

## ZERO-TRUNCATED NEGATIVE BINOMIAL | SAS DATA ANALYSIS EXAMPLES

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**Version info:** Code for this page was tested in SAS 9.3.

Zero-truncated negative binomial regression is used to model count data for which the value zero cannot occur and when there is evidence of over dispersion .

**Please Note:** The purpose of this page is to show how to use various data analysis commands. It does not cover all aspects of the research process which researchers are expected to do. In particular, it does not cover data cleaning and verification, verification of assumptions, model diagnostics and potential follow-up analyses.

### Examples of zero-truncated negative binomial

Example 1.

A study of the length of hospital stay, in days, as a function of age, kind of health insurance and whether or not the patient died while in the hospital. Length of hospital stay is recorded as a minimum of at least one day.

Example 2.

A study of the number of journal articles published by tenured faculty as a function of discipline (fine arts, science, social science, humanities, medical, etc). To get tenure faculty must publish, i.e., there are no tenured faculty with zero publications.

Example 3.

A study by the county traffic court on the number of tickets received by teenagers as predicted by school performance, amount of driver training and gender. Only individuals who have received at least one citation are in the traffic court files.

### Description of the data

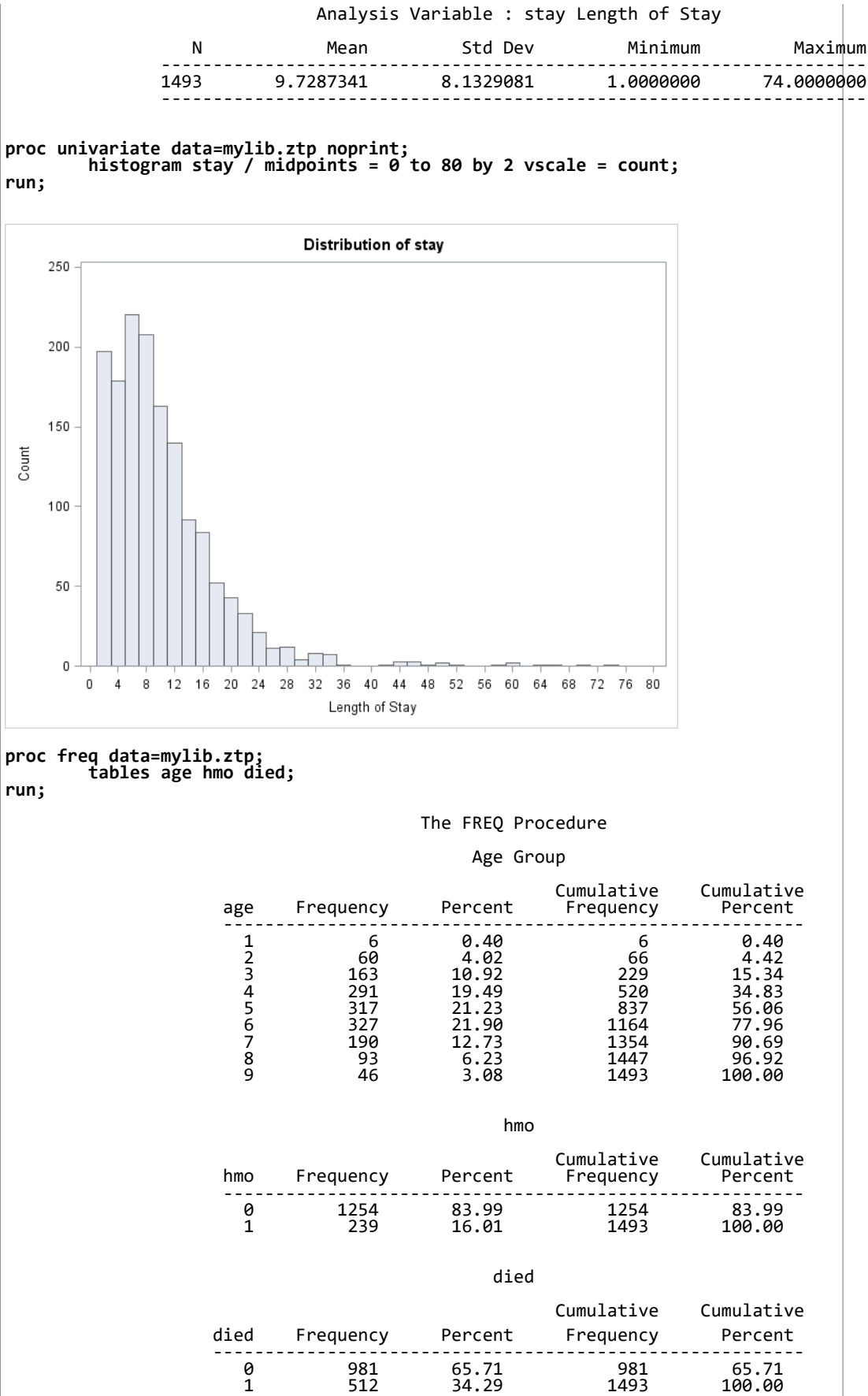
Let's pursue Example 1 from above.

