE
- Weighting: Weight each observation by the inverse of its propensity score, and then calculate ATE



- IPTW: Inverse probability of treatment weights
- Idea: Weight sample by propensity score so that we get a sample where there is no confounding
- Weights: $1/e(X)$ for treatment, $1/(1 - e(X))$ for control
- Can be shown that
 - $E[\frac{TY(1)}{e(X)} | T = 1] = E[Y(1)]$
 - $E[\frac{(1-T)Y(0)}{1-e(X)} | T = 0] = E[Y(0)]$

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MI Notation

- Y is our whole dataset. It will have i rows and j columns. Some of the covariates in the dataset will be completely observed, and others will have missingness.
- Y_j is a specific column of Y . Y_j is composed as $Y_j = (Y_{j,obs}, Y_{j,mis})$, where
 - $Y_{j,obs}$ is the data we have observed for covariate j
 - $Y_{j,mis}$ is the missing data covariate j
- Y_{obs} is all of the data that we have observed
- Y_{mis} is all the data that we have not observed
- R is a binary matrix the same size as Y where a 1 indicates we observed the data, and 0 means it is missing
- ψ is a vector of parameters for the missing data model.
- The missing data model is given as $p(R|Y_{obs}, Y_{mis}, \psi)$
- θ is a vector of the parameters for the full model of Y
- ϕ_j is the unknown parameters of the imputation model

Missing data Mechanisms

Now, we may discuss the three main types of missing data mechanisms.

- MCAR: Missing completely at random:

$$P(R = 0 | Y_{obs}, Y_{mis}, \psi) = P(R = 0 | \psi)$$

- The missingness in the data is not at all related to any of the data that we do or don't have
- MAR: Missing at random:

$$p(R = 0 | Y_{obs}, Y_{mis}, \psi) = p(R = 0 | Y_{obs}, \psi)$$

- The missingness we have is related to something in the data
- MNAR: Missing not at random:

$$p(R = 0 | Y_{obs}, Y_{mis}, \psi)$$

does not simplify

- and the missingness depends on data that we have as well as have not collected

Full Conditional Specification (FCS)

- Assume MAR missing data mechanism
- Missing data is imputed iteratively on a variable by variable basis
- Drawing from $p(Y, R|\theta)$ through the full conditionals $p(Y_j|Y_{-j}, R, \theta_j)$
- Requires no distributional assumptions
- Idea: Specify k one dimensional models to impute on the missing data columns

1. Specify an imputation model $P(Y_j^{\text{mis}}|Y_j^{\text{obs}}, Y_{-j}, R)$ for variable Y_j with $j = 1, \dots, p$.
2. For each j , fill in starting imputations \dot{Y}_j^0 by random draws from Y_j^{obs} .
3. Repeat for $t = 1, \dots, T$:
4. Repeat for $j = 1, \dots, p$:
5. Define $\dot{Y}_{-j}^t = (\dot{Y}_1^t, \dots, \dot{Y}_{j-1}^t, \dot{Y}_{j+1}^{t-1}, \dots, \dot{Y}_p^{t-1})$ as the currently complete data except Y_j .
6. Draw $\dot{\phi}_j^t \sim P(\phi_j^t|Y_j^{\text{obs}}, \dot{Y}_{-j}^t, R)$.
7. Draw imputations $\dot{Y}_j^t \sim P(Y_j^{\text{mis}}|Y_j^{\text{obs}}, \dot{Y}_{-j}^t, R, \dot{\phi}_j^t)$.
8. End repeat j .
9. End repeat t .

Figure 6: FCS imputation pseudocode, taken from [1]

FCS Pros and Cons

Pros

- Flexible
- Easy to specify models
- Handles mixed continuous categorical

Cons

- No guarantee that full conditionals are compatible
- Slow
- Gets much harder as sample size increases to specify models

Setting Up The Model

- Specify the models
- Specify the predictors for each model
- Determine number of iterations and datasets to impute
 - This is a topic of hot debate
 - Old literature suggested 5 imputations, 5 iterations, but more now

Checking The Imputations

Convergence

- Chains should be freely intermingled with no pattern
- Convergence when variance between chains is no larger than variance within each chain
- Formal tests like Gelman/Rubin \hat{R} proposed to check convergence

Validation

- “Does the data look like it could have come from real data had it not been missing”?
 - Requires intimate knowledge of the data
- Graphical checks
 - Density plots
 - Conditional scatter plots
 - Box and whisker
 - etc.

