Chiy Analysis of Categorical Data In many cases, the data we collect is categorical in nature i.e. the observations full into one of soveral distinct categories and we record the court or proportion c.g. smoker/formersmoker/nonrsmoker e.g. SA/A/N/0/SD (5-pt. Likert scale for surveys) The no. of observations that full into each category follows a multinomial dist. Recall the multinomial experiment (p. 279) 1 experiment has a identical trials Doutcome of each firel falls into one of k classes or cells 3 pi= P(outcome falls mcolli) i= 1 tok Py+ P2+ 11- + PK=1 (1.e. pi constant for every tral) 5 Yi = no. of outcomes that full in celli is I tok Y1+ 42+ ...+ 46 20

The jt. prob. fet (from ch.4)

P(41, ..., 7k) = 1 1 12 ... Pk

of course, we are generally interested in conducting inference on Ri (proportion of smokers / former smokers; proportion of defectives (seconds, etc.)

compersome >> Karl Penson developed the X2 test which

is an approximate test.

14.2 The Chi-square Test

as we have shown before, Z~N(0,1) then W= 22- x2,17

. We also have the CLT ...

Let Y, ... Yn he cid w/ E(Y)= 4 and V(Y)= +2 ~

12 = 1 - N = 2 - N (011)

Greature It & 0 H(011) monig x - A 5 0 2 Kg (1) ;

1st, recall the definition of convergence in dist.

Def

Let EYn3 be a sequence of RVs and let Y be a RV. Let C [Fow]

denote the set of all pts where Fly) is continuous. We say that

Yn converges in dist to y if

How from + from An e cl tros)

denoted Yn - Y

CLT says that Yn = ET: - NM O MICILD

=> lim Fyn(y) = Q(y), Q(s) is the cat of a M(0,1)

Let $X_n = (Y_n)^2$ and Let $H_n(x) = P(X_n \le x)$ be the of X_n $\Rightarrow H_n(x) = P(X_n \le x) = P(Y_n^2 \le x) = P(-\sqrt{x}) \le Y_n \le \sqrt{x}$ $= F_n(\sqrt{x}) - F_n(-\sqrt{x})$

> lim Haix = lim Fa(JE) - lim Fa(-JE)

= $\Phi(JR) - \Phi(-JR)$ note: $\Phi(y)$ cont. everywhere

Symmetry $\frac{1}{2}$, $\frac{1}{\sqrt{2\pi i}}$ e dy

4. y2 = du= 2y dy = du= 2 50 dy

= 2. 5 1 e 2. du

= (1/2)-21/2 u e du

cdf of x~GAM(x=1/2, β=2)~χ²(1)

=> Xn = (Yn)2 -> 1/2(1)

Consider a binomial dist. (2 categories S/F)

Let y ~ Bin (n, p) and 12= n-y, => 12 ~ Bin (n, p2)

$$= \frac{(Y_1 - n_1 p_1)^2}{n p_1} + \frac{(Y_2 - n_2)^2}{n p_2}$$

What do we have here?

note: This is a large-sample approximation using CLT.
It is not an exact test!

We have shown this for k=2, and it can be shown in

	general for k categories that
	$Q = \frac{1}{2} \left(\frac{O(1-E(1)^2)}{E(1-1)} \right)$, $E(1-N)$
	Q = 2 = x (x-1) , E:= N.P. [=1 E: marginal EVs
	note: Kealls of data, subject to one
	linear constraint (pi+ - + pk=1)
	14.3 Goodness-of-fit test > df = k-1
	e.g. Mendel Genetics
	R: round pea Y yellow note: round and yellow
	r: wrinkled pea y: green carry the dominant genes
	If we cross two heterogenous parents: (Rryg x Rryg)
	we get the classic 9:3:3:1 ratio
	Yellow/ green genes from each purent
1	Re RY RY RY Ry
Count/wrinkle	
gence from each parent	
•	CC CY CY CY
	Our hypothesis then is for every 16 plants:
	9 round/gellow 3 round/green 3 winkled/gellow wrinkled green
0	16: P= 9/16 P2 = P3 = 3/16 P4= 1/16
0	Ha: at least one pi 7 po
	The state of the s