We have a hypothetical data file, **ztp.sas7bdat** with 1,493 observations available [here](#) ([/data/ztp.sas7bdat](#)) . The variable describing length of hospital visit is **stay**. The variable **age** gives the age group from 1 to 9 which will be treated as interval in this example. The variables **hmo** and **died** are binary indicator variables for HMO insured patients and patients who died while in hospital, respectively. These are the same data as were used in the [ztp](https://stats.idre.ucla.edu/sas/dae/zero-truncated-poisson-regression/) (<https://stats.idre.ucla.edu/sas/dae/zero-truncated-poisson-regression/>) example.

Let's look at the data.

```
proc means data=mylib.ztp;  
    var stay;  
run;
```

The MEANS Procedure



## Analysis methods you might consider

Before we show how you can analyze these data with a zero-truncated negative binomial analysis, let's consider some other methods that you might use.

- Zero-truncated Negative Binomial Regression – The focus of this web page.
- Zero-truncated Poisson Regression – Useful if there is no overdispersion in the zero truncated variable. See the Data Analysis Example for [ztp](https://stats.idre.ucla.edu/sas/dae/zero-truncated-poisson-regression/) (<https://stats.idre.ucla.edu/sas/dae/zero-truncated-poisson-regression/>).
- Negative Binomial Regression – Ordinary negative binomial regression will have difficulty with zero-truncated data. It will try to predict zero counts even though there are no zero values.
- Poisson Regression – The same concerns as for negative binomial regression, namely, ordinary poisson regression will have difficulty with zero-truncated data. It will try to predict zero counts even though there are no zero values.
- OLS Regression – You could try to analyze these data using OLS regression. However, count data are highly non-normal and are not well estimated by OLS regression.

