A diagnostic plot for assessing model fit in count data models

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Rennes, 5th July 2016





Introduction

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- Assume that a routine to obtain estimates $\hat{\mu}_i = \hat{E}(Y_i|x_i)$ and $\hat{\theta}_i$ is readily available.
- ▶ Denote N(k), for k = 0, 1, 2, ..., the number of observed counts k in $y_1, ..., y_n$.
- ▶ Idea: check whether, for each count k = 0, 1, 2, ..., the number N(k) is 'plausible' under the distribution $F(\hat{\mu}_i, \hat{\theta}_i)$.

Poisson-Binomial distribution

► The random variable N(k) follows a Poisson–Binomial distribution with parameters $p_1(k), \ldots, p_n(k)$, where

$$p_i(k) = P(k|\mu_i, \theta_i)$$

is the probability of observing the count k under covariate x_i and model F (Chen and Liu, 1997).

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- ▶ The $p_i(k)$ can be estimated by $\hat{p}_i(k) = P(k|\hat{\mu}_i, \hat{\theta}_i)$ from the fitted model.
 - For instance, in the special case that $F(\mu_i, \theta_i)$ corresponds to $Pois(\mu_i)$, one has $\hat{p}_i(k) = \exp(-\hat{\mu}_i)\hat{\mu}_i^k/k!$.
 - This scenario was discussed in the previous talk with focus on the case k = 0.
 - ▶ This talk generalizes those ideas to general *k* and *F* and proposes a generic diagrammatic tool.

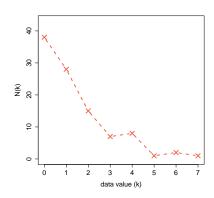
Plausibility intervals for N(k)

- ▶ Knowing the distribution of N(k), one can derive intervals of plausible values of N(k) by considering appropriate quantiles from this distribution.
- For fixed k, appropriate lower and upper quantiles, say $q_{\alpha/2}(k)$ and $q_{1-\alpha/2}(k)$ of the Poisson–Binomial distribution can be computed using the R package poibin (Hong, 2013).
- ▶ Do this for a range of values of k, and plot intervals $(q_{\alpha/2}(k), q_{1-\alpha/2}(k))$ alongside observed values N(k) as a function of k.

Example: simulated data

▶ n = 100 observations y_1, \ldots, y_n simulated from a Zero–inflated Poisson (ZIP) distribution with Poisson parameter $\mu = 1.5$ and zero–inflation parameter p = 0.2

k	N(k)	
0	38	
1	28	
2	15	
3	7	
4	8	
5	1	
6	2	
7	1	



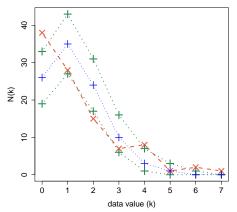
Example: simulated data

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- ▶ Consider $F(\mu) \sim \text{Pois}(\mu)$ with $\hat{\mu} = \bar{y}$.

k	N(k)	$q_{0.05}(k)$	$q_{0.95}(k)$	
0	38	19	33	- 4 - X
1	28	27	43	8 + 1
2	15	17	31	* ,
3	7	6	16	N(K)
4	8	1	7	* +
5	1	0	3	2-
6	2	0	1	# T.
7	1	0	0	0 1 2 3 4 5 6 7
				data value (k)

Median-adjustment

- ► The previous graph can be difficult to read if the sample size is large, and so the bounds get very tight.
- We therefore adjust it by subtracting the medians M(k) = med(N(k)) from all values, where the median is taken wrt to the Poisson-Binomial distribution of N(k).



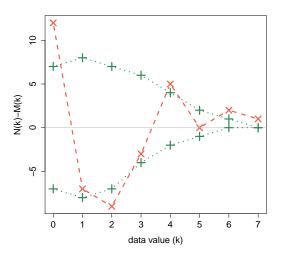
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k	N(k)	M(k)	N(k)-M(k)	$q_{0.05}(k)$ – $M(k)$	$q_{0.95}(k) - M(k)$
0	38	26	12	-7	7
1	28	35	-7	-8	8
2	15	24	-9	-7	7
3	7	10	-3	-4	6
4	8	3	5	-2	4
5	1	1	0	-1	2
6	2	0	2	0	1
7	1	0	1	0	0

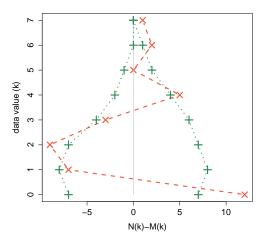
Median-adjusted bounds

▶ Diagnostic plot for the accuracy of the Poisson assumption.



