# Predict and analyze patient outcome from medical record

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#### Looking at data



#### 5 datasets with 5 trialss (studyA-E)

#### Variable:

- 1. Study A character indicating which of the five studies the data represents.
- 2. Country The country where the assessment was conducted.
- 3. PatientID An identification number given to each unique patient.
- 4. SitelD An identification number given to each unique assessment site.
- 5. RaterID An identification number given to each unique rater.
- 6. AssessmentiD An idenfication number given to each unique assessment conducted.
- 7. TxGroup A string corresponding to the patient's (randomly) assigned treatment group.
- 8. VisitDay An integer corresponding to the number of days that have passed since the baseline assessment.
- 9. P1-P7 The scores corresponding to each of the 7 positive symptoms of the assessment.
- 10. N1-N7 The scores corresponding to each of the 7 negative symptoms of the assessment.
- 11. G1-G16 The scores corresponding to each of the 16 general psychopathology symptoms of the assessment.
- 12. PANSS\_Total The sum of of the ratings across the 30 PANSS items.
- 13. LeadStatus A string indicating whether the assessment's audit passed, was flagged, or was assigned to a CS (i.e. clinical specialist).

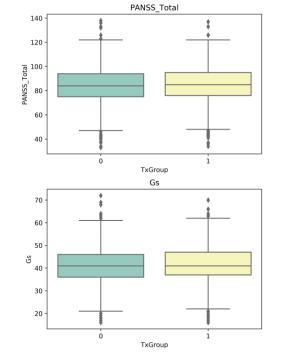


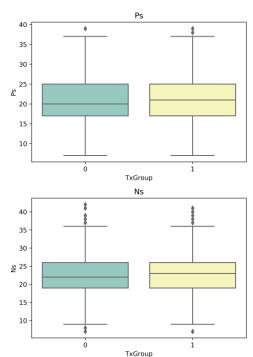
#### 5 datasets with 5 trialss (studyA-E)

Study	Country	PatientID	SiteID	RaterID	AssessmentiD	TxGroup	VisitDay	P1	P2	 G9	G10	G11	G12	G13	G14	G15	G16	PANSS_Total	LeadStatus
Α	USA	10001	20035	30076	100679	Control	0	5	5	 5	3	3	4	3	3	3	5	107	Assign to CS
Α	USA	10001	20035	30076	101017	Control	11	5	5	 5	3	3	4	3	3	3	5	109	Assign to CS
Α	USA	10001	20035	30076	102177	Control	18	4	4	 4	2	2	3	3	2	3	4	91	Passed
Α	USA	10001	20035	30076	101533	Control	25	3	3	 3	2	2	3	3	2	3	4	80	Flagged
Α	USA	10001	20035	30076	100930	Control	39	3	3	 3	2	2	3	3	2	3	4	77	Flagged
Α	USA	10001	20035	30076	100471	Control	53	3	3	 3	2	2	3	3	2	3	4	75	Flagged
Α	USA	10001	20035	30076	102347	Control	67	4	2	 3	2	2	3	3	2	3	4	72	Flagged
Α	USA	10002	20011	30016	100597	Control	0	5	5	 5	2	1	3	3	3	3	5	85	Passed
Α	USA	10002	20011	30016	100270	Control	7	5	5	 5	3	1	3	3	1	3	5	85	Passed
Α	USA	10002	20011	30016	101211	Control	9	5	5	 5	3	1	3	3	1	3	5	94	Passed
Α	USA	10003	20031	30058	101799	Treatment	0	5	5	 5	3	3	4	3	3	3	4	97	Flagged
Α	USA	10003	20031	30058	100330	Treatment	11	6	5	 6	4	4	4	4	4	5	5	128	Flagged
Α	USA	10003	20031	30058	101749	Treatment	18	6	5	 6	4	4	3	4	3	5	5	126	Flagged
Α	USA	10003	20031	30058	101301	Treatment	25	5	5	 5	4	4	4	4	4	4	5	119	Flagged
Α	USA	10003	20031	30058	101615	Treatment	39	4	4	 4	4	4	3	3	4	4	4	101	Flagged



Before the analysis: Ensure the treatment and control groups are substantially the same before the study begin – look at the summary statistic for the treatment and control groups