## Zero-truncated negative binomial regression using proc nlmixed

In order to use **proc nlmixed** to perform truncated negative binomial regression, we must supply it with a likelihood function. The probability that an observation has count ( $y$ ) under the negative binomial distribution (without zero truncation) is given by the equation:  $P(Y=y) = \frac{\Gamma(\alpha+1)}{\Gamma(\alpha+1)\Gamma(y+1)} \left(\frac{\mu}{1+\mu}\right)^{\alpha+1} \left(\frac{1}{1+\mu}\right)^y$  where ( $\alpha$ ) is the overdispersion parameter and ( $\mu$ ) is the mean of the negative binomial distribution. With zero truncation, we calculate the probability that ( $Y=y$ ) conditional on ( $Y>0$ ), that is, that ( $Y$ ) is observed as 0 values are not observed. The probability of a zero count under the negative binomial distribution is  $P(Y=0) = \frac{\Gamma(\alpha+1)}{\Gamma(\alpha+1)\Gamma(1)} \left(\frac{\mu}{1+\mu}\right)^{\alpha+1} \left(\frac{1}{1+\mu}\right)^0$ . The conditional probability is then:  $P(Y=y|Y>0) = \frac{P(Y=y)}{1-P(Y=0)} = \frac{\frac{\Gamma(\alpha+1)}{\Gamma(\alpha+1)\Gamma(y+1)} \left(\frac{\mu}{1+\mu}\right)^{\alpha+1} \left(\frac{1}{1+\mu}\right)^y}{1 - \frac{\Gamma(\alpha+1)}{\Gamma(\alpha+1)\Gamma(1)} \left(\frac{\mu}{1+\mu}\right)^{\alpha+1} \left(\frac{1}{1+\mu}\right)^0}$ . The log-likelihood function for the zero-truncated negative binomial distribution is thus:

$$l = \sum_{i=1}^n \left[ \log \Gamma(\alpha+1) - \log \Gamma(y_i+1) - \log \Gamma(\alpha+1) - \frac{\alpha}{\mu} \log(1+\mu) + y_i \log(\mu) - y_i \log(1+\mu) - \log \left( 1 - \left( \frac{\mu}{1+\mu} \right)^{\alpha+1} \right) \right]$$

In negative binomial regression, we model ( $\log(\mu)$ ), the log of the mean (expected counts), as a linear combination of a set of predictors:  $\log(\mu) = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \beta_3 x_3$ . We supply the last two equations to **proc nlmixed** to model our data using a zero-truncated negative distribution. Additionally, **proc nlmixed** does not support a **class** statement, so categorical variables should be dummy-coded before running the analysis. Unlike other SAS procs, by default the first group is the reference group by default in **proc nlmixed**.

```
proc nlmixed data = mylib.ztp;
  log_mu = intercept + b_age*age + b_died*died + b_hmo*hmo;
  mu = exp(log_mu);
  het = 1/alpha;
  ll = lgamma(stay+het) - lgamma(stay + 1) - lgamma(het) - het*log(1+alpha*mu)
      + stay*log(alpha*mu) - stay*log(1+alpha*mu) - log(1 - (1 + alpha * mu)**-het);
```

run;									
model stay ~ general(11);									
The NL MIXED Procedure									
Specifications									
Data Set					MYLIB.ZTP				
Dependent Variable					stay				
Distribution for Dependent Variable					General				
Optimization Technique					Dual Quasi-Newton				
Integration Method					None				
Dimensions									
Observations Used					1493				
Observations Not Used					0				
Total Observations					1493				
Parameters					5				
Parameters									
intercept	b_age	b_died	b_hmo	alpha	NegLogLike				
1	1	1	1	1	10136.7274				
Iteration History									
Iter	Calls	NegLogLike	Diff	MaxGrad	Slope				
1	3	5203.75757	4932.97	1718.332	-825332				
2	6	5130.65185	73.10572	212.6078	-12208.4				
3	8	4922.88698	207.7649	1701.184	-735.733				
4	9	4862.95248	59.9345	176.3689	-177.538				
5	11	4851.81702	11.13546	393.0774	-13.9647				
6	12	4838.27102	13.546	179.7832	-7.96192				
7	16	4788.46175	49.80926	168.3697	-26.6674				
8	17	4774.94754	13.51421	105.3687	-117.309				
9	18	4759.72531	15.22222	77.4436	-25.9074				
10	20	4755.95435	3.77096	85.88275	-22.2361				
11	22	4755.3438	0.610557	39.18804	-2.65095				
12	24	4755.29354	0.050252	30.83521	-0.14278				
13	26	4755.28066	0.012889	3.944229	-0.03589				
14	28	4755.28014	0.000512	0.44716	-0.00416				
15	29	4755.27964	0.0005	0.195745	-0.00109				
16	31	4755.27962	0.000028	0.007496	-0.00006				
17	33	4755.27962	1.109E-7	0.030916	-4.12E-7				
NOTE: GCONV convergence criterion satisfied.									
The SAS System				09:40 Monday, June 4, 2012 5					
The NL MIXED Procedure									
Fit Statistics									
-2 Log Likelihood				9510.6					
AIC (smaller is better)				9520.6					
AICC (smaller is better)				9520.6					
BIC (smaller is better)				9547.1					
Parameter Estimates									
Parameter	Estimate	Standard Error	DF	t Value	Pr >  t	Alpha	Lower	Upper	Gradient
intercept	2.4083	0.07198	1493	33.46	<.0001	0.05	2.2671	2.5495	-0.00749
b_age	-0.01569	0.01311	1493	-1.20	0.2314	0.05	-0.04140	0.01002	-0.03092
b_died	-0.2178	0.04616	1493	-4.72	<.0001	0.05	-0.3083	-0.1272	0.000972
b_hmo	-0.1471	0.05922	1493	-2.48	0.0131	0.05	-0.2632	-0.03090	-0.00018
aAlpha	0.5663	0.03123	1493	18.13	<.0001	0.05	0.5050	0.6276	-0.00461

