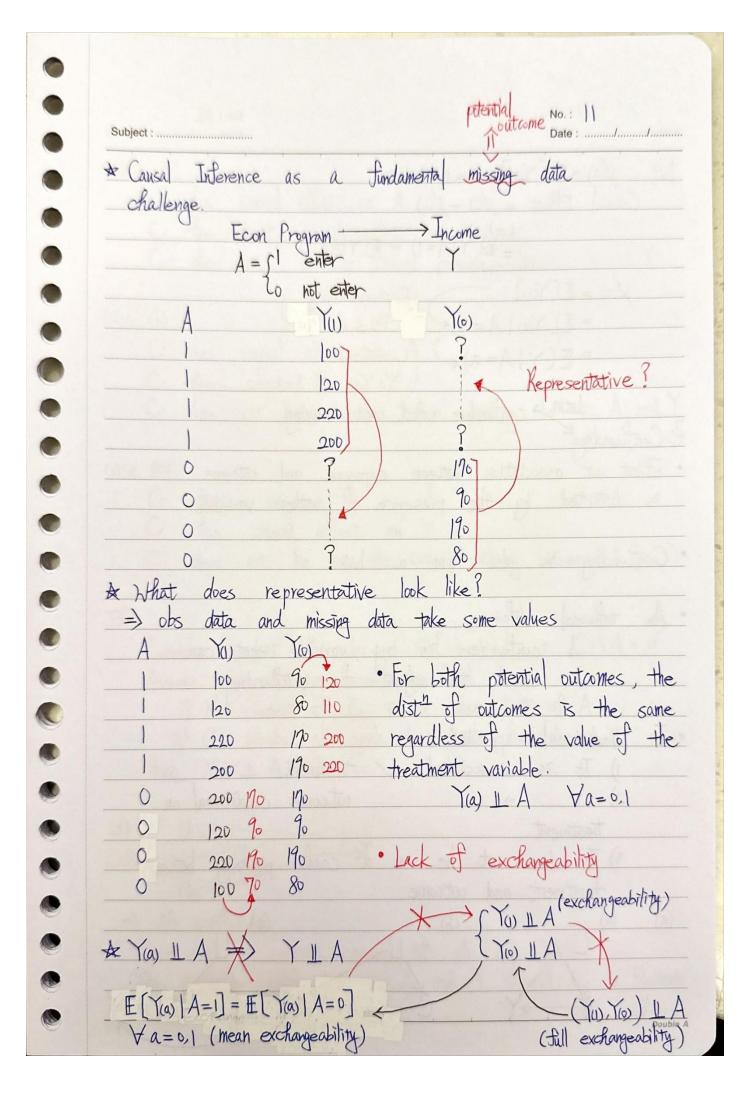
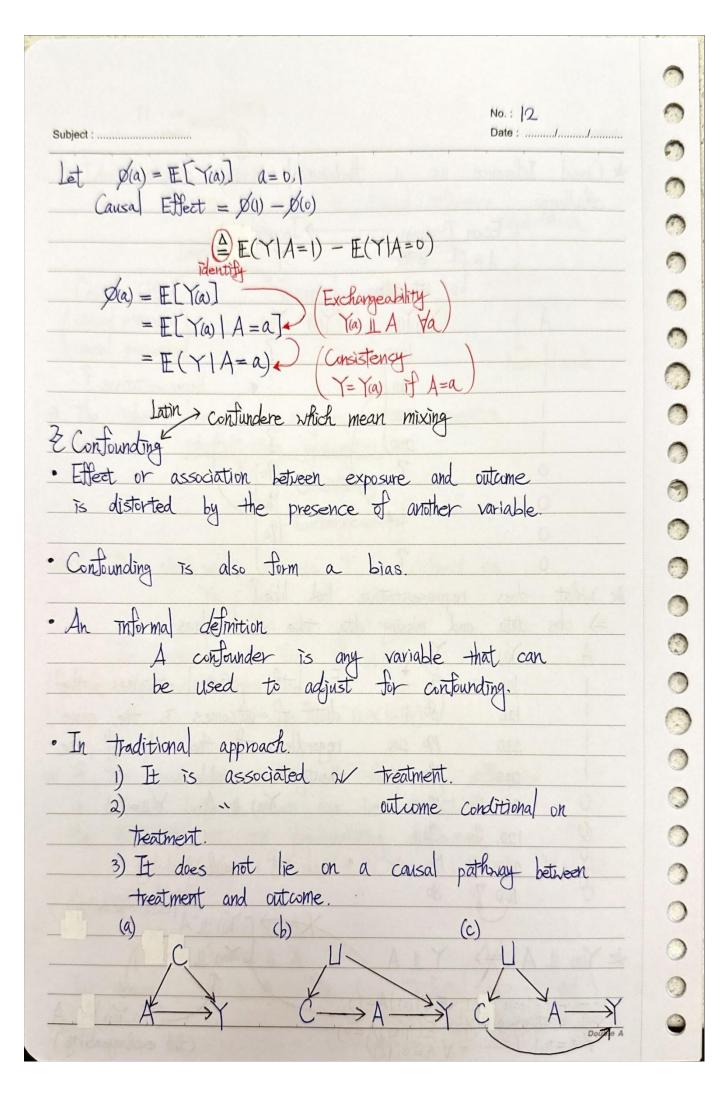
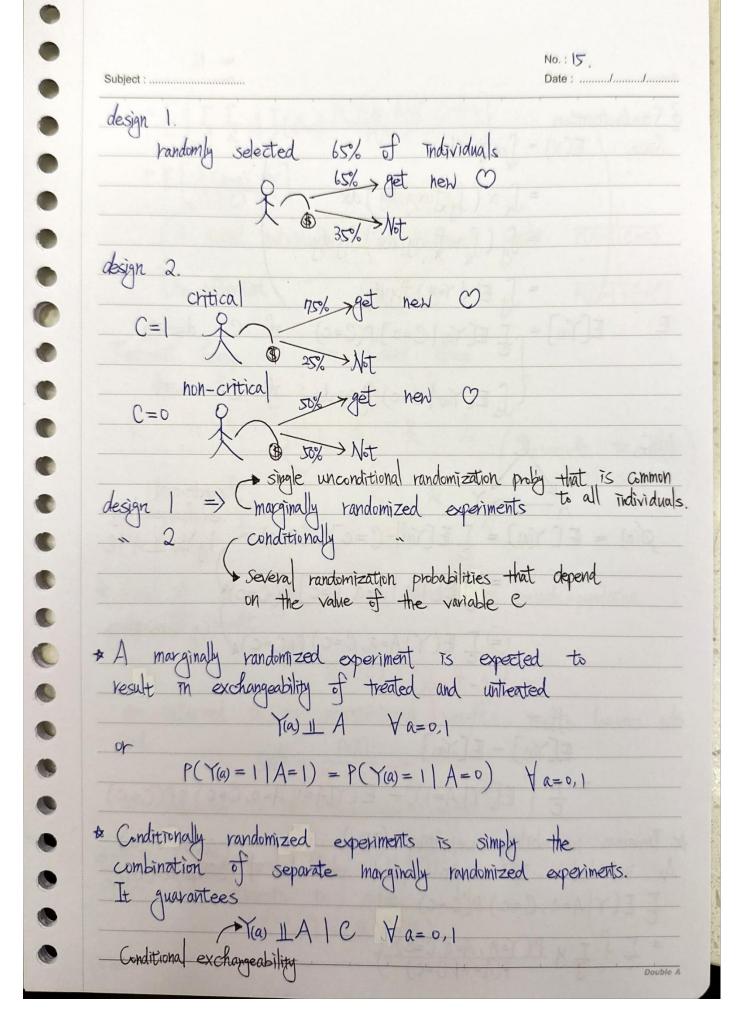
Subject:	No. :   <i>O</i>
& Identification	twent to mitlaged to
The key step in caus	sal Tifference
	Method/Tormula
Identication	Estimation
Causal Effect Statistical parame	Estimator Apply Listimate data
(causal estimand)	
(causal parameter)	Numerical result
· + -1	I WIND TO LET
* The identification proc require certain	ain assumption,
known as identification assumption.	
T Consistency	# # # N
Exchangeabilit	4
00	
otef. The observable outcome Y is	delined as
$Y = \begin{cases} Y(0) & \text{if } A = 1 \\ Y(0) & \text{if } A = 0 \end{cases}$	1 = lef = III summer b
In general, Y=Y(a) if A= Remark paired definition	
hemark. paired definition $Y = Y_0 A + Y_0 (1-$	1
1-10) A + 10)CI-	-n) /
A A tip groups are exchangeable that	n any difference
If two groups are exchangeable there in outcomes between them can be	attributed to
the treatment hat to non-existing	difference
the treatment, not to pre-existing $P(Y_{(a)}=1 \mid A=1)=P(Y_{(a)}=1 \mid A=1)$	$(a) = 1 \mid A = 0 \rangle  \forall a = 0.1$
or equivalently	(a) 1 1/1-0) V 12-0/
S You IL A	A Side - MADE
You IL A	X 4 A 3-3 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
1	unterfactual outcome and)
In general, Yas II A Ya (Co	wher actual outcome was

