

Example 3: Predicting Count Outcomes using 4 Types of Poisson and Negative Binomial Models (syntax and output available for SAS, STATA, and R electronically)

The data for this example come from a study about the effects of emotion regulation strategy (none=control, cognitive reappraisal, or suppression) in predicting the aggressive verbalizations of persons with or without a history of perpetrating intimate partner violence (IPV). The planned analysis was a two-way between-groups ANOVA for 3 levels of strategy condition by 2 levels of IPV history. Here is the paper published about these data (with similar results, although their models included covariates and so their sample differed slightly):

Maldonado, R. C., DiLillo, D., & Hoffman, L. (2015). [Can college students alter their intimate partner aggression-risk behaviors using emotion regulation strategies? An examination using I3 Theory](#). *Psychology of Violence*, 5(1), 46-55.

This example will examine the results of the same linear predictor using a general linear model (identity link + normal conditional distribution), as well as four types of generalized linear models with log links: Poisson, negative binomial, zero-inflated Poisson, and zero-inflated negative binomial. The probability of being an extra zero is predicted with a logit link in the two zero-inflated variants. Because the relevant STATA options (using GLM to get conditional distribution fit, also using NBREG, ZIP, and ZINB here) do not have denominator degrees of freedom, they were set to "none" in SAS GLIMMIX so that the SAS Wald test results (still labeled as t or F) will match those of STATA (using z or χ^2). In R, I am using the base R function GLM, the `glm.nb` function from package MASS, and the `zeroinfl` function from package pscl (each also using z or χ^2). In SAS (as shown in the online files only), I am still using GLIMMIX (even though these are not mixed-effects models) to get conditional distribution fit, as well as GENMOD for the zero-inflated model variants.

STATA Syntax for Importing and Preparing Data for Analysis:

```
// Defining global variable for file location to be replaced in code below
// \\Client\ precedes path in Virtual Desktop outside H drive;
global filesave "\\Client\C:\Dropbox\23_PSQF6270\PSQF6270_Example3"

// Import Example 3 Excel data
import excel "$filesave\Excel_Example3.xlsx", clear

// STATA code to create indicator-dummy-coded predictor variables
gen NvC=. // Make 2 new empty variables
gen NvS=.
replace NvC=0 if ercond==1 // Replace if ercond=1=None
replace NvS=0 if ercond==1
replace NvC=1 if ercond==2 // Replace if ercond=2=CogR
replace NvS=0 if ercond==2
replace NvC=0 if ercond==3 // Replace if ercond=3=Supp
replace NvS=1 if ercond==3
label variable IPV "IPV: Inter-Partner Violence (0=N,1=Y)"
label variable ercond "ercond: 1=None, 2=CogR, 3=Supp"
label variable aggr "aggr: Aggressive Verbalizations"
label variable NvC "NvC: Condition None=0 vs. CogR=1"
label variable NvS "NvS: Condition None=0 vs. Supp=1"
// Filter to only cases complete on all variables to be used below
egen nummiss = rowmiss(aggr IPV ercond)
drop if nummiss>0
```

R Syntax for Importing and Preparing Data for Analysis (after loading packages *readxl*, *TeachingDemos*, *psych*, *multcomp*, *prediction*, *MASS*, and *pscl*, as shown online):

```
# Define variables for working directory and data name
filesave = "C:\\Dropbox\23_PSQF6270\PSQF6270_Example3/"
filename = "Excel_Example3.xlsx"
setwd(dir=filesave)
```

```
# Import Example 3 Excel data
Example3 = read_excel(paste0(filesave,filename))
# Convert to data frame without labels to use for analysis
Example3 = as.data.frame(Example3)

# R code to create indicator-dummy-coded binary predictors
Example3$NvC=NA; Example3$NvS=NA # Make 2 new empty variables
Example3$NvC[which(Example3$ercond==1)]=0 # Replace if ercond=None=1
Example3$NvS[which(Example3$ercond==1)]=0
Example3$NvC[which(Example3$ercond==2)]=1 # Replace if ercond=CogR=2
Example3$NvS[which(Example3$ercond==2)]=0
Example3$NvC[which(Example3$ercond==3)]=0 # Replace if ercond=Supp=3
Example3$NvS[which(Example3$ercond==3)]=1
# Label variables as comments only (not actually added to data)

# Filter to only cases complete on all variables to be used below
Example3 = Example3[complete.cases(Example3[, c("aggr","IPV","ercond")]),]
```