The output looks very much like the output from an OLS regression:

- Close to the top is the iteration log giving the values of the log likelihoods starting with a model that has no predictors. The last value in the log (4755.27962) is the final value of the negative log

likelihood for the full model and is repeated below.

- Next comes a number of fit statistics, which can be used to compare the fit of nested models.
- Below the header you will find the zero-truncated negative binomial coefficients for each of the variables along with standard errors, t-values, p-values and 95% confidence intervals for each coefficient.
- Below that, is the the overdispersion parameter **alpha** along with its standard error, t-value, p-value, and 95% confidence interval.

Looking through the results we see the following:

- The value of the coefficient for **age**, -.01569, suggests that the log count of stay decreases by .01569 for each unit increase in age group. This coefficient is not statistically significant.
- The coefficient for **hmo**, -.1471, is significant and indicates that the log count of stay for HMO patient is .1471 less than for non-HMO patients.
- The log count of stay for patients that **died** while in the hospital was .2178 less than those patients that did not die.
- The value of the constant (**intercept**), 2.4083 is log count of the stay when all of the predictors equal zero.
- The estimate for **alpha** is .5663. For comparison, a model with an **alpha** of zero is equivalent to a zero-truncated poisson model. In this model, **alpha** is statistically different from zero, suggesting that the negative binomial model is a better choice than a poisson model.

We can also use **estimate** statments to help understand our model, by examining the predicted or expected length of stay of patients with varying covariate values. For example we can predict the expected number of days spent at the hospital across age groups for the two hmo statuses for patients who died. The **estimate** statement for **proc nlmixed** works slightly differently from how it works within other procs. Here, each parameter must be explicitly multiplied by the value at which is to be held for that **estimate** statment, and expressions are allowed, such as exponentiation (see code below). Because we would like to predict actual number of days rather than log number of days, we need to exponentiate the estimate. Additionally, the following expected counts are unconditional, meaning these are the expected lengths of stay for patients with the given covariate values in the entire population, not for those patients who we know have stayed at least one day in the hospital (the conditional expectation).

```
proc nlmixed data = mylib.ztp;
  log_mu = intercept + b_age*age + b_died*died + b_hmo*hmo;
  mu = exp(log_mu);
  het = 1/alpha;
  ll = lgamma(stay+het) - lgamma(stay + 1) - lgamma(het) - het*log(1+alpha*mu)
      + stay*log(alpha*mu) - stay*log(1+alpha*mu) - log(1 - (1 + alpha * mu)**-het);
```

```

model stay ~ general(ll);
estimate 'age 1 died 1 hmo 0' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 0);
estimate 'age 1 died 1 hmo 1' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 1);
estimate 'age 3 died 1 hmo 0' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 0);
estimate 'age 3 died 1 hmo 1' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 1);
estimate 'age 5 died 1 hmo 0' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 0);
estimate 'age 5 died 1 hmo 1' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 1);
estimate 'age 7 died 1 hmo 0' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 0);
estimate 'age 7 died 1 hmo 1' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 1);
estimate 'age 9 died 1 hmo 0' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 0);
estimate 'age 9 died 1 hmo 1' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 1);
run;
< **SOME OUTPUT OMITTED** >

