

Tentative solutions to TMA4267 Linear statistical models, 19 May 2017 – English

Problem 1 Random vector

a) Find a constant matrix C such that Y = CX.

$$\boldsymbol{C} = \begin{pmatrix} \frac{1}{3} & \frac{1}{3} & \frac{1}{3} \\ 1 & -1 & 0 \end{pmatrix}$$

Find $E(\mathbf{Y})$ and $Cov(\mathbf{Y})$.

$$E(\mathbf{Y}) = \mathbf{C} E(\mathbf{X}) = \mathbf{C} \begin{pmatrix} 0 \\ 0 \\ 0 \end{pmatrix} = \begin{pmatrix} 0 \\ 0 \\ 0 \end{pmatrix}$$

$$Cov(\mathbf{Y}) = \mathbf{C} Cov(\mathbf{X}) \mathbf{C}^{T} = \begin{pmatrix} \frac{1}{3} & \frac{1}{3} & \frac{1}{3} \\ 1 & -1 & 0 \end{pmatrix} \begin{pmatrix} 1 & \rho & \rho \\ \rho & 1 & \rho \\ \rho & \rho & 1 \end{pmatrix} \begin{pmatrix} \frac{1}{3} & 1 \\ \frac{1}{3} & -1 \\ \frac{1}{3} & 1 \end{pmatrix}$$

$$= \begin{pmatrix} \frac{1}{3} & \frac{1}{3} & \frac{1}{3} \\ 1 & -1 & 0 \end{pmatrix} \begin{pmatrix} \frac{1}{3} + \frac{2}{3}\rho & 1 - \rho \\ \frac{1}{3} + \frac{2}{3}\rho & -(1 - \rho) \\ \frac{1}{3} + \frac{2}{3}\rho & 0 \end{pmatrix} = \begin{pmatrix} \frac{1}{3} + \frac{2}{3}\rho & 0 \\ 0 & 2(1 - \rho) \end{pmatrix}$$

What is the distribution of Y?

Y is a vector of linear combination of a multivariate normal random vector and is therefore multivariate normal, with mean and covariance matrix given above.

Are Y_1 and Y_2 independent?

Yes, since $Cov(Y_1, Y_2) = 0$ and \mathbf{Y} is multivariate normal, then Y_1 and Y_2 are independent.

16: focus on D

I is positive definite when CTIC>O for all column vectors C ≠ 0, and this is the case when all eigenvalues of I are possible. We have

M= 1+29, so A1>0 if 1+29>0 8>-2

De= 25=1-9, so 22>01f 1-9>0 9<1

This means $g \in (-\frac{1}{2}, 1)$ will give I possible definite

We went I to be positive definite, because then any linear combination of our X's my have vanance >0.

CTX has Car(CIX). CIZC = Venerce of CIX

so if CTZC >0 then this is sakstied.

Let $P = [e_1 e_2 e_3]$ be the matrix with the eigenvectors of I as column vectors. We have

 $Z = \begin{bmatrix} e_i T Z \\ e_i T Z \end{bmatrix} = P^T X$. $\begin{bmatrix} e_i T Z \\ e_j T Z \end{bmatrix}$ We also have the spectral decomposition of Σ as $\Sigma = P \wedge P^T$, where $\Lambda = diag(\lambda_i)$.

[0 2 8]

We know that if I'mon, then AX+bis also muN, so Z is multivensherenal with E(Z)=0 since E(X)=0

So, $(Z) = P^T \sum_{i} P_i = P^T P_i P_i P_i P_i A_i$ (0)

Finally: $e_{1}^{T}X+e_{2}^{T}X+e_{3}^{T}X\sim N_{1}(0,(428)+(1-8)+(1-8))$ $=N_{1}(0,3+28-8-8)=N_{1}(0,3)$ $P(W>4)=1-P(W=4)=1-P(\frac{4-0}{\sqrt{3}})$ =1-P(231)=1-69896=0.0104

Note: Var(W) = 3 for all choices of g.

I did write g=0.5 so that it was possible to show eq, ez, ez, and not confuse the sudent on the missing value of g. Hopethy noone got confused by that...