Syntax and Condensed Output for Data Description:

```
display "STATA Cell Means for Aggressive Verbalizations"
bysort ercond IPV: tabstat aggr, statistics(n max sd mean semean)

display "STATA Histogram for Aggressive Verbalizations"
hist aggr, percent normal discrete width(1) start(0)
graph export "$filesave\STATA Overall Histogram.png", replace

display "STATA Histogram for Aggressive Verbalizations by Cell"
hist aggr, by(IPV ercond) percent normal discrete width(1) start(0)
graph export "$filesave\STATA By Cell Histogram.png", replace

print("R Cell Means for Aggressive Verbalizations Outcome")
describeBy(x=Example3$aggr,list(Example3$IPV,Example3$ercond))

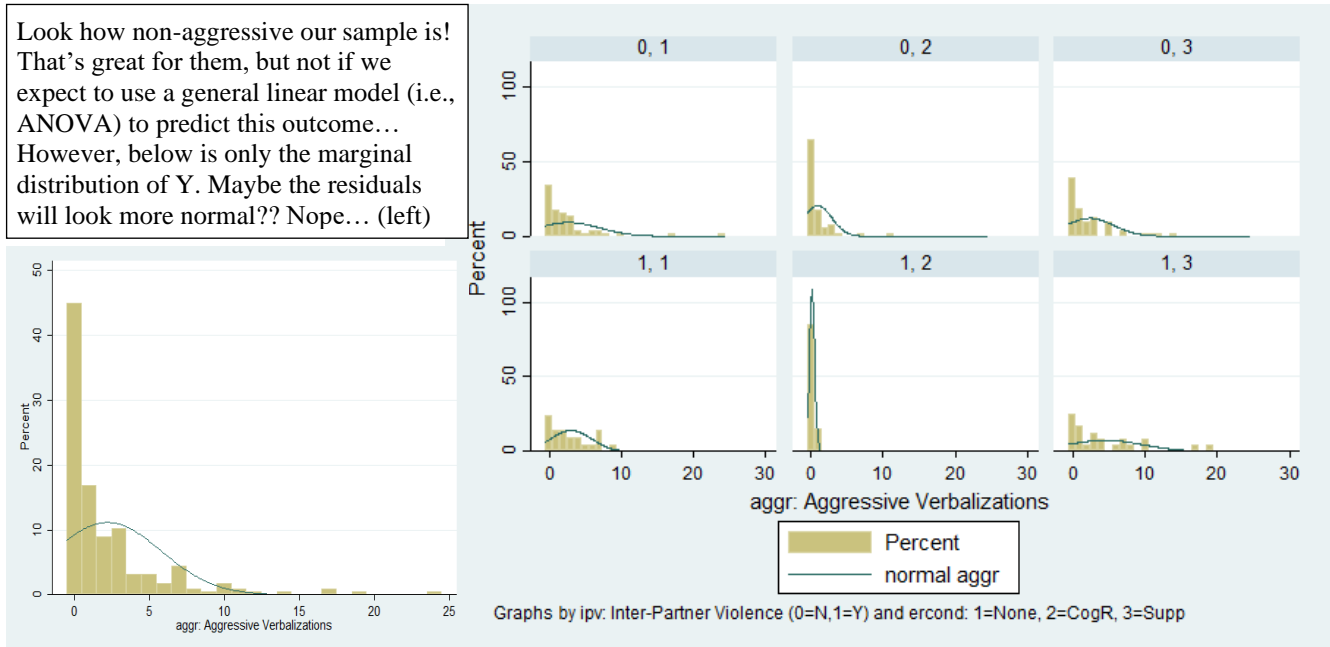
# to save a plot: open a file, create the plot, then close the file
png(file = "R Histogram for Aggressive Verbalizations.png") # open file
hist(x=Example3$aggr, freq=FALSE,
     ylab="Density",xlab="aggr: Aggressive Verbalizations") # axis labels
dev.off() # close file
# I did not figure out how to make a separate histogram for each cell
```

		Exp(Log Link)										
		Raw Data					Normal Dist		Poisson Dist		NegBin Dist	
IPV	Cond	N	Max	SD	Mean	SE	Mean	SE	Mean	SE	Mean	SE
IPV=No	None	53	24	4.32	2.72	0.59	2.72	0.47	2.72	0.23	2.72	0.52
IPV=Yes	None	21	9	2.82	3.05	0.62	3.05	0.74	3.05	0.38	3.05	0.92
IPV=No	CogR	53	11	1.95	0.92	0.27	0.92	0.47	0.92	0.13	0.92	0.21
IPV=Yes	CogR	20	1	0.04	0.15	0.08	0.15	0.76	0.15	0.09	0.15	0.10
IPV=No	Supp	54	14	3.30	2.35	0.45	2.35	0.46	2.35	0.21	2.35	0.46
IPV=Yes	Supp	24	19	5.27	4.46	1.08	4.46	0.70	4.46	0.43	4.46	1.23

Above are the outcome **cell means** for each combination of IPV by emotion regulation condition that our model is trying to capture, along with the per-cell maximum and SD.

What we will see in this example is that the **cell means will stay the same** across models (because we will use the same linear predictor across all models).

What will change are the inferences about their differences (which come from their standard errors, which result from the conditional distribution chosen).



Same Linear Predictor to be used across ALL models:

$$\widehat{Aggr}_i = \beta_0 + \beta_1(IPV_i) + \beta_2(NoneVsCogR_i) + \beta_3(NoneVsSupp_i) + \beta_4(IPV_i)(NoneVsCogR_i) + \beta_5(IPV_i)(NoneVsSupp_i)$$

Model-implied slope of IPV history (no vs. yes) per condition:

$$IPV\ slope = \beta_1 + \beta_4(NoneVsCogR_i) + \beta_5(NoneVsSupp_i)$$

Model-implied slope for condition differences (none, cognitive reappraisal, suppression) per IPV:

$$None\ vs.\ CogR\ slope = \beta_2 + \beta_4(IPV_i)$$

$$None\ vs.\ Supp\ slope = \beta_3 + \beta_5(IPV_i)$$

$$CogR\ vs.\ Supp\ slope = [\beta_3 + \beta_5(IPV_i)] - [\beta_2 + \beta_4(IPV_i)]$$

STATA GLM: Model using an Identity Link and a Normal Conditional Distribution

This model would otherwise be known as ANOVA if it were estimated using ordinary least squares, which is equivalent to residual ML (REML). Although the LL value is using ML estimation, the SEs and Wald tests use the REML estimate of the residual variance instead for comparability with R GLM.

```
display "STATA Link=Identity Dist=Normal Model using glm"
display "SEs use REML=OLS instead for Comparability with R"
glm aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, ml link(identity) family(gaussian) nolog
```

Generalized linear models	No. of obs	=	225
Optimization : ML	Residual df	=	219
Deviance = 2544.228359	Scale parameter	=	11.61748
Pearson = 2544.228359	(1/df) Deviance	=	11.61748
Variance function: V(u) = 1	(1/df) Pearson	=	11.61748 → REML res variance
Link function : g(u) = u	[Gaussian]		
	[Identity]		
Log likelihood = -592.1279267	AIC	=	5.316693 → not usual AIC!
	BIC	=	1358.102 → not usual BIC!