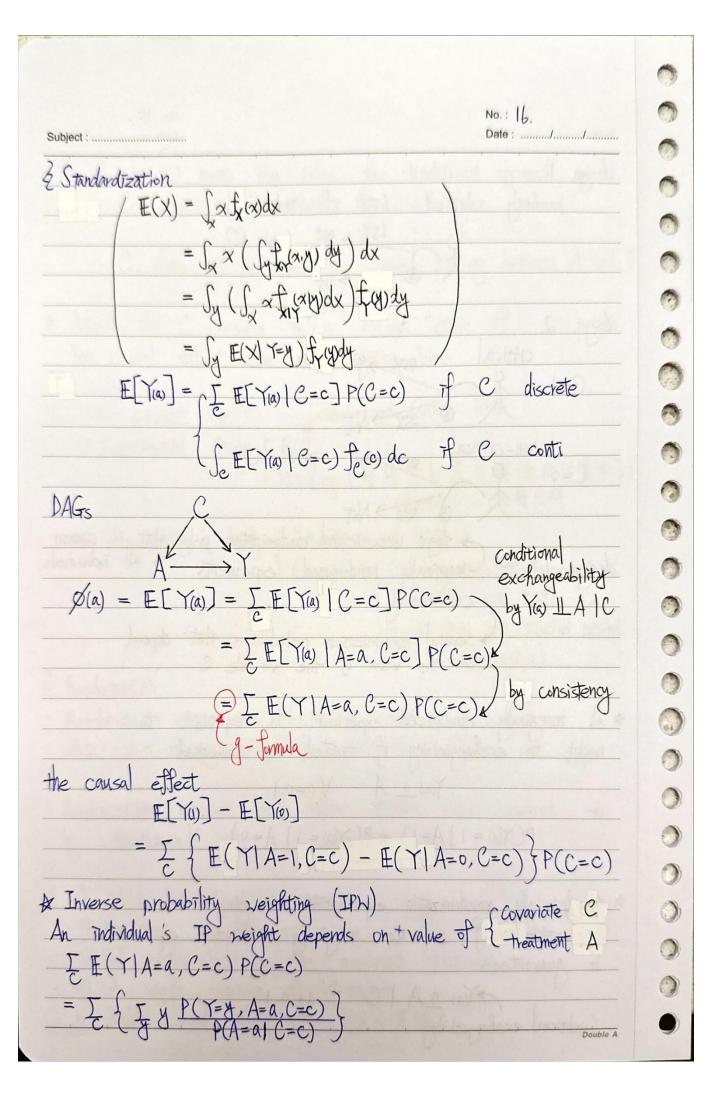




Subject:	No.: 13 Date:
Case (a)  C has causal effect on A  C has direct causal effect on Y  C does not lie on causal pathway	between A and Y
Case cb)  C has causal effect on A  C shares cause U W Y  C does not lie on the causal paths	vay between A and Y
case (c)  C shares cause U N/A  C has causal effect on Y  C does not lie on the causal paths	vay between A and Y
Thm.  Consider a set of nodes C = {\mu_0-,\mu_0}  Assume that following conditions hold:  i) No element of C is a descent ii) C blocks all back door paths  then Y(a) IL A   C Ya	dent of A
(d) U1  Collider C A Y  => No confounding  There are no common causes of A a  The backdoor path between A and Y  because C is a collider.	