- We now have m imputed datasets
- Run the analysis on each of the m complete datasets
- But we want one analysis, not m

Rubin's Rules

Let

- \hat{Q}_i be the scientific estimand from the i^{th} MI dataset
- U_i be the variance-covariance matrix of the i^{th} MI estimand

Then

- The MI estimate is given by $\bar{Q} = \frac{1}{m} \sum_{i=1}^m \hat{Q}_i$
- The MI “within” variance is given by $\bar{U} = \frac{1}{m} \sum_{i=1}^m U_i$
- the MI “between” variance is given by
 $B = \frac{1}{m-1} \sum_{i=1}^m (\hat{Q}_i - \bar{Q})(\hat{Q}_i - \bar{Q})'$
- Total variance given by [2]

$$T = \bar{U} + B + \frac{B}{m}$$

Inference with Rubin's Rules

- Assume that with complete data, inference on the estimand Q would be based on the statement $(Q - \hat{Q}) \sim N(0, U)$
 - \hat{Q} is the statistic estimating Q
 - U is the variance-covariance of $(Q - \hat{Q})$
- Since true T is not known, then

$$\frac{Q - \hat{Q}}{\sqrt{T}} \sim t_\nu$$

- ν is given by [6]

$$\nu = \frac{\nu_{old}\nu_{obs}}{\nu_{old} + \nu_{obs}}$$

- Where $\nu_{obs} = \frac{\nu_{com}+1}{\nu_{com}+3}\nu_{com}(1 - \frac{B+B/m}{T})$
- ν_{com} is the hypothetical complete sample degrees of freedom
- $\nu_{old} = \frac{m-1}{(\frac{B+B/m}{T})^2}$

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Kaplan-Meier in the MI Setting

- Clearly define the population, groups, and events of interest
- Ensure that we have non-informative censoring
- Algorithm: Pool the complimentary log-log of the Kaplan-Meier curve, get estimates, back transform [7]

Median Survival Time

- Want a measure of central tendency
 - Survival distributions often skewed, so mean is poor choice
- Median: smallest time such that $\hat{S}(t) \leq .5$
- Algorithm: Take ML Kaplan-Meier curve, observe first time it goes below 50%
- Confidence interval at median: the median of the upper and lower confidence bands

Log Rank Test

- Under no tied times, the score test on Cox Regression on a treatment is equivalent to the log rank test
 - And very similar under tied times
- Idea: Derive log rank test from Cox model
 - Pooling LRT and Score test is unstable [7]
 - Wald test is asymptotically equivalent
- Final Solution: Run the Wald test on Cox model as an approximation

Cox Model in the MI Setting

- Goal: To get a “baseline” Cox model, then add treatment variables
- Need to check for proportional hazards assumption
 - Problem: MI Cox model doesn't have residuals
 - Solution: Check assumptions (Schoenfeld residuals) on stacked dataset or each MI dataset individually
- Cox model is normally distributed, use Rubin's Rules to pool
- Add treatment covariates, rerun models, pool

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Propensity Score in MI Setting

- Mitra and Reiter propose two methods [8]
- Within: Work with propensity score on each of the m MI datasets
- Across: Average propensity scores across the m datasets and then analyze with the averaged propensities
- Which to use: Dependent on your data

Propensity Score: Questions To Ask

- Generating the propensity score
 - What to put into the model
 - How to compute it: Logistic regression, Probit regression, CART, GBM, etc.
- How to use the propensity score in the ML setting
 - Matching, weighting, covariate adjustment, stratification
- What estimand do we care about?

Verifying Balance

- Need the distribution of the groups to be similar
 - Standardized bias: $|\bar{X}_{k1} - \bar{X}_{k0}|/\hat{\sigma}_k$
 - Kolmogorov-Smirnov test
- Need to be sure that propensity scores are between 0 and 1

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Data Explanation

- 1514 MD Anderson patients who had brain mets from breast cancer between October 2009 and December 2012
- 1242 usable cases
- 90 covariates
 - Missingness from 0 to 65%

Type	Example
Subject data	Age range, race, date of birth
Breast Cancer data	TNM staging, type, receptor status
Pre brain mets data	Treatment types
Post brain mets clinical observations	Seizures, headache, nausea
Post brain mets data	Treatment type, type of brain mets
Survival data	Survival time after brain mets, censoring indicator

Want to explore...