Space the observed counts are: 0, = 219 0, = 81 03 = 64 04 = 31 (n=400) Is this duta consistent w/ our hapothesis? E = 400(9/6) = 225 E2 = E3 = 3/6 (400) = 75 Ey = 1 (400) = 25 $Q = \sum_{i=1}^{4} \frac{(0i-e_i)^2}{e_i} \frac{(219-225)^2}{225} + \frac{(81-75)^2}{75}$ $+ (69-75)^2 + (31-25)^2 = 2.56 \sim \chi^2$ P3.05 = 7.81 do not i.e. the observed dute does not reject Ho give us any evidence younst the 9:3:3:1 ratio 7.51 note: Q = 0 means the observed counts matched the expected counts too closely (fisher/Mondel controversy) e.g. Death by horse kick (Prussian Calvary) · Ten army corps observed over 20 yrs (1875-1894) . no. of deaths by horse kick were recorded each year for each corps, giving us 200 observations

			1	
	no. of deaths	count	Pi	Ei
	0	109	.54335	108.67
		65	,33145	66.29
	2	22	.10110	20.22
	3	3	.02055	4.11
ollapse	٧ -	1	.00315	.63
	5 or more	0	P000.	. 08

Does this data follow a poisson Dist?

Use T to estimate) ...

1.e. on my, we observed , 61 who per year

$$\chi^2 = \frac{(109 - 108.67)^2}{108.67} + \dots + \frac{(4 - 4.82)^2}{4.82} = .3223$$

note: duta almost matches too well!

14.4 Contingency Tables

In many cases we are interested in determining if there is a relationship between 2 variables. When the variables are numerical, we can use linear regression techniques.

e.s. X= wt. of car Y= MPG

X = SAT Y= College GPA

When our variables are categorical in nature, we need to rely agam on counts are proportions in each cell

Example 14.3

Spec we want to evaluate the effectives of the flu vaccine in a small community. Data is collected and summarised in a contingency table

IV

No vaccine One shot Two shots

Flu 24 9 13

No Flu 289 100 565

In this case, we would like to know if the no. of shots has an influence on a patient's chance of getting flu.

We can think of this as a test of independence (between flw/no. of shots) or as a test for homogeniety of proportions i.e. the proportion of people who get the flu should be similar for no shot, I shot, or 2 shots if the flu vaccine is ineffective.

	Observed proportions of flu for each leve	el of shots:
	no vaccine one shat	
	Fly .077 .083	
	no Flu 923 .917	
	nd The	
	By inspection, we might survive the	at 2 shote siznificantly
	reduces someone's chance of contract	
	know if these differences are toola	
	to rankom chance?	
	The same of the sa	
	Consider on Txc contingency tabl	•
	columns	
	1 2 + 1 - 1 - 6	7
		n ₁ .
	2	N2.
الاسع	nr;	A .
		A
	r	Nr.
	Not Noz No	^
		,
	ncj = count in cell (i, i) > roul,	cal.
	No = Eth row total	
		rand total
	2 74	
		\(\frac{1}{2}\) \(\frac{1}{2}\
	n.; = jth col. total	, , , , , , , , , , , , , , , , , , ,
	= \(\frac{1}{2} \) \(\frac{1}{2} \)	
	(2)	