aggr	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]		
IPV	.3306379	.8670718	0.38	0.703	-1.368792	2.030067	Beta1
NvC	-1.792453	.6532266	-2.74	0.006	-3.072753	-.5121523	Beta2
NvS	-.3651293	.6501953	-0.56	0.574	-1.639489	.9092302	Beta3
c.IPV#c.NvC	-1.105166	1.237154	-0.89	0.372	-3.529944	1.319611	Beta4
c.IPV#c.NvS	1.775844	1.196816	1.48	0.138	-.5698726	4.12156	Beta5
_cons	2.716981	.4619009	5.88	0.000	1.811672	3.62229	Beta0

```
display "-2LL= " e(11)*-2 // Print -2LL for model
-2LL= 1184.2559
```

```
// DF=5 Multiv Wald Test of Model
test (c.IPV=0) (c.NvC=0) (c.NvS=0) (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
```

```
chi2( 5) = 27.82
Prob > chi2 = 0.0000
```

```
// DF=2 Multiv Wald Test of Interaction
test (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
```

```
chi2( 2) = 5.70
Prob > chi2 = 0.0579
```

In the intended ANOVA model, this interaction using denominator DF was $F(2, 219) = 2.85, p = .0600$

Btw, $F * \# \text{ slopes} = \chi^2$

```
// Yhat cell means in original count model scale per condition
margins, at(c.IPV=(0(1)1) c.NvC=0 c.NvS=0) predict(xb) // None
margins, at(c.IPV=(0(1)1) c.NvC=1 c.NvS=0) predict(xb) // CogR
margins, at(c.IPV=(0(1)1) c.NvC=0 c.NvS=1) predict(xb) // Supp
```

```
// Simple slopes of IPV per condition
lincom c.IPV*1 + c.IPV#c.NvC*0 + c.IPV#c.NvS*0 // No vs Yes IPV: None
lincom c.IPV*1 + c.IPV#c.NvC*1 + c.IPV#c.NvS*0 // No vs Yes IPV: CogR
lincom c.IPV*1 + c.IPV#c.NvC*0 + c.IPV#c.NvS*1 // No vs Yes IPV: Supp
```

```
// Simple slopes of condition per IPV
lincom c.NvC*1 + c.NvS*0 + c.IPV#c.NvC*0 + c.IPV#c.NvS*0 // None vs CogR: IPV=No
lincom c.NvC*0 + c.NvS*1 + c.IPV#c.NvC*0 + c.IPV#c.NvS*0 // None vs Supp: IPV=No
lincom c.NvC*-1 + c.NvS*1 + c.IPV#c.NvC*0 + c.IPV#c.NvS*0 // CogR vs Supp: IPV=No
lincom c.NvC*1 + c.NvS*0 + c.IPV#c.NvC*1 + c.IPV#c.NvS*0 // None vs CogR: IPV=Yes
lincom c.NvC*0 + c.NvS*1 + c.IPV#c.NvC*0 + c.IPV#c.NvS*1 // None vs Supp: IPV=Yes
lincom c.NvC*-1 + c.NvS*1 + c.IPV#c.NvC*-1 + c.IPV#c.NvS*1 // CogR vs Supp: IPV=Yes
```

```
// Simple slopes for interaction contrasts
lincom c.IPV#c.NvC*1 + c.IPV#c.NvS*0 // No/Yes IPV differ btw None/CogR
lincom c.IPV#c.NvC*0 + c.IPV#c.NvS*1 // No/Yes IPV differ btw None/Supp
lincom c.IPV#c.NvC*-1 + c.IPV#c.NvS*1 // No/Yes IPV differ btw CogR/Supp
```

(Voluminous STATA output for margins and lincom not shown; see R output below)

R GLM: Model using an Identity Link and a Normal Conditional Distribution

```
print("R Link=Identity Dist=Normal Model using glm for ML Estimation")
print("Uses t instead of z for Univariate Wald tests")
ModelNorm = glm(data=Example3, family=gaussian(link="identity"),
  formula=aggr~1+IPV+NvC+NvS +IPV:NvC +IPV:NvS)
print("Print results with -2LL"); summary(ModelNorm)
```

```
Coefficients:
Estimate Std. Error t value Pr(>|t|)
(Intercept) 2.71698 0.46819 5.8032 0.00000002262 Beta0
IPV 0.33064 0.87887 0.3762 0.707126 Beta1
NvC -1.79245 0.66211 -2.7072 0.007321 Beta2
NvS -0.36513 0.65904 -0.5540 0.580123 Beta3
IPV:NvC -1.10517 1.25399 -0.8813 0.379110 Beta4
IPV:NvS 1.77584 1.21310 1.4639 0.144658 Beta5
(Dispersion parameter for gaussian family taken to be 11.617481) → REML residual variance
```

Null deviance: 2867.40 on 224 DDF → **sum of squared Pearson residuals: empty model**
 Residual deviance: 2544.23 on 219 DDF → **sum of squared Pearson residuals: this model**

-2*logLik(ModelNorm)

'log Lik.' 1184.2559 (df=7) → **-2LL**

print("DF=5 Multiv Wald Test of Model with 8 digits")

NormR2 = glht(model=ModelNorm, linfct=c("IPV=0", "NvC=0", "NvS=0", "IPV:NvC=0", "IPV:NvS=0"))

print(summary(NormR2, test=Chisqtest()), digits="8") # Joint chi-square test

	Chisq	DF	Pr(>Chisq)
1	27.817321	5	0.000039517063

print("DF=2 Multiv Wald Test of Interaction with 8 digits")