- Chemotherapeutic drugs: Capecitabine vs other chemotherapeutic agents
- HER2 directed therapies (Lapatinib, Trastuzumab) in HER2+ subjects
- Note: treatment not determined at time of diagnosis, need to landmark (2 months)

Important Covariates

Name	Percent Missing	Meaning
hrher2	5	Categorical variable: The hormonal receptor and HER2 receptor status of the subject
agebrainmet	0	Indicator: Age greater or less than 60 at time of brain mets
timedx	1	Indicator: Time (years) from breast cancer diagnosis to brain mets diagnosis greater or less than 6 years
site5	1	Indicator: First metastasis was to brain
race2	0	Categorical: White, Black, Hispanic, other
priorn	0	Indicator: Number of prior treatments in metastatic setting before brain mets
braintype	4	Categorical: Single, multiple, Leptomeningeal disease
controlled	12	Indicator: Extracranial progression of brain mets
capeothno	18	Indicator: Capecitabine, other, or no chemotherapeutic treatment. Treatment variable 1
lapatrasno	18	Indicator: Lapatinib, Trastuzumab, or no HER2 treatment. Treatment variable 2
os	0	Overall survival (months)
dead	0	Indicator: death indicator
her2	10	Indicator: HER2 receptor status

Table 2: Table of important covariates to be used in the analysis

Visualization of Missingness



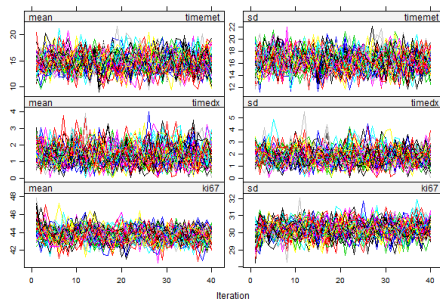
Figure 7: Visualization of missingness in the cancer dataset

Imputation

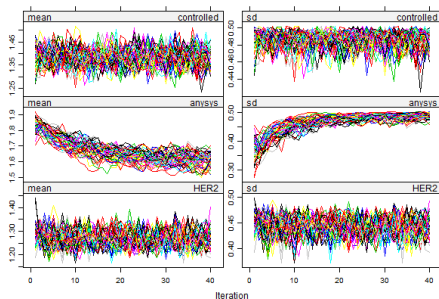
- MAR assumption seems reasonable
- FCS over JM due to nature of data
- Need to set up models and predictors
- Check for convergence and validity