Poi = noi poi = noi poi = noi = 1 1 = 5 1:5 The joint dist. of (ni, niz, ..., nrc) is multinomial with Pi; for [=1 to r and j > 1 to C We want to test that rows are independent of columns (r.e. pe; = pe = P.;) note: no cell subject to one linear constraint from before ... 55 PUS =1 =26 = 10-1 $Q = \sum_{i=1}^{\infty} \sum_{j=1}^{\infty} (x_{ij} - n \cdot p_{ij})^2 \xrightarrow{D} \chi^2(x_{i-1})$ The reason we don't use this test statistiz, is that we don't know Pig. However, under Ho: Pig= Pi. x Pi 5 malependence Q(A/B) = P(A). P(B) So we need to estimate pio (1=1 tor) and poj (j=1 toc) Since & Pi = 1 and & Pis I we only need to estimate (1-1) + (1-1) parameters. Find the MLE estimates of pij under Ho T(billbis) = (u" ...uce). bil. ... bic a uniteromial 2-ros. fet

lul = la (nu ma)+ nulupin + · · · + nre lupre subject to constraint: [[[] pi; =] Under Ho: Pij = Pi. x Pij => ln L = ln ()+ n [ln p 1. + ln p . 1] + n 12 [ln p 1. + ln p . 2] + ··· + nre [Infro + Imfre] Subject to constructs: & Pi. =1 Iloj=1 To mak luly w.r.t. Pio's and Pojis which are subject to constraints, use La Grange multipliers Let K = In L R + X (1- \(Pi.) + M (1- \(Pi.) \) Take partials of k wint all pris, pijes, and I and u gh: bo gh: bil - m = 0

$$\sum_{i=1}^{n} n_i = \sum_{i=1}^{n} \lambda \cdot p_i, \quad \Rightarrow \quad n = \lambda \cdot \left(\sum_{i=1}^{n} p_i, \lambda \right) \Rightarrow n = \lambda$$

Let Ocj = "observed count in row i, coli" (Ocj = nij)

$$\Rightarrow Q = \frac{\sum_{i=1}^{n} \sum_{j=1}^{n} (Ni\hat{s} - N.\hat{p}_i.\hat{p}_i)^2}{N.\hat{p}_i.\hat{p}_i.\hat{p}_i}$$

	$r_{c-1} - (r_{-1}) - (c_{-1}) = r_{c-r-c} + 1 = (r_{-1}) \times 1$	((-1)
	r(-1-(r-1)-12-1) - 12-1-2+1 - (1-1)	cc-1 _j
	This is the x2 test statistic from 2040/2050!	
	It is based on the CLT and is an approximation	1
	DI 15 Sages of The CET and 15 mor approximation	
	Back to the flue example:	
	we compute the expected courts (in parenthese	()
		3,
	using Ec; = 12 x cs	
	2 2 2 2	
_	no vaccine I shot 2 shots	46
_	Fly 24 (14.4) 9 (5.0) 13 (26.6)	
	10 Flu 284 (248.6) 100 (104.0) 565 (551.4)	954
	313 109 578	1000
	Ho: Independence between vaccine and contractor	ig the Fly
	the: Relationship between vaccine and contracting	
	2 (24-14.4)2 (9-5)2 (565-55	1,412
	X = 14.4 5 551.4	- Indiana
	2 17.35	
	x=== 5.491	
i	1-(2),05	

17.3575.991 = Reject Ho What do we conclude?

	going back to the observed proportions I would conclude
	no vaccine 1 shot 2 shots
	Flu .077 .083 ,0225
	no Plu ,923 ,917 ,9775
	Taking 2 flu shots significantly reduces your chance
	of contracting the flu. Taking the flu shot only once
	shows no exidence of being effective.
	STOUS NO EVIDENZE DI SETTIO
	Rolationship to large-sample 2-test for pivs.pz
	Example 2
	e.g. Is there a relationship between smoking and lung concer?
	Lung problem
	Yes No
	Yas (22.5) 35 (22.5)10 45
Smoking	No (27.5) 15 (27.5) 40 55
0	50 50 100
	note: ps = 35 = .778 pns = 55 = .273
	12 test: Ho: independence between smoking and ling problems
	Ha: relationship exists between smoking and lung problems
	$\gamma^2 = (35 - 22.5)^2 + (10 - 22.5)^2 + (15 - 27.5)^2$ $22.5 \qquad 22.5 \qquad 27.5$
	22.5 22.5 27.5
	(40-27.5)2 = 25.2525 p-value = .0000005
	, +92