Problem 2: Modelling systolic blood pressure

a) nodel A: Q?'s

BHI: Std. Error pussing

Std. Error 15 SD (Beni), where

Cu(B) = (XTX)-1 or and

design

motific

Ver ($\hat{\beta}$ wii) is the corresponding aragonal denont of $Cov(\hat{\beta})$, call this $Cij \cdot C^2$.

50 (p'eni) = (Gj · C

where $\hat{\sigma}^2 = \frac{SSE}{n-p}$ and $SSE = \frac{1}{2}(Y_1 - Y_1)^2$

vector of X(XIX)-'XI
response.

Numerical value: $T_0 = \frac{\hat{R}_1}{\hat{SD}(\hat{r}_1)} \Longrightarrow \hat{SD}(\hat{r}_2) = \frac{\hat{R}_1}{\hat{T}_2} = \frac{1.01050}{10.129}$

 $so(\hat{r}_{j}) = 0.0998 \approx 0.1$

This is the enhanted standard deviction of the enhanced regression coefficient.

Pr(> It1) missing for SEX

We want to test

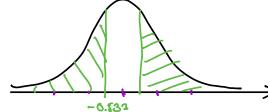
and use as test statistic (let j densk SEX)

$$T_{j} = \frac{\beta_{j} - 0}{\text{so}(\beta_{j})}$$

 $T_{ij} = \frac{\beta_{ij} - 0}{\beta_{ij}} \quad \text{where } \beta_{ij} \text{ is the 5th element}$ $SD(\beta_{ij}) \quad \text{of } \beta_{ij} = (X^{T}X)^{-1}X^{T}Y$ and SD(B) as given on previous page.

The p-value of the text is "Pr(>|t1)" and calculated 2. P(Tj > It, I) her to is the observed value of the test stetistic.

When its is true Ti, N thep. We have observed



$$t_1 = -0.533$$
 end $n-p=2593$

The +- distribution with 2593 degrees of freedom is very close to the WW,1) distribution.

This prake is larger than any sensible drove of significance level, so we do not reject the and believe SEX does not influence SYSBP in our model. Ho: Box = 0 W H. Box \$0

Model fit of model A:

- OR2 = 25% which is low, but for a medical problem we might not be able to get much higher.
- o The regression is significent, that is, Ho: B1=B2= --= B6=0 13 Hi: at least one +0 15 rejected, p-valu < 2.2.10-16.
- o ofodel requiements: looking at the residual plat in the left penel of Figre 2, g on x-axis

-linearly of cors oh? & on y-axis may be poblem with homosc. it bobs roughly rendom. It might be slight enviolet) down werds tend end lengt venerlager small values of g, but ther is not clear. the ag-plat (right penel) does not look like a straight line the teils are denoting a lot - which may imply that the assumption of normality of errors is violated also Andrsonfergin Rg2.
reject this mend.

2b) model A: SYSBP is cons.
B: - I us cons.

Residual plots: left penul of Figure 2 and 4 are not very different, maybe less of a downward trand in model B-plot than model A. The gg-plot for model B is very good - we can not reject normally of residuals.

The would prefer model B.

Anderson Derling 3

A full model night include venebles that do not influence the response, and thereby for noise insheed of signal, thus overfitting might be a problem. This will in perticular give a bad performance for prediction because over fitting increases the venere of \$\beta_j^2 \sqrt{s}.

In best subset selection we exemine all possible regression models, that is, with 6 coverates (we use have) we have $2^6 = 64$ possible ways of including or not the 6 different coverates.

We stert by looking at all models of equal size, and select the best based on SSE or R2. In the printent the best model for each of the sizes 1-6 is giran. E.g. best model with one corenste include aug BrnEDS, but with 2 include AGE+BPTEDS, atc.

After the best model for each size is found we need to choose between models of different sizes, and for that we could use SSE because SSE will never decrease when new coverleted are added to the nodel. Instead we use a penalized version by adding penally term for include many cover else. The BIC criterion is based on adding a penalty to -2 lighthelihood of the fitted rocal.

BIC = $n \cdot (n(\delta^2) + (n(n)(|M| + 1))$ for σ^2 penelty size of model =

term number of overal.

We choose the model with the lowest Bic. In our case this is the model with 4 coversus, but this model is not very different from the model with 3 coverses.