Convergence

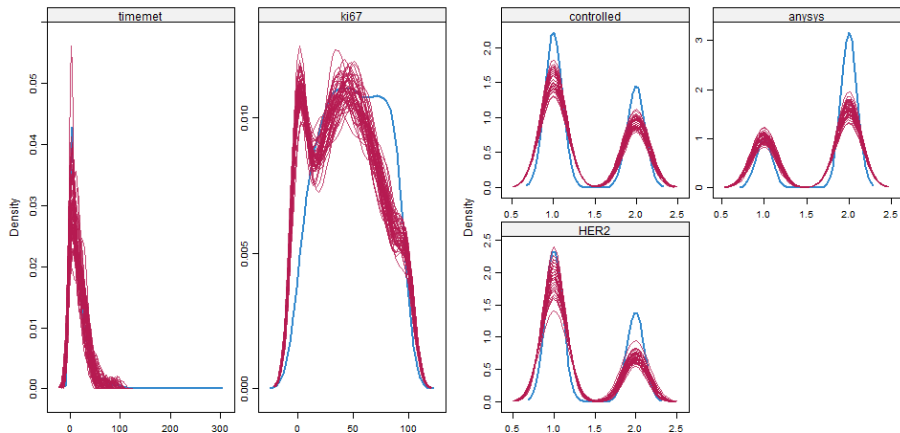
Traceplots, continuous



Traceplots, binary



Validity Checks



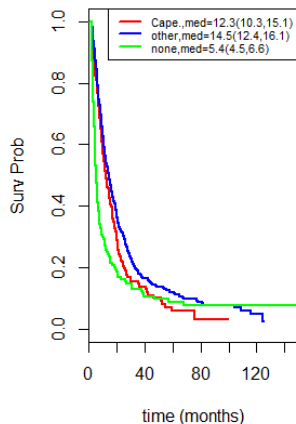
MI data Breakdown

	Sys therapy available case	Sys therapy MI	No Sys therapy available case	No Sys therapy MI
Age (mean,sd)	51.4(10.8)	51.2(10.9)	52.7(11.9)	52.9(11.4)
Breast Cancer subtype				
HR+/HER2-	27%	31%	28%	33%
HR+/HER2+	19%	18%	12%	13%
HR-/HER2+	22%	20%	15%	12%
Triple negative	32%	32%	45%	42%
Prior therapies for stage 4	1(0-3)	2(0-4)	2(0-4)	2(0-4)
Single brain lesion	25%	23%	23%	20%
Controlled extra-cranial	40%	40%	35%	36%
ECOG 0-1	84%	70%	53%	40%
Local Therapy				
Resection Alone	5%	5%	9%	7%
SBRT alone	13%	12%	9%	8%
WBRT	60%	59%	52%	53%
Resection/SBRT+WBRT	12%	14%	10%	8%
no local therapy	10%	10%	20%	23%

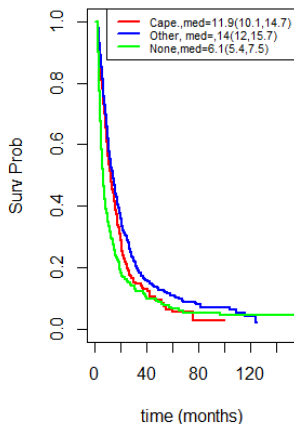
Table 3: Characteristics of available case data versus MI data ▶

Chemo KM and Log Rank Test

**Available case OS for chemo
2 month landmark**



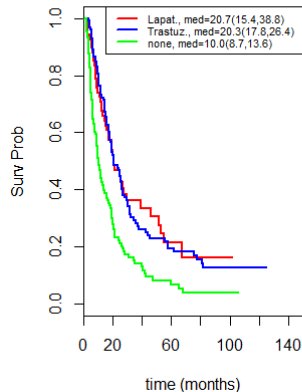
**MI OS for chemo
2 month landmark**



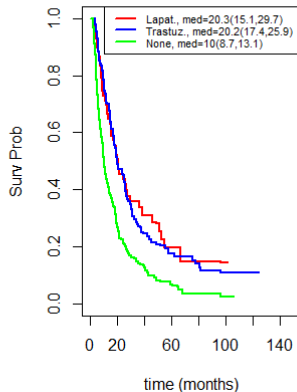
	Chemo	
	AC	MI
cape/other/none	<.0001	<.0001
cape/other	0.0321	0.033
cape/none	0.00039	.0016
other/none	<.0001	<.0001

HER2 Directed KM and Log Rank Test

**AC - OS for HER2 therapy
2 month landmark**

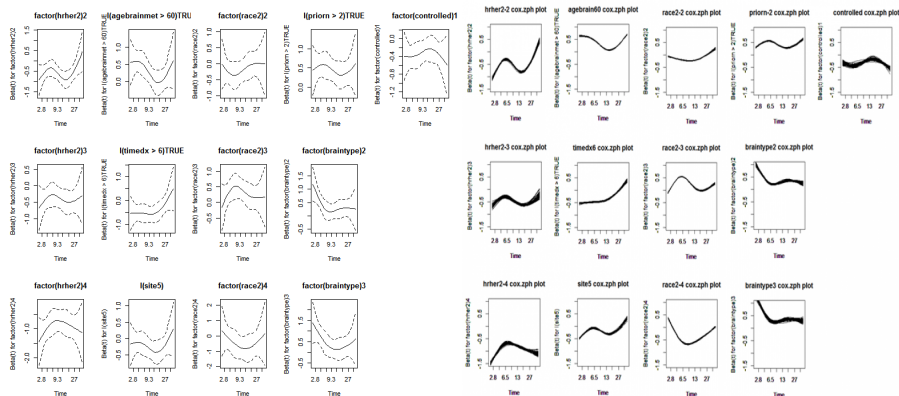


**MI - OS for HER2 therapy
2 month landmark**



	HER2	
	AC	MI
Lapat./Trastuz./none	<.0001	<.0001
Lapat./Trastuz	.87	.81
Lapat./none	.00017	.00018
Trastuz./none	<.0001	<.0001