27.5

3) Reject to i.e. lung problems and smoking are related
2-test for pivs. pa.
140: Ps = PMS
Ha: Ps 7 PMS
$\hat{p}_{s} = \frac{35}{15} = .778$ $\hat{p}_{AS} = \frac{15}{55} = .273$
A = 35+15 = .50
$\frac{(.778273)}{2} = 5.025189 p-value = .0000005$ $\frac{1}{45} = \frac{1}{55}$
$\frac{1}{(.5)(.5)}\left(\frac{1}{45} + \frac{1}{55}\right)$
2 ² = (5.025189) = 25.2525
2 = (5.025184) = 25.2525
These are the same tests! 22 ~ X2(1)

ı	
	14.5 rxc Tables with Fixed low or Column Totals
	In many applications, we might want to fix the no.
	we have in each column.
İ	e.g. R.S. a fixed no. of people who had no vaccine,
1	
	one shot, or two shots
	Long the transfer of the trans
1	What does this do to our test statistic? Keep in mind that
1	we still want to tost if the proportion of those who get flu
1	B the same as we move from one column to the next.
	In this case, we could consider the no. who get the flue, nis,
İ	
İ	to be binongal. ?
+	No vaccine one shot Two shots
	fla MII NIZ MI3
1	no flu
	n., n.2 n.3
Ī	
1	Fxed
	N11~Bin (N-1, PIN) N12~Bin (N-2, P12) N13~Bin (N-3, P13)
+	
	Under 16: p11 = p12 = p13 = p
	If we extend this to more than 2 rows, then we have
	the courte or each column following a multipose last

In this case, Ho: Pii=Pi., Pri=Pr.

Solving (+1) egas. simultaneously

This is the same statistic!

Survey of voter sentiment in 4 mideity political wards

Is there sufficient evidence to indicate that voting preferences differ in the 4 awards?

Ward
(.38) (.245) (.245) (.245) (.245) Favor A 76 (59) 53(59) 59 (59) 48 (.245)
don't favor 124 (141) 147 (141) 141 (141) 152 564 (.705)
4 200 200 200 800
note: Observed proportions: .38 vs265 vs295 vs24 One there differences too large to be attributable to chance alone?
Ho: p=p2=p3=p4 Ita: at least one pi is different
x2 = (76-54)2 (53-54)2 + + (152-141)2
59 59 141
= 10.72
23,05 = 7.815 => Reject to i.e. we halieve there
are real differences in voter preferences
between wards.

¥

16.2 Bayesian Priors, Poseteriors and Estymators

To this point, we have assumed we have no information regarding the unknown parameter &, until we collect sample information.

e.g. We want to estimate the proportion of crystal math addicts in Utah County. We know this should be small.

The idea of Bayesian statistics, is that if we make an mittal assumption regarding Q, (r.e. utilize "prior info on O"), then our estimator after collecting sample duta will be hetter than if we had assumed no prior info at all on O was available.

Bayesian process:

Let L(y, yz, ..., yn) a) denote the likelihood fet for a r.s. of site n. This represents the density of our sample data gives Q.

In Bayesian analysis, O is viewed as a RV itself with prior dist. or prior density fet g(B).

=> f(y1, ..., yn, 0) = L(y1, y2, ..., yn 10) x g(0) is the junt density fet. of y1 ... yn, 0.

Recall: f(12/31) = f(41,42) f(41) = f(41,42) dy2

=> the morganal density of our sample data (4, -- 7n) is

m(y, ... yn) = \$ L(y, ... yn 10) x 2(0) lo

Therefore, the posterior density of @ | 31 ... In is

5"(0 | 51,42, ...,4n) = L(41, ...,4n) = > (0) × 9(0)

note: The posterior density contains all pertinent into about &: (1) prior mto on & (2) sample mto on &

Example 16.1

we would like to estimate p = proportion of pop afflicted with a certain disease. We need to assume a prior dist. for p, and proportions are often modeled with a Beta dist.