NormInt = glht(model=ModelNorm, linfct=c("IPV:NvC=0", "IPV:NvS=0"))

print(summary(NormInt, test=Chisqtest()), digits="8") # Joint chi-square test

	Chisq	DF	Pr(>Chisq)
1	5.6992729	2	0.057865353

print("Yhat cell means in original count model scale per condition")

NormPredN = prediction(model=ModelNorm, type="response", at=list(IPV=0:1, NvC=0, NvS=0))

NormPredC = prediction(model=ModelNorm, type="response", at=list(IPV=0:1, NvC=1, NvS=0))

NormPredS = prediction(model=ModelNorm, type="response", at=list(IPV=0:1, NvC=0, NvS=1))

summary(rbind(NormPredN, NormPredC, NormPredS))

at (IPV)	at (NvC)	at (NvS)	Prediction	SE	z	p	lower	upper
0	0	0	2.7170	0.4682	5.8032	0.000000006506	1.799354	3.635
1	0	0	3.0476	0.7438	4.0975	0.000041771484	1.589831	4.505
0	1	0	0.9245	0.4682	1.9747	0.048301670177	0.006901	1.842
1	1	0	0.1500	0.7438	0.2017	0.840173402631	-1.307788	1.608
0	0	1	2.3519	0.4682	5.0233	0.000000507826	1.434225	3.269
1	0	1	4.4583	0.7438	5.9941	0.000000002046	3.000545	5.916

print("Simple slopes: condition per IPV, IPV per condition, interactions")

print("SEs match SAS using REML instead")

NormSlopes = (summary(glht(model=ModelNorm,

linfct=rbind(

"No vs Yes IPV: None" = c(0,1, 0,0, 0,0),

"No vs Yes IPV: CogR" = c(0,1, 0,0, 1,0),

"No vs Yes IPV: Supp" = c(0,1, 0,0, 0,1),

"None vs CogR: IPV=No" = c(0,0, 1,0, 0,0),

"None vs Supp: IPV=No" = c(0,0, 0,1, 0,0),

"CogR vs Supp: IPV=No" = c(0,0, -1,1, 0,0),

"None vs CogR: IPV=Yes" = c(0,0, 1,0, 1,0),

"None vs Supp: IPV=Yes" = c(0,0, 0,1, 0,1),

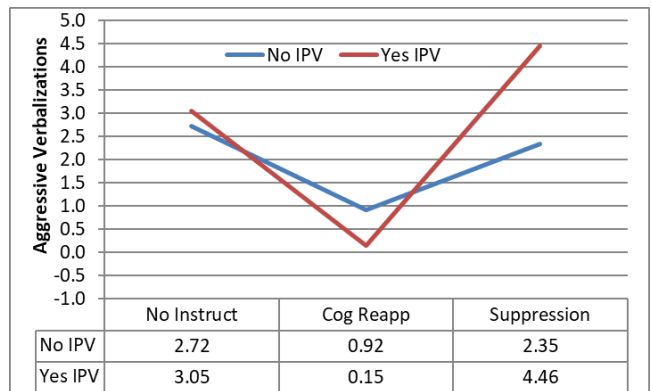
"CogR vs Supp: IPV=Yes" = c(0,0, -1,1, -1,1),

"No/Yes IPV diff btw None/CogR" = c(0,0,0,0, 1,0),

"No/Yes IPV diff btw None/Supp" = c(0,0,0,0, 0,1),

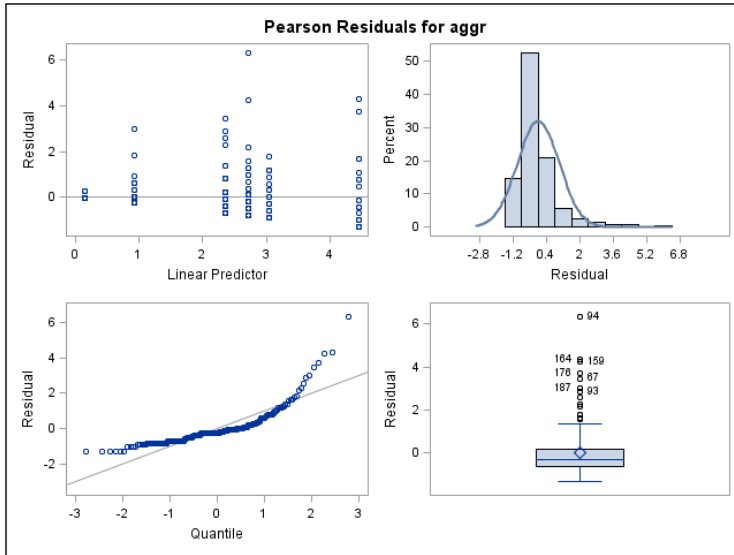
"No/Yes IPV diff btw CogR/Supp" = c(0,0,0,0, -1,1))),

test=adjusted("none"))); NormSlopes



Linear Hypotheses:

	Estimate	Std. Error	z value	Pr(> z)
No vs Yes IPV: None == 0	0.33064	0.87887	0.3762	0.706762
No vs Yes IPV: CogR == 0	-0.77453	0.89447	-0.8659	0.386539
No vs Yes IPV: Supp == 0	2.10648	0.83618	2.5192	0.011763
None vs CogR: IPV=No == 0	-1.79245	0.66211	-2.7072	0.006786
None vs Supp: IPV=No == 0	-0.36513	0.65904	-0.5540	0.579558
CogR vs Supp: IPV=No == 0	1.42732	0.65904	2.1658	0.030330
None vs CogR: IPV=Yes == 0	-2.89762	1.06494	-2.7209	0.006510
None vs Supp: IPV=Yes == 0	1.41071	1.01847	1.3851	0.166011
CogR vs Supp: IPV=Yes == 0	4.30833	1.03196	4.1749	0.00002981
No/Yes IPV differ btw None/CogR == 0	-1.10517	1.25399	-0.8813	0.378144
No/Yes IPV differ btw None/Supp == 0	1.77584	1.21310	1.4639	0.143224
No/Yes IPV differ btw CogR/Supp == 0	2.88101	1.22445	2.3529	0.018627



RESIDUAL Output (from SAS):
 What about that whole non-normal residuals thing? Yep, it's still an issue... in addition, **the variance appears to grow with the mean.**

A data transformation is not going to make this better. What should we do instead? **We get a new model.** Let's try using a log link (to keep the predicted counts positive) and start with a Poisson conditional distribution (in which the conditional variance is supposed to be the same as the conditional mean, which means it is non-constant across the predicted count outcome).