Schoenfeld Residual Splines



Base model

			AC n= 845			MI		
Variable	Contrast	HR	95% CI	pvalue		HR	95% CI	pvalue (t test)
HR/HER2	-/+ vs. -/-	0.57	(0.46,0.71)	<0.0001		0.59	(0.48,0.72)	<0.0001
	+/- vs. -/-	0.66	(0.54,0.81)	<0.0001		0.63	(0.52,0.76)	<0.0001
	+/+ vs. -/-	0.4	(0.31,0.50)	<0.0001		0.4	(0.32,0.50)	<0.0001
Age	>60 vs. <60	1.37	(1.13,1.65)	0.0011		1.45	(1.22,1.72)	<0.0001
Dx to BM	>6 vs. <6	0.66	(0.54,0.82)	0.00013		0.71	(0.59,0.86)	0.0002
First DM	Brain vs. Oth	0.8	(0.66,0.97)	0.026		0.83	(0.70,0.99)	0.02
Race	Hisp. Vs. White	0.85	(0.68,1.07)	0.17		0.88	(0.71,1.08)	0.11
	Black vs. White	1.31	(1.06,1.63)	0.014		1.25	(1.02,1.52)	0.015
	Other vs. White	0.65	(0.40,1.04)	0.075		0.7	(0.45,1.07)	0.05
# prior Rx	>2 vs. 0-2	1.58	(1.31,1.91)	<0.0001		1.53	(1.29,1.82)	<0.0001
BM type	Mult. Vs. Single	1.45	(1.20,1.76)	<0.0001		1.48	(1.24,1.76)	<0.0001
	LMD vs. Single	1.6	(1.21,2.13)	0.001		1.58	(1.25,2.00)	<0.0001
Sys. Cont.	Yes vs. No	0.71	(0.61,0.83)	<0.0001		0.73	(0.63,0.85)	<0.0001

Table 4: AC and MI baseline Cox model

MI Cox Model, Chemo

			AC n= 745			MI		
Variable	Contrast	HR	95% CI	p-value		HR	95% CI	p-value (t test)
HR/HER2	-/+ vs. -/-	0.62	(0.49,0.79)	<.0001		0.63	(0.51,0.77)	<.0001
	+/- vs. -/-	0.65	(0.53,0.81)	0.00011		0.64	(0.53,0.78)	<.0001
	+/+ vs. -/-	0.41	(0.31,0.53)	<.0001		0.42	(0.34,0.53)	<.0001
Age	>60 vs. <60	1.34	(1.10,1.64)	0.0041		1.44	(1.21,1.72)	<.0001
Dx to BM	>6 vs. <6	0.72	(0.58,0.90)	0.0032		0.71	(0.58,0.86)	0.00039
First DM	Brain vs. Oth	0.77	(0.63,0.95)	0.014		0.81	(0.68,0.96)	0.016
Race	Hisp. Vs. White	0.77	(0.61,0.98)	0.034		0.86	(0.69,1.06)	0.15
	Black vs. White	1.29	(1.02,1.63)	0.032		1.23	(1.01,1.51)	0.043
	Other vs. White	0.76	(0.47,1.25)	0.28		0.7	(0.45,1.08)	0.11
# prior Rx	>2 vs. 0-2	1.61	(1.32,1.98)	<.0001		1.53	(1.28,1.82)	<.0001
BM type	Mult. Vs. Single	1.46	(1.20,1.78)	0.00017		1.51	(1.27,1.81)	<.0001
	LMD vs. Single	1.45	(1.04,2.03)	0.029		1.41	(1.11,1.80)	0.0049
Sys. Cont.	Yes vs. No	0.57	(0.48,0.68)	<.0001		0.69	(0.59,0.80)	<.0001
Chemo	Cape. vs. none	0.69	(0.53,0.89)	0.0046		0.75	(0.60,0.95)	0.018
	other vs. none	0.52	(0.42,0.65)	<.0001		0.58	(0.47,0.71)	<.0001