The median value is slightly different for Ps, Gs, Ns

→ the difference between the current score and the previous one for the same patient)



#### Before the analysis: Ensure the treatment and control groups are substantially the same before the study begin – look at the summary statistic for the treatment and control groups

		(	OLS Re	egress:	ion R	esults		
Dep. Variable	:	PAI	NSS_To	otal	R-sq	uared:		0.000
Model:				OLS	Adj.	R-squared:		0.000
Method:		Least	t Squa	ares	F-st	atistic:		1.36
Date:		Fri, 11	Sep 2	2020	Prob	(F-statistic):		0.24
Time:			14:34	4:37	Log-	Likelihood:		-12469
No. Observati	ons:			3000	AIC:			2.494e+04
Df Residuals:				2998	BIC:			2.495e+04
Df Model:				1				
Covariance Ty	pe:		nonrob	bust				
	coet	f std	err		t	P> t	[0.025	0.975
Intercept	84.4539	9	.396	213	501	0.000	83.678	85.229
TxGroup	0.6589	9	.564	1.	167	0.243	-0.448	1.769
Omnibus:			20.	.596	Durb	in-Watson:		1.10
Prob(Omnibus)	:		0.	.000	Jarq	ue-Bera (JB):		22.667
Skew:			-0.	.156	Prob	(JB):		1.20e-09
Kurtosis:			3.	.290	Cond	. No.		2.66

			(	DLS H	egress	ion R	esults		
Dep. Variabl	e:				Ps	R-sq	uared:		0.000
Model:					OLS	Adj.	R-squared:		-0.000
Method:			Least	t Squ	ares	F-st	atistic:		0.1662
Date:			ri, 11	Sep	2020	Prob	(F-statistic):		0.684
Time:				14:3	2:23	Log-	Likelihood:		-9597.3
No. Observat	ions				3000	AIC:			1.920e+04
Df Residuals					2998	BIC:			1.921e+04
Df Model:					1				
Covariance T	ype:			nonro	bust				
		coef	std	err		t	P>  t	[0.025	0.975]
Intercept	20.	7346	0	152	136	.524	0.000	20.437	21.032
TxGroup	0.	0883	0.	217	0	.408	0.684	-0.337	0.513
Omnibus:				34	.085	Durb:	in-Watson:		1.110
Prob(Omnibus	:):			9	.000	Jarq	ue-Bera (JB):		22.328
Skew:				9	.060	Prob	(JB):		1.42e-05
Kurtosis:				2	.595	Cond	. No.		2.60

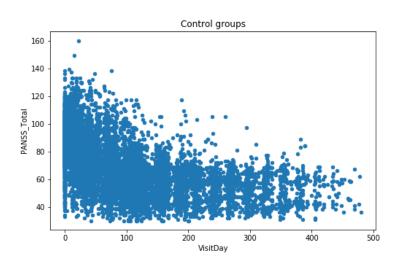
Null hypothesis test:
the difference between the
control and treatment groups is not statistically significant

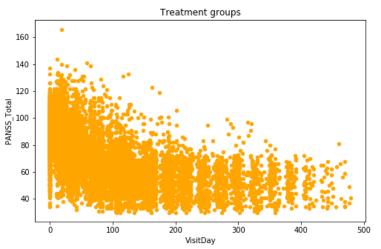
				OL	S Re	gress	sion R	esults			
Dep. Variab	le:					Gs	R-sq	uared:			0.001
Model:						OLS	Adj.	R-squa	red:		0.000
Method:			Lea	st	Squa	res	F-st	atistic	:		2.127
Date:			Fri. 1	1 S	ep 2	020	Prob	(F-sta	tistic):		0.145
Time:				1	4:33	:03	Log-	Likelih	ood:		-10639.
No. Observat	tions:				3	999	AIC:				2.128e+04
Df Residuals	s:				2	998	BIC:				2.129e+04
Df Model:						1					
Covariance 1	Type:			no	nrob	ust					
		coef	st	d e	rr		t	P>	t	[0.025	0.975]
Intercept										40.698	
TxGroup	0.	4472		0.3	07	1	1.458	0.	145	-0.154	1.048
Omnibus:					13.	534	Durb	in-Wats	on:		1.229
Prob(Omnibus	s):				0.	001	Jarq	ue-Bera	(JB):		14.705
Skew:					-0.	119	Prob	(JB):			0.000641
Kurtosis:					3.	246	Cond	No.			2.60