Choose no del

AGE + Bril + TOTCHOL+ BPMEDS

with lowest BIC

20: Teshng linear hypotheses

$$f_{obs} = \frac{\frac{1}{r} \Delta SSE}{\frac{88E}{n-p}}$$

Since de mpsse, we may use of from

r=3 runder of thep SSE = 11 model B SSEHS = 11 read C

ΔSSE- SSENG-SJE N-p= 19 noael O (= 2593 her)

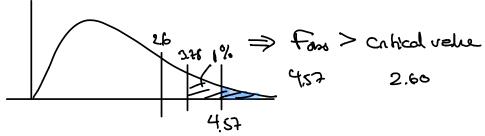
when to is true Foos or Fe, n=p

SSE=
$$0.005819^2 \cdot 2593 = 0.08480$$

SSEH,= $0.005831^2 \cdot 2596 = 0.08826$

$$F_{obs} = \frac{\frac{88E_{Hb} - 88E}{3}}{\frac{88E}{2592}} = \frac{0.0812L - 6.08780}{3} = 4.57$$

F(3, 2593) with ere 0.05 to the right: 2.60 3.78



(0.008 according to R) => reject Ho

we prefor model B

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Problem 3 Design of experiments

What type of experiment is this?

We see that we have a full factorial design in the factors A, B, C, but there is a fourth factor D added. This is a half fraction of a 2^4 design, also called a 2^{4-1} -design.

	Α	В	С	D	ABC
1	-1	-1	-1	1	-1
2	1	-1	-1	-1	1
3	-1	1	-1	-1	1
4	1	1	-1	1	-1
5	-1	-1	1	-1	1
6	1	-1	1	1	-1
7	-1	1	1	1	-1
8	1	1	1	-1	1

What is the generator and the defining relation for the experiment?

The generator for the design is D = -ABC (which is seen from the table above after the ABC column is added). The defining relation is then I = -ABCD.

What is the resolution of the experiment?

The resolution of the design equals the number of letters in the defining relation and is given using Roman numerals. Thus, the resolution is IV.

Write down the alias structure of the experiment.

Main effects and 3-factor interactions: A = -BCD, B = -ACD, C = -ABD, D = -ABC 2-factor interactions with eachother: AB = -CD, AC = -BD, AD = -BC

Why perform the experiments in random order?

To minimize the potential influence of external factors not part of the experimental plan.

Problem 4: Underfitting

Y = In out Ex underfitted model

i) Show that H is idempoked and find the trace of Hy

Hi Hy = Xy (Xy Xy) Xy Xy (Xy Xy) - Xy = Xy (Xy Xy) - Xy = Hy

tr (Hy) = tr (Xy (Xy Xy) - Xy) = tr (Xy Xy (Xy Xy))

| kry nxh wh

= tr (F) = k

tr (AB) = tr (BA)

Since this idempolent, I-Hy is also idempolent.

Hut: brece formula: E(XTAX)= tr(AI) + pTAp E(X)= p Car(X)= E.

=
$$G^{2}$$
 tr($T-H_{1}$) + $(X_{1}B_{1}+X_{2}\beta_{2})^{T}(J-H_{1})(J-H_{1})(X_{1}P_{1}+X_{2}P_{2})$
 $\beta_{2}TX_{2}^{T}(J-H_{1})(J-H_{1})X_{2}\beta_{2}$

$$E\left(\begin{array}{c}SSE_{4}\\N-h\end{array}\right)=O^{2}+\frac{\beta IX_{2}^{T}(I-H_{1})X_{2}\beta_{2}}{n-h}$$

Problem 5: Independence of linear combinations

$$X \sim N_P(\mu, Z)$$
, $M_R(t) = \exp(\mu t + \frac{1}{2}t^T Z t)$
A and B
The rep

i) Show that YN Nger using naf.

$$M_{\gamma}(t) = E(exp(t^{T}Y)) = E(exp(t^{AX}))$$

$$1 \times (r+q)$$

ii) Condition for when AI end BI ere independent.

AX and BX are all components AX are independent of all components of BX all components of BX

Components se O

A IBT = O and BZAT = (AZBT)T

That is, A IBT = O is a condition for when

AX and BX are independent

(The condition is necessary and sufficient)