Table 5: AC and MI Cox model with Chemo Treatment

AC and MI Cox Model with HER2 Treatment

			AC n=292			MI n between 391 and 415		
Variable	Contrast	HR	95% CI	p-value		HR	95% CI	p-value (t test)
HR/HER2	+/+ vs. -/+	0.65	(0.49,0.87)	0.0036		0.66	(0.51,0.85)	0.0015
Age	>60 vs. <60	1.38	(0.95,2.01)	0.092		1.58	(1.15,2.18)	0.0054
Dx to BM	>6 vs. <6	0.64	(0.43,0.97)	0.033		0.69	(0.49,0.99)	0.041
First DM	Brain vs. Oth	0.84	(0.58,1.20)	0.34		0.86	(0.62,1.17)	0.34
Race	Hisp. Vs. White	0.69	(0.46,1.02)	0.064		0.76	(0.53,1.09)	0.14
	Black vs. White	1.41	(0.94,2.11)	0.1		1.43	(1.00,2.04)	0.047
	Other vs. White	0.7	(0.32,1.53)	0.38		0.83	(0.46,1.52)	0.55
# prior Rx	>2 vs. 0-2	1.88	(1.34,2.63)	0.00028		1.71	(1.28,2.28)	0.00028
BM type	Mult. Vs. Single	1.3	(0.92,1.86)	0.14		1.25	(0.91,1.70)	0.16
	LMD vs. Single	2.15	(1.20,3.88)	0.011		1.77	(1.10,2.83)	0.018
Sys. Cont.	Yes vs. No	0.73	(0.55,0.97)	0.029		0.78	(0.60,1.01)	0.063
HER2 therapy	Lapat vs. none	0.47	(0.32,0.69)	0.00015		0.52	(0.37,0.75)	0.00036
	Trastuz vs. none	0.45	(0.33,0.61)	<.0001		0.51	(0.38,0.68)	<.0001

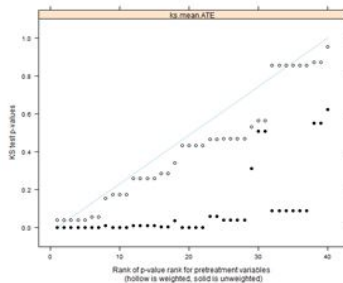
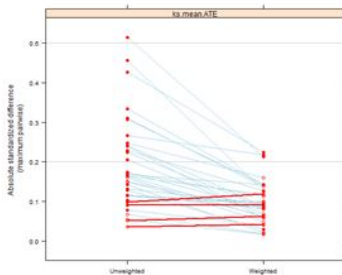
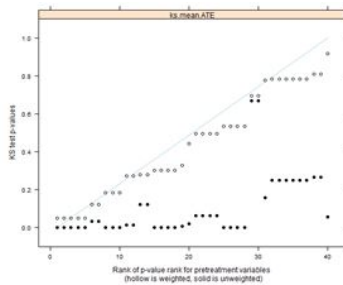
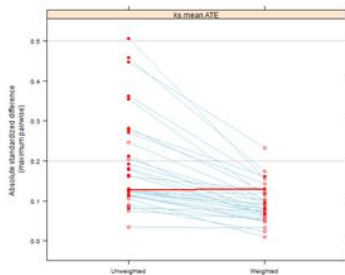
Table 6: AC and MI Cox model with HER2 Treatment

- Need to get propensity score from *pretreatment* covariates
- Check the balance and standardized bias
- Run IPTW analysis on each dataset
- Use Mitra and Reiter's within method to get each MI dataset ATE [8]
- Pool via Rubin's Rules

IPTW Weighting in MI Setting

- Set the propensity score model - GBM
 - Confounding factors: stage, race, IDC, breast cancer surgery, HR/HER2 status, breast cancer radiation, first met site, number of prior treatments, ECOG score, localized brain mets treatment, age at brain met, type of brain mets, brain met controlled
- Weight observations by propensity score
- Check to ensure balance achieved -KS and standardized bias
- Run IPTW weighted Cox regression on treatment
- Pool results via Rubin's rules

Balance Checks



Results of IPTW: Chemotherapeutic

	AC Unweighted		AC IPTW		MI unweighted		MI IPTW	
	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI
Cape. vs none	0.396	(0.325,0.482)	0.655	(0.481,0.894)	0.484	(0.400,0.585)	0.702	(0.543,0.906)
Other vs none	0.336	(0.287,0.394)	0.567	(0.46,0.754)	0.413	(0.354,0.481)	0.593	(0.470,0.748)
Cape. vs other	1.179	(0.983,1.416)	1.156	(0.966,1.383)	1.173	(0.981,1.402)	1.183	(0.998,1.404)

Table 7: Chemotherapeutic ATE with IPTW weights, AC and MI

!! Need to put in doubly robust table here!!!