STATA GLM: Model using a Log Link and a Poisson Conditional Distribution

I am using STATA GLM (instead of the direct POISSON function) to obtain the Pearson χ^2 / DF statistic that indicates how well our residuals fit a Poisson conditional distribution.

```
display "STATA Link=Log Dist=Poisson Model using glm"
glm aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, ml link(log) family(poisson) nolog
```

```
Generalized linear models                               No. of obs   =           225
Optimization      : ML                               Residual df   =           219
                                                         Scale parameter =           1
Deviance          = 793.6859856                       (1/df) Deviance = 3.624137
Pearson          = 1028.539634                       (1/df) Pearson  = 4.696528 → too high (1=good)
Variance function: V(u) = u                          [Poisson]
Link function    : g(u) = ln(u)                      [Log]
                                                         AIC          = 5.190062 → Not usual AIC!
Log likelihood   = -577.881941                       BIC          = -392.44 → Not usual BIC!
```

aggr	Coef.	OIM Std. Err.	z	P> z	[95% Conf. Interval]		
IPV	.1148393	.1502313	0.76	0.445	-.1796087	.4092872	Beta1
NvC	-1.077993	.1653862	-6.52	0.000	-1.402144	-.7538419	Beta2
NvS	-.1443183	.1217311	-1.19	0.236	-.3829069	.0942702	Beta3
c.IPV#c.NvC	-1.933488	.6134418	-3.15	0.002	-3.135811	-.7311636	Beta4
c.IPV#c.NvS	.5247327	.1994724	2.63	0.009	.1337739	.9156915	Beta5
_cons	.9995214	.0833333	11.99	0.000	.8361911	1.162852	Beta0

```
display "-2LL= " e(11)*-2 // Print -2LL for model
-2LL= 1155.7639
```

// DF=5 Multiv Wald Test of Model R2

```
test (c.IPV=0) (c.NvC=0) (c.NvS=0) (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
      chi2( 5) = 111.08
      Prob > chi2 = 0.0000
```

From previous normal model: $\chi^2(5) = 27.82, p < .0001$

// DF=2 Multiv Wald Test of Interaction

```
test (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
      chi2( 2) = 20.61
      Prob > chi2 = 0.0000
```

From previous normal model: $\chi^2(2) = 5.70, p = .0579$

// Yhat cell means in log count model scale per condition

```
margins, at(c.ipv=(0(1)1) c.NvC=0 c.NvS=0) predict(xb) // None
margins, at(c.ipv=(0(1)1) c.NvC=1 c.NvS=0) predict(xb) // CogR
margins, at(c.ipv=(0(1)1) c.NvC=0 c.NvS=1) predict(xb) // Supp
```



```
// Y hat cell means in expected count data scale per ercond
margins, at(c.ipv=(0(1)1) c.NvC=0 c.NvS=0) // None
margins, at(c.ipv=(0(1)1) c.NvC=1 c.NvS=0) // CogR
margins, at(c.ipv=(0(1)1) c.NvC=0 c.NvS=1) // Supp
```

Exp(count) margins matches pred means from previous normal model

(All lincom statements are the same as before; see R output below)

```
display "STATA Link=Log Dist=Poisson Model using glm"
display "Request Incidence-Rate Ratios (via eform or irr)"
glm aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, eform ml link(log) family(poisson)
```

I also added the option irr to these lincom statements (online)

R GLM: Model using a Log Link and a Poisson Conditional Distribution

```
# R: save sample size and DDF for conditional distribution fit
DDFn=225 # What SAS GLIMMIX uses by default
DDFk=DDFn-6 # What STATA and SAS GENMOD use (N - # fixed effects)
```

```
print("R Link=Log Dist=Poisson Model using glm")
ModelPoisson = glm(data=Example3, family=poisson(link="log"),
  formula=aggr~1+IPV+NvC+NvS +IPV:NvC +IPV:NvS)
print("Print results with -2LL"); summary(ModelPoisson)
```

```
Coefficients:
              Estimate Std. Error z value      Pr(>|z|)
(Intercept)  0.999521    0.083333  11.9943    < 2.2e-16  Beta0
IPV           0.114839    0.150231   0.7644    0.444619  Beta1
NvC          -1.077993    0.165386  -6.5180  0.00000000007123  Beta2
NvS          -0.144318    0.121731  -1.1856    0.235800  Beta3
IPV:NvC      -1.933488    0.613442  -3.1519    0.001622  Beta4
IPV:NvS       0.524733    0.199472   2.6306    0.008523  Beta5
```

(Dispersion parameter for poisson family taken to be 1)

```
-2*logLik(ModelPoisson)
'log Lik.' 1155.7639 (df=6)
```

```
print("Pearson Chi-Square / DF Index of Fit")
sum(residuals(ModelPoisson, type="pearson")^2)/DDFk # STATA
[1] 4.696528
```

I put these Wald test results inside a print function to ensure enough precision in its reporting for homework.

```
print("DF=5 Multiv Wald Test of Model with 8 digits")
PoissonR2 = glht(model=ModelPoisson,
  linfct=c("IPV=0","NvC=0","NvS=0","IPV:NvC=0","IPV:NvS=0"))
print(summary(PoissonR2, test=Chisqtest()), digits="8") # Joint chi-square test
```

```
Chisq DF Pr(>Chisq)
1 111.07725 5 2.4255681e-22
```