```

Additional Estimates									
Label	Estimate	Standard Error	DF	t Value	Pr >  t	Alpha	Lower	Upper	
age 1 died 1 hmo 0	8.8010	0.6291	1493	13.99	<.0001	0.05	7.5668	10.0351	
age 1 died 1 hmo 1	7.5974	0.6545	1493	11.61	<.0001	0.05	6.3135	8.8813	
age 3 died 1 hmo 0	8.5290	0.4385	1493	19.45	<.0001	0.05	7.6688	9.3893	
age 3 died 1 hmo 1	7.3627	0.5212	1493	14.13	<.0001	0.05	6.3404	8.3850	
age 5 died 1 hmo 0	8.2655	0.3256	1493	25.39	<.0001	0.05	7.6268	8.9042	
age 5 died 1 hmo 1	7.1352	0.4498	1493	15.86	<.0001	0.05	6.2530	8.0174	
age 7 died 1 hmo 0	8.0101	0.3430	1493	23.35	<.0001	0.05	7.3373	8.6830	
age 7 died 1 hmo 1	6.9147	0.4540	1493	15.23	<.0001	0.05	6.0242	7.8052	
age 9 died 1 hmo 0	7.7627	0.4586	1493	16.93	<.0001	0.05	6.8630	8.6623	
age 9 died 1 hmo 1	6.7011	0.5200	1493	12.89	<.0001	0.05	5.6811	7.7211	

The expected stay for non-HMO patients in age group 1 who died was 8.8010 days, while it was 7.5974 days for HMO patients in age group 1 who died.

We can also test whether the effect of HMO is significant at each level of age for patients who died. We can simply subtract the two estimates that vary by hmo at each level of age.

```

proc nlmixed data = mylib.ztp;
log_mu = intercept + b_age*age + b_died*died + b_hmo*hmo;
mu = exp(log_mu);
het = 1/alpha;
ll = lgamma(stay+het) - lgamma(stay + 1) - lgamma(het) - het*log(1+alpha*mu)
+ stay*log(alpha*mu) - stay*log(1+alpha*mu) - log(1 - (1 + alpha * mu)**-het);
model stay ~ general(ll);
estimate 'age 1 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 0);
estimate 'age 3 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 0);
estimate 'age 5 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 1);
estimate 'age 7 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 0);
estimate 'age 7 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 1);
estimate 'age 7 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 0);
estimate 'age 7 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 1);
estimate 'age 9 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 0);
estimate 'age 9 died 1 hmo 0 vs 1' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 1);
run;
< **SOME OUTPUT OMITTED** >

```

Additional Estimates									
Label	Estimate	Standard Error	DF	t Value	Pr >  t	Alpha	Lower	Upper	
age 1 died 1 hmo 0 vs 1	1.2036	0.4698	1493	2.56	0.0105	0.05	0.2820	2.1251	
age 3 died 1 hmo 0 vs 1	1.1664	0.4511	1493	2.59	0.0098	0.05	0.2816	2.0512	
age 5 died 1 hmo 0 vs 1	1.1303	0.4350	1493	2.60	0.0095	0.05	0.2770	1.9837	
age 7 died 1 hmo 0 vs 1	1.0954	0.4215	1493	2.60	0.0094	0.05	0.2686	1.9222	
age 9 died 1 hmo 0 vs 1	1.0616	0.4103	1493	2.59	0.0098	0.05	0.2568	1.8664	

The effect of **hmo** is significant for each **age** group tested.