Results of IPTW: HER2 Directed

	AC Unweighted		AC IPTW		MI Unweighted		MI IPTW	
	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI
Lapat. vs none	0.467	(0.355,0.616)	0.571	(0.381,0.855)	0.474	(0.362,0.622)	0.485	(0.304,0.775)
Trastuz. vs none	0.488	(0.398,0.597)	0.566	(0.421,0.759)	0.506	(0.417,0.614)	0.480	(0.313,0.735)
Lapat. vs Trastuz.	0.958	(0.693,1.324)	1.009	(0.680,1.496)	0.927	(0.673,1.28)	1.011	(0.763,1.338)

Table 8: HER2 directed ATE with IPTW weights, AC and MI

	AC Unweighted		AC IPTW		MI Unweighted		MI IPTW	
	HR	95% CI	HR	95% CI	HR	95% CI	HR	95% CI
Lapat. vs none	0.468	(0.316,0.692)	0.514	(0.331,0.798)	0.524	(0.367,0.747)	0.410	(0.257,0.652)
Trastuz. vs none	0.447	(0.328,0.6089)	0.456	(0.328,0.632)	0.511	(0.381,0.685)	0.388	(0.249,0.602)
Lapat. vs Trastuz.	1.048	(0.704,1.560)	1.128	(0.726,1.754)	1.026	(0.713,1.477)	1.057	(0.788,1.417)

Table 9: HER2 directed ATE with IPTW weights, double robust

Outline

1 Introduction

- The Problem
- Missing data
- Survival Analysis
- Causal Analysis

2 Methods

- Imputation
- Survival
- Causal Analysis

3 Application

- Breast Brain Mets Example

4 Conclusion

- Discussion

- MI doubters and method critiques
- Assumptions in the survival section
- Propensity score in general and model choices

Further research

- Differing imputation methods
- Competing risks
- AFT models
- Differing propensity score methods
- Estimating counterfactuals as an MI problem in MI setting

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- My committee - For assembling on such short notice
- Dr. Hess - For advising me
- Dr. Ibrahim and Dr. Bugano - For letting me work with their data and advising on project matters
- My family - For always supporting me
- My friends - For always believing in me and helping critique my thesis

Any Questions?

- Any questions?

Propensity Score Issues

- Unmeasured confounders
- Choice of pretreatment covariates in the propensity score model
- Different models and methods may lead to different conclusions

Joint Modelling (JM)

- Assume ignorable MAR missing data mechanism
- Missing data imputed by sampling from a user specified distribution
- A lot of theory developed for Normal, not much else
 - Normal imputation has been shown to perform well, even under non normality [9]
- Idea: pull imputations by missing data row pattern

1. Sort the rows of Y into S missing data patterns $Y_{[s]}, s = 1, \dots, S$.
2. Initialize $\theta^0 = (\mu^0, \Sigma^0)$ by a reasonable starting value.
3. Repeat for $t = 1, \dots, T$:
 4. Repeat for $s = 1, \dots, S$:
 5. Calculate parameters $\dot{\phi}_s = \text{SWP}(\hat{\theta}^{t-1}, s)$ by sweeping the predictors of pattern s out of $\hat{\theta}^{t-1}$.
 6. Calculate p_s as the number missing data in pattern s . Calculate $o_s = p - p_s$.
 7. Calculate the Choleski decomposition C_s of the $p_s \times p_s$ submatrix of $\dot{\phi}_s$ corresponding to the missing data in pattern s .
 8. Draw a random vector $z \sim N(0, 1)$ of length p_s .
 9. Take $\dot{\beta}_s$ as the $o_s \times p_s$ submatrix of $\dot{\phi}_s$ of regression weights.
 10. Calculate imputations $\dot{Y}_{[s]}^t = Y_{[s]}^{\text{obs}} \dot{\beta}_s + C_s' z$, where $Y_{[s]}^{\text{obs}}$ is the observed data in pattern s .
11. End repeat s .
12. Draw $\hat{\theta}^t = (\hat{\mu}, \hat{\Sigma})$ from the normal inverted-Wishart distribution according to Schafer (1997, p. 184).
13. End repeat t .