From previous normal model: $\chi^2(5) = 27.82, p < .0001$

```
print("DF=2 Multiv Wald Test of Interaction")
PoissonInt = glht(model=ModelPoisson, linfct=c("IPV:NvC=0","IPV:NvS=0"))
print(summary(PoissonInt, test=Chisqtest()), digits="8") # Joint chi-square test
```

```
Chisq DF Pr(>Chisq)
1 20.614194 2 0.000033395245
```

From previous normal model: $\chi^2(2) = 5.70, p = .0579$

```
print("Yhat cell means in log count model scale per condition")
PoissonLogN = prediction(model=ModelPoisson, type="link", at=list(IPV=0:1,NvC=0,NvS=0))
PoissonLogC = prediction(model=ModelPoisson, type="link", at=list(IPV=0:1,NvC=1,NvS=0))
PoissonLogS = prediction(model=ModelPoisson, type="link", at=list(IPV=0:1,NvC=0,NvS=1))
summary(rbind(PoissonLogN,PoissonLogC,PoissonLogS))
```

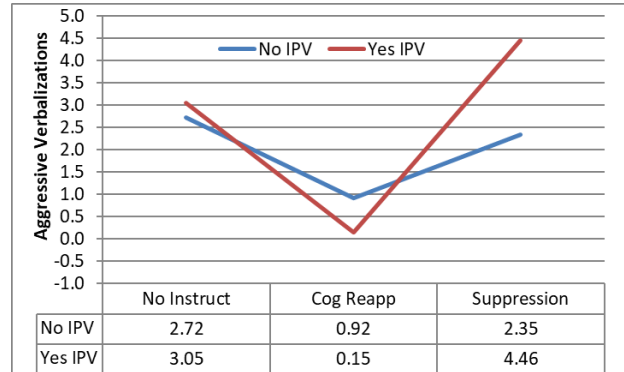
at (IPV)	at (NvC)	at (NvS)	Prediction	SE	z	p	lower	upper
0	0	0	0.99952	0.08333	11.9943	3.808e-33	0.8362	1.16285
1	0	0	1.11436	0.12500	8.9149	4.883e-19	0.8694	1.35936
0	1	0	-0.07847	0.08333	-0.9417	3.464e-01	-0.2418	0.08486
1	1	0	-1.89712	0.12500	-15.1770	5.025e-52	-2.1421	-1.65212
0	0	1	0.85520	0.08333	10.2624	1.040e-24	0.6919	1.01853
1	0	1	1.49478	0.12500	11.9582	5.882e-33	1.2498	1.73977

```
print("Yhat cell means in expected count data scale per condition")
PoissonCountN = prediction(model=ModelPoisson, type="response", at=list(IPV=0:1, NvC=0, NvS=0))
PoissonCountC = prediction(model=ModelPoisson, type="response", at=list(IPV=0:1, NvC=1, NvS=0))
PoissonCountS = prediction(model=ModelPoisson, type="response", at=list(IPV=0:1, NvC=0, NvS=1))
summary(rbind(PoissonCountN, PoissonCountC, PoissonCountS))
```

at (IPV)	at (NvC)	at (NvS)	Prediction	SE	z	p	lower	upper
0	0	0	2.7170	0.2264	12.0000	3.553e-33	2.2732	3.1607
1	0	0	3.0476	0.3810	8.0000	1.244e-15	2.3010	3.7943
0	1	0	0.9245	0.2264	4.0833	4.439e-05	0.4808	1.3683
1	1	0	0.1500	0.3810	0.3938	6.938e-01	-0.5967	0.8967
0	0	1	2.3519	0.2264	10.3873	2.831e-25	1.9081	2.7956
1	0	1	4.4583	0.3810	11.7031	1.228e-31	3.7117	5.2050

Exp(count)
response
matches pred
means from
previous
normal model

```
print("Simple slopes: condition per IPV,  
IPV per condition, interactions")
PoissonSlopes = (summary(gllt(model=ModelPoisson,  
linfct=rbind(  
"No vs Yes IPV: None" = c(0,1, 0,0, 0,0),  
"No vs Yes IPV: CogR" = c(0,1, 0,0, 1,0),  
"No vs Yes IPV: Supp" = c(0,1, 0,0, 0,1),  
"None vs CogR: IPV=No" = c(0,0, 1,0, 0,0),  
"None vs Supp: IPV=No" = c(0,0, 0,1, 0,0),  
"CogR vs Supp: IPV=No" = c(0,0,-1,1, 0,0),  
"None vs CogR: IPV=Yes" = c(0,0, 1,0, 1,0),  
"None vs Supp: IPV=Yes" = c(0,0, 0,1, 0,1),  
"CogR vs Supp: IPV=Yes" = c(0,0,-1,1,-1,1),  
"No/Yes IPV differ btw None/CogR" = c(0,0,0,0, 1,0),  
"No/Yes IPV differ btw None/Supp" = c(0,0,0,0, 0,1),  
"No/Yes IPV differ btw CogR/Supp" = c(0,0,0,0,-1,1))),  
test=adjusted("none"))); PoissonSlopes  
print("IRR effect sizes for simple slopes")  
data.frame(OR=exp(PoissonSlopes$test$coefficients))
```



Comparing results across models: Identify/Normal vs. Log/Poisson:

Model Slope in Log Count	Identity Link, Normal Dist			Log Link, Poisson Dist		
	Est	SE	p-value	Est	SE	p-value
No vs Yes IPV: None	0.33	0.88	.707	0.11	0.15	.445
No vs Yes IPV: CogR	-0.77	0.89	.387	-1.82	0.59	.002
No vs Yes IPV: Supp	2.11	0.84	.012	0.64	0.13	.000
None vs CogR: IPV=No	-1.79	0.66	.007	-1.08	0.17	.000
None vs Supp: IPV=No	-0.37	0.66	.580	-0.14	0.12	.236
CogR vs Supp: IPV=No	1.43	0.66	.030	0.93	0.17	.000
None vs CogR: IPV=Yes	-2.90	1.06	.007	-3.01	0.59	.000
None vs Supp: IPV=Yes	1.41	1.02	.166	0.38	0.16	.016
CogR vs Supp: IPV=Yes	4.31	1.03	.000	3.39	0.59	.000
No/Yes IPV differ by None/CogR	-1.11	1.25	.378	-1.93	0.61	.002
No/Yes IPV differ by None/Supp	1.78	1.21	.143	0.52	0.20	.009
No/Yes IPV differ by CogR/Supp	2.88	1.22	.019	2.46	0.61	.000

The Poisson distribution has only one parameter—the mean, which is supposed to also be the conditional variance. But our Pearson $\chi^2 / DF = 4.697$ result says that the average residual is 4.697 times as large as the Poisson distribution predicts (from conditional SD), indicating that this distribution fit is not good enough yet.

In count data it is often more reasonable to allow the variance to differ from the mean (usually to be greater, known as “over-dispersion”). There are multiple ways to do this; here we will use a negative binomial model to allow the residual variance to change as a quadratic function of the mean (called “NB-2”), which seems to be the most accepted approach.

STATA: Model using a Log Link and a Negative Binomial Conditional Distribution

```
display "STATA Link=Log Dist=Negative Binomial Model using nbreg"
display "nbreg gives LRT for scale factor that distinguishes NB from Poisson"
nbreg aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, nolog
```

```
Negative binomial regression                Number of obs    =          225
                                           LR chi2(5)      =          42.88
Dispersion = mean                          Prob > chi2      =          0.0000
Log likelihood = -409.78835                 Pseudo R2       =          0.0497
```

aggr	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]		
IPV	.1148393	.3591874	0.32	0.749	-.5891552	.8188337	Beta1
NvC	-1.077993	.2962567	-3.64	0.000	-1.658645	-.4973406	Beta2
NvS	-.1443183	.2732663	-0.53	0.597	-.6799104	.3912737	Beta3
c.IPV#c.NvC	-1.933488	.7700748	-2.51	0.012	-3.442807	-.4241687	Beta4
c.IPV#c.NvS	.5247327	.4925367	1.07	0.287	-.4406215	1.490087	Beta5
_cons	.9995214	.1927489	5.19	0.000	.6217405	1.377302	Beta0

/lnalpha	.4706335	.1516165			.1734706	.7677963	log(k)

alpha	1.601008	.2427392			1.189426	2.155012	k dispersion

LR test of alpha=0: $\chi^2(01) = 336.19$ Prob >= $\chi^2 = 0.000 \rightarrow$ NB wins

```
display "STATA Link=Log Dist=Negative Binomial Model using glm"
display "glm gives conditional fit, scale factor estimated by ML"
glm aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, ml link(log) family(nbinomial ml)
```

```
Generalized linear models                No. of obs      =          225
Optimization : ML                        Residual df     =          219
                                           Scale parameter =          1
Deviance = 219.006464                    (1/df) Deviance = 1.00003
Pearson = 244.6296746                    (1/df) Pearson = 1.11703  $\rightarrow$  Close to 1, hooray!
Variance function: V(u) = u+(1.601)u^2  [Neg. Binomial]
Link function : g(u) = ln(u)           [Log]
                                           AIC             = 3.695896  $\rightarrow$  not usual AIC!
Log likelihood = -409.7883514           BIC             = -967.1195  $\rightarrow$  not usual BIC!
```

aggr	Coef.	OIM Std. Err.	z	P> z	[95% Conf. Interval]		
IPV	.1148393	.3591874	0.32	0.749	-.5891552	.8188337	Beta1
NvC	-1.077993	.2962567	-3.64	0.000	-1.658645	-.4973406	Beta2
NvS	-.1443183	.2732663	-0.53	0.597	-.6799104	.3912737	Beta3
c.IPV#c.NvC	-1.933487	.7700748	-2.51	0.012	-3.442806	-.4241685	Beta4
c.IPV#c.NvS	.5247327	.4925367	1.07	0.287	-.4406215	1.490087	Beta5
_cons	.9995214	.1927489	5.19	0.000	.6217405	1.377302	Beta0

Note: Negative binomial parameter estimated via ML and treated as fixed once estimated.

```
display "-2LL=" e(11)*-2 // Print -2LL for model
-2LL = 819.5767
```

(All margins and lincom statements are the same as for Poisson)

```
// DF=5 Multiv Wald Test of Model R2
test (c.IPV=0) (c.NvC=0) (c.NvS=0) (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
      chi2( 5) = 41.22
      Prob > chi2 = 0.0000
```

From normal model: $\chi^2(5) = 27.82, p < .0001$
 From Poisson model: $\chi^2(5) = 111.08, p < .0001$

```
// DF=2 Multiv Wald Test of Interaction
test (c.IPV#c.NvC=0) (c.IPV#c.NvS=0)
      chi2( 2) = 10.47
      Prob > chi2 = 0.0000
```

From normal model: $\chi^2(2) = 5.70, p = .0579$
 From Poisson model: $\chi^2(2) = 20.61, p < .0001$