It may be illustrative for us to plot the predicted number of days stayed as a function of age and hmo status. To do this, we must tell SAS to save this table of predicted values as a dataset. Tables and graphics

produced by procedures are given names upon creation. We will need the name of this prediction table to tell SAS to save it. Place **ods trace on** and **ods trace off** statements around the procedure which produced this table to obtain its name. Output from the **ods trace** statements is located in the log, not the output.

```
ods trace on;
proc nlmixed data = mylib.ztp;
  log_mu = intercept + b_age*age + b_died*died + b_hmo*hmo;
  mu = exp(log_mu);
  het = 1/alpha;
  ll = lgamma(stay+het) - lgamma(stay + 1) - lgamma(het) - het*log(1+alpha*mu)
      + stay*log(alpha*mu) - stay*log(1+alpha*mu) - log(1 - (1 + alpha * mu)**-het);
  model stay ~ general(ll);
  estimate 'age 1 died 1 hmo 0' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 0);
  estimate 'age 1 died 1 hmo 1' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 1);
  estimate 'age 3 died 1 hmo 0' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 0);
  estimate 'age 3 died 1 hmo 1' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 1);
  estimate 'age 5 died 1 hmo 0' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 0);
  estimate 'age 5 died 1 hmo 1' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 1);
  estimate 'age 7 died 1 hmo 0' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 0);
  estimate 'age 7 died 1 hmo 1' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 1);
  estimate 'age 9 died 1 hmo 0' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 0);
  estimate 'age 9 died 1 hmo 1' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 1);
run;
ods trace off;

<***SOME OF THE LOG OMITTED***>

Output Added:
-----
Name:      AdditionalEstimates
Label:     Additional Estimates
Template:  Stat.NLM.AdditionalEstimates
Path:      Nlmixed.AdditionalEstimates
-----
NOTE: PROCEDURE NLMIXED used (Total process time):
      real time           0.23 seconds
      cpu time            0.17 seconds

105  ods trace off;
```

Towards the end of the log we find the name of this table, which as expected by its heading in the output above, is "AdditionalEstimates". We can now tell SAS to save this output table as the dataset "mylib.addest" using an **ods output** statement.

```
ods output AdditionalEstimates = mylib.addest;
proc nlmixed data = mylib.ztp;
  log_mu = intercept + b_age*age + b_died*died + b_hmo*hmo;
  mu = exp(log_mu);
  het = 1/alpha;
  ll = lgamma(stay+het) - lgamma(stay + 1) - lgamma(het) - het*log(1+alpha*mu)
      + stay*log(alpha*mu) - stay*log(1+alpha*mu) - log(1 - (1 + alpha * mu)**-het);
  model stay ~ general(ll);
  estimate 'age 1 died 1 hmo 0' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 0);
  estimate 'age 1 died 1 hmo 1' exp(intercept * 1 + b_age * 1 + b_died * 1 + b_hmo * 1);
  estimate 'age 3 died 1 hmo 0' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 0);
  estimate 'age 3 died 1 hmo 1' exp(intercept * 1 + b_age * 3 + b_died * 1 + b_hmo * 1);
  estimate 'age 5 died 1 hmo 0' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 0);
  estimate 'age 5 died 1 hmo 1' exp(intercept * 1 + b_age * 5 + b_died * 1 + b_hmo * 1);
  estimate 'age 7 died 1 hmo 0' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 0);
  estimate 'age 7 died 1 hmo 1' exp(intercept * 1 + b_age * 7 + b_died * 1 + b_hmo * 1);
  estimate 'age 9 died 1 hmo 0' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 0);
  estimate 'age 9 died 1 hmo 1' exp(intercept * 1 + b_age * 9 + b_died * 1 + b_hmo * 1);
run;
```