			OLS Re	gress	ion Resui	lts		
Dep. Variabl	e:			Ns	R-square	ed:		0.000
Model:				OLS	Adi. R-	squared:		-0.000
Method:		Le	ast Squa	ares	F-statis	stic:		0.4466
Date:		Fri.	11 Sep :	2020	Prob (F	statistic):		0.504
Time:			14:3	3:23	Log-Like	elihood:		-9107.9
No. Observat	ions:				AIC:			1.822e+04
of Residuals					BIC:			1.823e+04
Of Model:				1				
Covariance T	vpe:		nonrol	bust				
						P> t		
Intercept								
TxGroup	0.12	229	0.184	0	. 668	0.504	-0.238	0.484
Omnibus:			29	. 535	Durbin-	Natson:		1.862
Prob(Omnibus	):		0			Bera (JB):		33.761
Skew:			0	.186	Prob(JB)	):		4.66e-08
(urtosis:			3	.364	Cond. No	o.		2.66



#### Data visualization: Control groups v.s treatment groups: The plot looks similar







Original analysis: Used PANSS\_Total\_diff (the difference between the current PANSS\_total score and the previous one for the same patient) to form regression against Visitday and treatment

Because the PANSS\_total score varied from patient to patient, this ensures that we are evaluating the effects of the treatment, on the changes in the PANSS core.

The linear regression is: PANSS\_Total\_diff= $\beta 0+\beta 1TxGroup+\beta 2VisitDay+\beta 3TxGroup*VisitDay+\epsilon$ 

TxGroup =1 (Treatment) PANSS\_Total\_diff = $\beta 0+\beta 1+\beta 2 VisitDay +\beta 3 VisitDay +\epsilon$ =  $\beta 0+\beta 1 + (\beta 2+\beta 3) VisitDay +\epsilon$ TxGroup =0 (Control) PANSS\_Total\_diff = $\beta 0+\beta 2 VisitDay +\epsilon$ 

Null-hypothesis: β3=0

If the p-value for TxGroup\*VisitDay is < 0.05: statistical significance



OLS Regression Results

Dep. Variable:	PANSS_Total_diff_1	R-squared:	0.023
Model:	OLS	Adj. R-squared:	0.023
Method:	Least Squares	F-statistic:	155.7
Date:	Sat, 31 Oct 2020	Prob (F-statistic):	8.91e-100
Time:	21:05:42	Log-Likelihood:	-68247.
No. Observations:	19962	AIC:	1.365e+05
Df Residuals:	19958	BIC:	1.365e+05
Df Model:	3		
Covariance Type:	nonrobust		

	coef	std err	t	P> t	[0.025	0.975]
Intercept VisitDay TxGroup TxGroup:VisitDay	-3.6212	0.110	-32.834	0.000	-3.837	-3.405
	0.0122	0.001	15.307	0.000	0.011	0.014
	-0.0239	0.157	-0.152	0.879	-0.331	0.283
	-3.638e-06	0.001	-0.003	0.997	-0.002	0.002

Omnibus:	0.148	Durbin-Watson:	2.053
Prob(Omnibus):		Jarque-Bera (JB):	48996.418
Skew:		Prob(JB):	0.00
Kurtosis:	10.669	Cond. No.	543.

#### Warnings

p-value for TxGroup\*VisitDay = 0.997:

not statistical significance

<sup>[1]</sup> Standard Errors assume that the covariance matrix of the errors is correctly specified.