JM Pros and Cons

Pros

- Fast
- Easy to derive posteriors with common distributions

Cons

- Inflexible
- Limited to known distributions
- How to deal with mixed categorical and continuous missing data

The Stack Method

- Rubin's Rules work well, but not always
 - Ex: partitioning the MI data on an imputed variable
 - Taking the average is not a good idea
- Solution: Stack the MI datasets on top of each other to get one huge dataset
 - Will get unbiased results
 - But sample size is falsely inflated, thus cannot trust variance



KM issues in the MI setting

- Issue: Kaplan-Meier is not normally distributed
 - Solution: Complimentary log log transformation, pool [7]
- Issue: Imputations leave one KM curve much shorter than the rest
 - Solution 1: Truncate all curves at the lowest time
 - Solution 2: Extend the curves out to the longest time
 - Solution 3: Use the stacked method

Log Rank Issues in MI setting

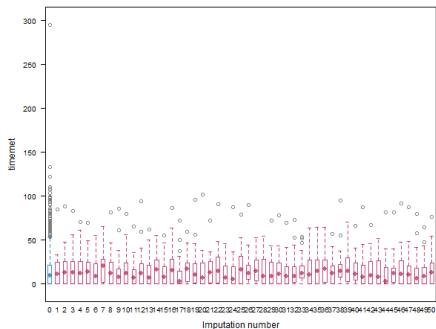
- Idea: Combine log rank tests from each MI dataset
 - Problem: Wastes information and is unstable [7]
 - Idea: Calculate log rank from the MI Kaplan-Meier curve
 - Problem: Risk set and deaths no longer meaningful

Setting up the model- Issues

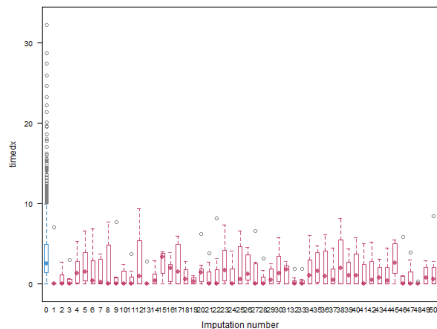
- Many categorical variables
- Collinearity between predictors
- Variables with poor influx/outflux [1]
- How many iterations and imputations to draw?

Validity Checks

bw plot, timemet by imputation



bw plot, timedx by imputation



Tabluar Checks

AC	os>10	
hrher2	FALSE	TRUE
-/-	0.246	0.0967
-/+	0.0916	0.0916
+/-	0.1917	0.1154
+/+	0.0712	0.0958
[[1]]	os>10	
hrher2	FALSE	TRUE
-/-	0.2432	0.0974
-/+	0.091	0.0878
+/-	0.2005	0.1184
+/+	0.0692	0.0926
[[2]]	os>10	
hrher2	FALSE	TRUE
-/-	0.2448	0.0966
-/+	0.0894	0.0886
+/-	0.1989	0.1192
+/+	0.0709	0.0918
[[3]]	os>10	
hrher2	FALSE	TRUE
-/-	0.244	0.095
-/+	0.0894	0.0894
+/-	0.1937	0.1208
+/+	0.0709	0.091

AC	controlled	
0	1	
0.6048	0.3952	
[[1]]	controlled	
0	1	
0.602254	0.3977	
[[2]]	controlled	
0	1	
0.605475	0.3945	
[[3]]	controlled	
0	1	
0.603865	0.3961	

AC	lapatrasno	
1	2	3
0.080709	0.1437	0.7756
[[1]]	lapatrasno	
1	2	3
0.0668	0.132	0.8011
[[2]]	lapatrasno	
1	2	3
0.0684	0.1337	0.7979
[[3]]	lapatrasno	
1	2	3
0.0741	0.1449	0.781







AC	capeothno	
1	2	3
0.206693	0.498	0.2953
[[1]]	capeothno	
1	2	3
0.190821	0.4686	0.3406
[[2]]	capeothno	
1	2	3
0.196457	0.4646	0.339
[[3]]	capeothno	
1	2	3
0.191626	0.471	0.3374

Figure 9: Selected tabluar checks

Issues with Propensity Score in our Setting

- Problem: Theory was developed for binary treatments, we have ternary
 - Solution: Run each treatment as binary, then compare groups
- Propensity score model specification
 - Solution: Boosting, subject to KS statistic minimization

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