Now we can use this predicted values for plotting. We need to add actual values of **age** and **hmo** to the dataset for plotting as well.

```

data mylib.addest;
  set mylib.addest;
  input age hmo;
  datalines;
1 0
1 1
3 0
3 1
5 0
5 1
7 0
7 1
9 0
9 1
;
run;

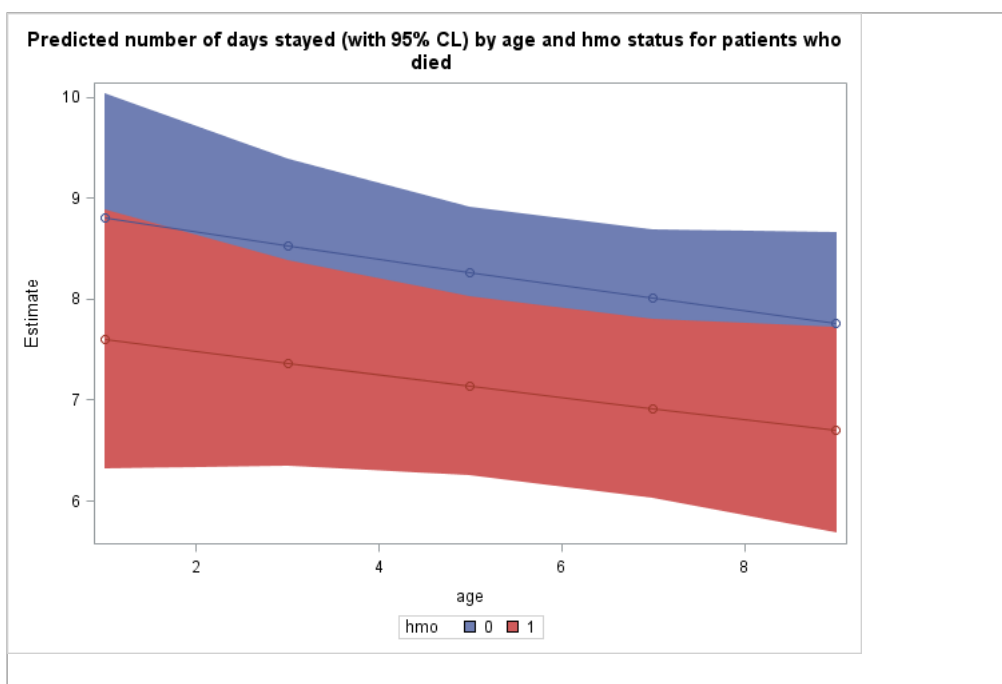
```

Finally, we use **proc sgplot** to plot our predicted number of days stayed as well as 95% confidence interval bands. The predicted values, lines connecting them, and confidence interval bands are all specified separately within the same **proc sgplot**. The **group** option will produce separate points, lines, and bands by the grouping variable.

```

proc sgplot data = mylib.addest;
  title 'Predicted number of days stayed (with 95% CL) by age and hmo status for patients who died';
  band x = age lower = lower upper = upper / group=hmo;
  scatter x= age y = estimate / group = hmo;
  series x = age y = estimate / group = hmo;
run;

```



## Things to consider

- Count data often use exposure variable to indicate the number of times the event could have happened. You can incorporate exposure into your model by including a log-linear term for exposure in the log-likelihood function specification.
- It is not recommended that zero-truncated negative binomial models be applied to small samples. What constitutes a small sample does not seem to be clearly defined in the literature.



## References

- Cameron, A. Colin and Trivedi, P.K. (2009). Microeconometrics using stata. College Station, TX: Stata Press.
- Cameron, A. Colin and Trivedi, P.K. (1998). Regression analysis of count data. Cambridge, UK: Cambridge University Press.
- Hilbe, J. M. (2007). Negative binomial regression. Cambridge, UK: Cambridge University Press.

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