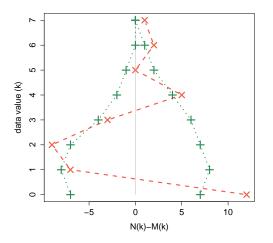
Median-adjusted bounds: Variant

Exchange horizontal and vertical axis:



Median-adjusted bounds: Variant

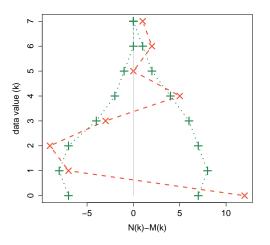
Exchange horizontal and vertical axis:



'Christmas tree diagram'.

Median-adjusted bounds: Variant

Exchange horizontal and vertical axis:



- 'Christmas tree diagram'.
- Adequate models have the 'decoration' inside the tree.



Example: Biodosimetry data

► Frequency of dicentric chromosomes in human lymphocytes after *in vitro* exposure to doses between 1 and 5Gy of 200kV X–rays. The irradiated blood was mixed with non–irradiated blood in a proportion 1:3 in order to mirror a partial body exposure scenario.

	Frequency of counts								
dose	0	1	2	3	4	5	6	7	8
1	2713	78	8	0	1	0	0	0	0
2	1302	71	22	5	0	0	0	0	0
3	1116	46	28	7	2	1	0	0	0
4	929	18	14	22	13	2	0	1	1
5	726	17	18	12	9	13	1	4	0



- ▶ These are n = 7200 observations of the type (dose_i, y_i), with y_i being a count in 0, ..., 8.
- X-rays are sparsely ionizing the literature suggests a quadratic dose model in this case.

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- Link function:
 - Cytogenists prefer identity link.
 - ▶ Being among Statisticians, I will use the log link.
- Response (count) distribution:
 - It is widely accepted that the number of dicentrics in irradiated blood samples is Poisson distributed.
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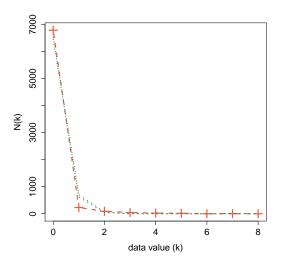
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 - However, under a partial body exposure scenario, we would expect a deviation from the Poisson assumption, towards zero-inflation.
- ▶ Consider the initial model $y_i | dose_i \approx Pois(\mu_i)$ with

$$\mu_i \equiv E(y_i | \mathsf{dose}_i) = \exp(\beta_0 + \beta_1 \mathsf{dose}_i + \beta_2 \mathsf{dose}_i^2)$$



Diagnostics for biodosimetry data

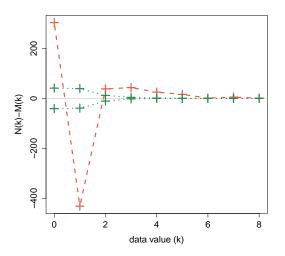
...without median— adjustment:



hard to see anything!

Diagnostics for biodosimetry data

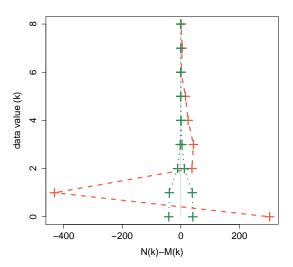
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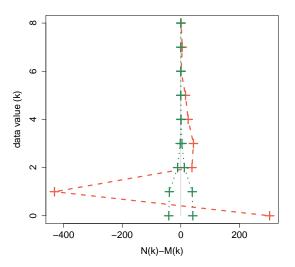
much better!