Recall: Y~ Beta (x, B)

f(y) = [(x+13) y (1-y) 5-1, 0<y<1

E(4)= x+13 (x+13)2(x+13+1) note: We will worry about the choice of of, B later Our sample Lota: Y, ... Yn Bars from YE - Bin (1,p) 1.1. each person either has the disease or they don't bearlb) = bas (1-b) 1-20, 12:2011 >> F(21 ... 2 = 16) = 1 b2: (1-6) -2. P (1-p) prior dist for p: g(p) = ((x+p) p" (1-p) B-1 (1d) [1B) so gt. density of Tinta, p is: f(y1 --- yn, p) = p = y= (1-p) - 1 (x+B) px-1 (1-p) B-1 => marginal density of 1, -- 4p is: m(y, ... y) = \ [(a+B) p (1-p) &p 5 (a). (B) let a = Egita, B'= n Egits

	= P(x+B) P(Isita). P(N-EsitB) note: loss not
	Pla)-PiB) P(nta+B) depend on P
	(11216)
	The posterior density of p given our sample eata:
	3*(p y, -yn)=
	55:+a-1 n-55:+B-1
	P(a+B) P (1-B)
	2181.187
	Plata Plating ((Esta) Planesy:+ ()
	[(a) [7]B) [(nta+b)
	7
	(2 y + a) -1 (n- 2 y + 3) -1
	[(n+x+B) (1-p)
	,06941
	[([5:+a) · [(n-27:+B)
nbeta (x', B')	
x== 25:+0	0
0 = n = 242+B	
30+0	T.e. the posterior dist of ply, yo is also a beta dist.
	This means that the seta dist. is a conjugate prior dist.
	for a bernoulli or binomial dist.
	TO A SECURICITY OF SIMOMARY CHET.
or or lesson	to E(alymona) = Evita
6 ozteren M	ent $E(p y_1y_n) = \frac{Ey_i + \alpha}{n + \alpha + \beta}$
	71) 51 (2
Noone a	× × × × × × × × × × × × × × × × × × ×
ALLIB! ME	E(P) = a+B

If sy: is high see a lot of people in our sample had the disease, the expected value of p 1 (vice versa of Eti coult) Spee we think p≈. 25 and use p~ seta (K=1, B=3) => E(p) = 1+3 =.25 , V(p) = 1/3) = .0375 If N=250, 245=10 (1.e. p=.40) => x = 10+1=11 | b = (25-10)+3 = 18 posterior menn = 11 = 379) posterior vurinnee = (.0078) note: \$= 10/25 = .40 V(\$) = .4(16) = .0096 If N= 100 | £4; = 40 >> x = 40+1=41 (3 = (100-40)+3=63 posterior mean = 41 = (394) posterior variance = (0023) note: as not, more wto is applied to sample daity Spee we think px.25 but use p~ beta (x=10, B=30) 57 E(p) = 10+30 = .25 , vcp) = (0(30) = 10+30+1) = .00457

If n=25, 241 = 10 20 x = 10+10=20 , B# = 25-10+30=45 20 +45 (307) posterior verionce 2,00323 # N=100, Ey =40 => 0x = 40+10=50 , B = (100-40) + 30 = 90 Pasterrar mean = 50 = (357) Pasterrar variance = (0016 note: using a prior dist. of beta (10,30) means that the posterior estimate for p will not be as "reactive" to the sample duta. However, it will move closer to the sample estimate as nT

see graphs mR

Def 16.2 Let Y, - In be a r.s. w/ likelihood fet L(y, yz, , sn) 0), and let @ have prior density g(0). The posterior Bayes estimator t10) is given by

The postorior Buyes estimator of & is