#### Analysis 1 + added categorical variables for different studies to the regression

In analysis 1, Different studies (trials) have different patient population. For example, the patient from study A is all from USA and the patient from study E is from China → Different studies may have bias result to the analysis

The linear regression become: PANSS\_Total\_diff=  $\beta 0+\beta 1TxGroup+\beta 2VisitDay +\beta 3TxGroup*VisitDay +\beta 4PatientID +\epsilon$ 

PatientID[T.50508]	2.7947	3.716	0.752	0.452	-4.489	10.078	
PatientID[T.50509]	2.9785	4.800	0.621	0.535	-6.429	12.386	
PatientID[T.50510]	2.5785	4.800	nan	nan	0.429	0	
PatientID[T.50511]	0	9	nan	nan	0	0	
PatientID[T.50512]	-3.4084	4.800	-0.710	0.478	-12.817	6.000	
PatientID[T.50513]	5.2562	8.031	0.654	0.513	-10.485	20.998	
TxGroup	2.8733	3.036	0.946	0.344	-3.077	8.824	
VisitDay	0.0180	0.001	18.008	0.000	0.016	0.020	
TxGroup:VisitDay	-0.0007	0.001	-0.514	0.608	-0.004	0.002	
=======================================						=	
Omnibus:	282	24.671 Dur	rbin-Watson:		2.27	0	
Prob(Omnibus):		0.000 Jar	rque-Bera (J	B):	41659.49	6	
Skew:		0.064 Pro	ob(JB):		0.0	0	
Kurtosis:	1	L0.076 Cor	nd. No.		2.31e+2	.0	
=======================================	========	.========		========		:=	

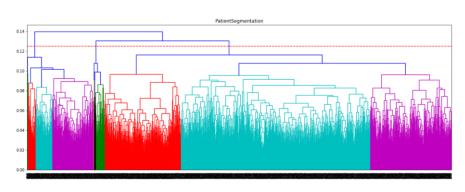
The p value for TxGroup variable is  $0.351 \rightarrow$  Fail to reject the null hypothesis

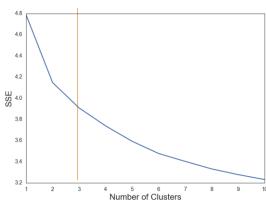


# Patient segmentation – understand each groups oh patients for their mental health status

#### Data preprocess: data is first normalized (The distance measures are affected by the scale of the variables) Data mining/evaluation: complete linkage clustering/ K-means clustering

- The red dotted line indicates the level at which the data is segmented. The dissimilarity between the three clusters created by seat the dotted line is greater than the ones that would have been created by cutting at a higher level (for two segments). Sub-segments below that are relatively similar with the other segments in their respective groups.
- Interpretable



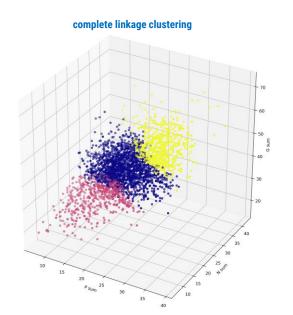


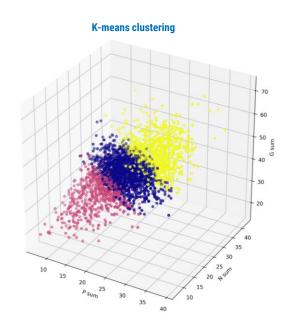


# Patient segmentation – understand each groups oh patients for their mental health status

Data preprocess: data is first normalized (The distance measures are affected by the scale of the variables)

Data mining/evaluation: complete linkage clustering and K-means clustering (k=3)





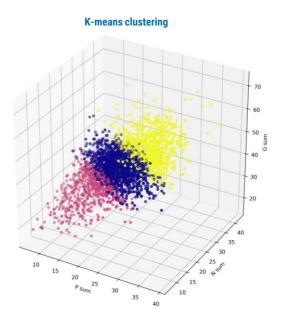
K-means clustering shows better separation of the individual clusters from each other.



# Patient segmentation – understand each groups oh patients for their mental health status

Data preprocess: data is first normalized (The distance measures are affected by the scale of the variables)

Data mining/evaluation: complete linkage clustering and K-means clustering (k=3)



- 1. Each cluster indicates a different combination of the sub-group scores. Moving from he 'Pink' cluster to the 'Blue' cluster requires an increase in all three sub-group scores.
- 2. To move from 'blue' cluster to 'yellow' cluster only requires a relatively smaller increase in the 'G' subgroup sum, whereas relatively similar changes in 'Ps' sum or 'Ns' sum doesn't result in crossing over to the next cluster.