Christmas tree diagram: Poisson hypothesis

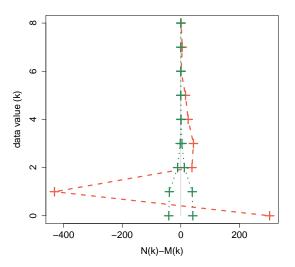


Christmas tree diagram: Poisson hypothesis



We clearly observe zero-inflation (and associated 1-deflation);

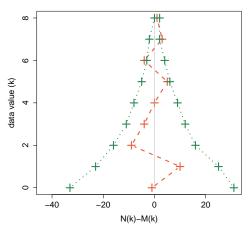
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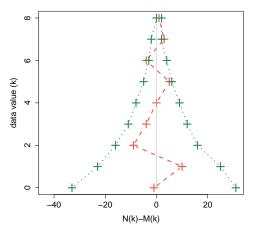
Christmas tree diagram: ZIP hypothesis

▶ Do all the same as before, but now compute $\hat{\mu}_i$, $\hat{\theta}_i$, and $\hat{p}_i(k)$, using the zero–inflated Poisson (ZIP) model as the hypothesized model.



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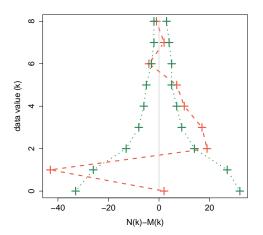
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indicates a good fit.

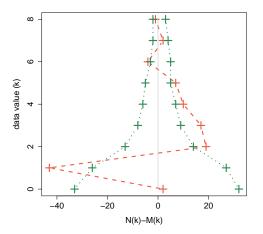
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Repeat the procedure using the negative Binomial model as the hypothesized model.



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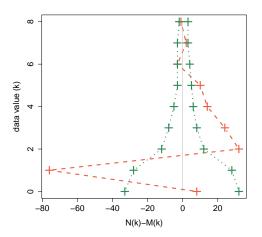


indicates that the NB model does not capture the data well.



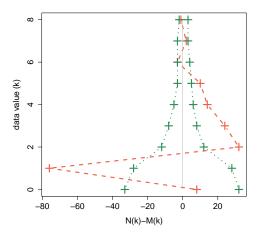
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▶ Repeat the procedure using the Poisson inverse Gaussian (PIG) model as the hypothesized model.



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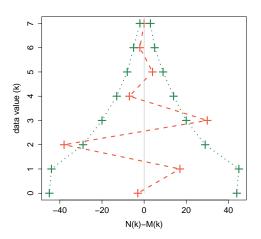


Alternative data set: Whole body exposure

Counts of dicentric chromosomes in 4400 blood cells after in vitro 'whole body' exposure with 200kV X-rays from 0 to 4.5Gy.

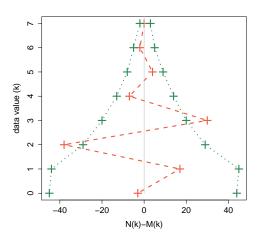
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indicates that Poisson model is fairly reasonable



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- That is, exceeding the boundary limits once or twice should not necessarily be interpreted as rejection of the hypothesized count distribution, as long as the 'decoration' is reasonably consistent with the tree.



Comparison with score tests

- ► Alternatively, one can carry out traditional score tests.
- ▶ For instance, consider H_0 : Poisson versus H_1 : ZIP or H_1 : NB.
- Score test statistic $T = S^T J^{-1} S$, where S and J are the score function and Fisher Information matrix (resp.) evaluated under the Poisson model. Asymptotically, $T \sim \chi^2(1)$.
- ▶ Resulting values of T, to be compared with $\chi^2_{1,0.95} = 3.84$ (Oliveira et al, 2016):

Test	Body exposure			
	Partial	Whole		
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- ► Confirms that Poisson is adequate for whole body exposure but inadequate for partial body exposure.
- ...but the score test does not tells us whether it's at all the zero's which cause the problem, nor whether the data are zero-inflated or -deflated!



Conclusion

- We have provided a simple diagrammatic tool to assess the adequacy of any given count data model.
- Essentially, it is verified whether the frequency, N(k), of each count, k, is plausible given the hyptothesized model.
- Can be used for with or without covariates.
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- ▶ Be aware of multiple testing: It is a diagram, not a test.



References

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