Subject:	No.: 14  Date:
The C in (d) meets the rules for -	traditional approach
The C in (d) meets the rules for -	✓ A "
"Y A HAT LIVE TO THE STATE OF T	u Y
C does not lie on the causa	
* Adjust for C results in a biased the causal effect due to seletion	ester of
the causal effect due to seletion	bias.
	CORP. All Separate Company
How to correct it? Randomized	Control Trials
U) Experimental Study (RC)	
[Matching (公票 ]	要知道有與趣的干護因子) Restriction/分層分析
The state of the s	lestration / 分層分析
(2) Observational Study & Stratification (:	模型法
G-method Sta	ndardization i g-firmula
	Wester and MSM
	estimation and structual nested
	el (Semi-pavametric)
& Randomization	well the surpli
· Randomized experiments, like other real	study geneval
data 2/ missing value of counterfactive	a outcome.
J J	HA LENY AS
· Randomization ensures that those missing	9 values occured
by chance.	
	Ale MOSSIES MINISTER
* A ideal randomized experiment	
7) No loss to follow-up	
ii) Full adherence to the assigned	treatment
iii) A single version of treatment	A STATE OF THE STA
iv) Double blind assignment (双直実	驗)
	Double A





No.: 17 Subject:....  $= \int_{C} \left\{ \int_{C} \int_{C}$  $= \mathbb{E}\left[\frac{\mathbb{I}_{\{A=a\}}}{P(A|C)}\right]$ FA treated Individual N/ C=C receives weight P(A=1|C=c) " C=c\* (An untreated Contro Group Treatment Group Reveight to align woverall sample & Goal of IP weight is to create a pesudo-population.

The IP weight are referred to as nonstablized weights.  $SN^A = \frac{P(A)}{P(A|C)}$ and are referred to as stablized veights. 2 Marginal Structural Model (MSM) The model attempts to solve E[Y(a)] = M(a) &)

Subject:	No.:  8 Date:
(1) Linear model	DEST / 1711=
$\mathbb{E}[Y(a)] = \Theta_0 + \Theta_1 A$	# 1 D 1 V
(3) Log-linear model  log { E[Ta]} = Go + G1a (	(=) E[Y(a)] = C + G/a
(3) Logistic model  Jogit { E[Yia]} = Bo + BIA (	-> F[Y0] = 60+819
regit ( Et la) - 20 12 loc c	1- exteria
We use IP weight to conduct pesudo and fit	-population
$\mathbb{E}(Y A) = \theta_0 + \theta_1 a$	1 4 /3 and .
to the pesudo-population data by us	ing IP WLS
<ul> <li>In obs. studies, we do not control assignment mechanism.</li> <li>Presence of measured and unmeasure unblanced between group.</li> <li>Some structural (often untestable) ass</li> </ul>	ed confounders
a conditionally randomized experiments	Ť.
1. Well-defined treatment (=) Consistency	
The treatments being compared and match the ver. in da	ta.
2. No unmeasured confounding (=> Ex	chameability
realment assign depends only	on observed covariates C
3. Positivity	
Every treatment has non-zero	proby for all value of C

* What	covariates should be	adjust for obs. study?
	Z	
P	A	
. 0		)
	is confounder => Yes	- Not suggest
	is pure random hoise = does not bias the esté	
	duce unnecessary variabi	
	is instrumental variable	
/Z	does not bias the esté	although (1)
It	Increases variability. Howeve	er, W
unme	asured confounding, include affect Y only => Yes	it amplifies bias
4. 11 0	iffect Y only => Yes	
Sino	e 1 is predictive to	γ,
Indu	ding them often improve	precision/
3.3	is collider => No	1.+ 2.+
=) We	should adjust for at ve bias and more idea	least 0 10
	ve dias and more lae	aug, surmer
	st for U to reduce	