Poisson model $-2LL = 1155.76$
 $-2\Delta LL(df = 1) = 1155.76 - 819.58$
 $= 336.19, p < .001$
 And the 1.12 fit above means the average residual is close what is predicted by NegBin!

```
// Save predicted counts per real person to dataset
predict predcount
corr predcount aggr // Get corr of pred count with aggr
display "R2=" r(rho)^2 // Print R2 relative to empty model
R2=.11270409

display "STATA Link=Log Dist=Negative Binomial Model"
display "Request Incidence-Rate Ratios (via eform or irr)"
glm aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, eform ml link(log) family(nbinomial ml) nolog
```

I also added the option `irr` to the `lincom` statements (online)

R: Model using a Log Link and a Negative Binomial Conditional Distribution

```
print("R Link=Log Dist=Negative Binomial Model")
print("Using glm.nb add-on to glm from MASS package")
ModelNegBin = glm.nb(data=Example3, link=log,
                      formula=aggr~1+IPV+NvC+NvS +IPV:NvC +IPV:NvS)
print("Print results with -2LL"); summary(ModelNegBin)

Coefficients:
              Estimate Std. Error z value Pr(>|z|)
(Intercept)  0.99952    0.19275  5.1856 0.000002153 Beta0
IPV           0.11484    0.35919  0.3197  0.749181 Beta1
NvC          -1.07799    0.29626 -3.6387  0.000274 Beta2
NvS          -0.14432    0.27327 -0.5281  0.597414 Beta3
IPV:NvC      -1.93349    0.77007 -2.5108  0.012047 Beta4
IPV:NvS      0.52473    0.49254  1.0654  0.286710 Beta5

(Dispersion parameter for Negative Binomial(0.6246) family taken to be 1) → is 1/k instead
      Theta: 0.6246
Std. Err.: 0.0947

-2*logLik(ModelNegBin)
'log Lik.' 819.5767 (df=7)

print("Scale factor in same k metric as SAS and STATA")
ModelNegBin$theta
[1] 1.6010081

print("Pearson Chi-Square / DF Index of Fit")
sum(residuals(ModelNegBin, type="pearson")^2)/DDFk # STATA
[1] 1.1170305

print("Likelihood Ratio Test for Poisson vs NegBin")
DevTest=-2*(logLik(ModelPoisson)-logLik(ModelNegBin))
RegPvalue=pchisq((DevTest), df=1, lower.tail=FALSE); MixPvalue=RegPvalue/2
print("Test Statistic, Regular and Mixture P-values for DF=1")
DevTest; RegPvalue; MixPvalue
'log Lik.' 336.18718 (df=6)
'log Lik.' 4.3176629e-75 (df=6)
'log Lik.' 2.1588315e-75 (df=6)

print("DF=5 Multiv Wald Test of Model with 8 digits")
NegBinR2 = glht(model=ModelNegBin,
                linfct=c("IPV=0", "NvC=0", "NvS=0", "IPV:NvC=0", "IPV:NvS=0"))
print(summary(NegBinR2, test=Chisqtest()), digits="8") # Joint chi-square test

      Chisq DF Pr(>Chisq)
1 41.217936 5 0.000000084780759
From normal model:  $\chi^2(5) = 27.82, p < .0001$ 
From Poisson model:  $\chi^2(5) = 111.08, p < .0001$ 

print("DF=2 Multiv Wald Test of Interaction")
NegBinInt = glht(model=ModelNegBin, linfct=c("IPV:NvC=0", "IPV:NvS=0"))
print(summary(NegBinInt, test=Chisqtest()), digits="8") # Joint chi-square test

      Chisq DF Pr(>Chisq)
1 10.47018 2 0.0053263459
From normal model:  $\chi^2(2) = 5.70, p = .0579$ 
From Poisson model:  $\chi^2(2) = 20.61, p < .0001$ 

# Syntax omitted for cell means, and simple effects (same syntax as for Poisson)
```

```
print("Save predicted counts and correlate with aggr")
Example3$PredCount = predict(ModelNegBin, type="response")
rPred = cor.test(Example3$PredCount, Example3$aggr, method="pearson")
print("R2"); rPred$estimate^2
0.11270409
```

Comparing results across models: Identify/Normal vs. Log/Poisson vs. Log/Negative Binomial:

Model Slope in Log Count	Identity Link, Normal Dist			Log Link, Poisson Dist			Log Link, Neg Bin Dist		
	Est	SE	p-value	Est	SE	p-value	Est	SE	p-value
No vs Yes IPV: None	0.33	0.88	.707	0.11	0.15	.445	0.11	0.36	.749
No vs Yes IPV: CogR	-0.77	0.89	.387	-1.82	0.59	.002	-1.82	0.68	.008
No vs Yes IPV: Supp	2.11	0.84	.012	0.64	0.13	.000	0.64	0.34	.058
None vs CogR: IPV=No	-1.79	0.66	.007	-1.08	0.17	.000	-1.08	0.30	.000
None vs Supp: IPV=No	-0.37	0.66	.580	-0.14	0.12	.236	-0.14	0.27	.597
CogR vs Supp: IPV=No	1.43	0.66	.030	0.93	0.17	.000	0.93	0.30	.002
None vs CogR: IPV=Yes	-2.90	1.06	.007	-3.01	0.59	.000	-3.01	0.71	.000
None vs Supp: IPV=Yes	1.41	1.02	.166	0.38	0.16	.016	0.38	0.41	.353
CogR vs Supp: IPV=Yes	4.31	1.03	.000	3.39	0.59	.000	3.39	0.70	.000
No/Yes IPV differ by None/CogR	-1.11	1.25	.378	-1.93	0.61	.002	-1.93	0.77	.012
No/Yes IPV differ by None/Supp	1.78	1.21	.143	0.52	0.20	.009	0.52	0.49	.287
No/Yes IPV differ by CogR/Supp	2.88	1.22	.019	2.46	0.61	.000	2.46	0.76	.001

Given the large amount of zero values, we should examine whether we have adequately addressed them—let’s compare our currently winning negative binomial model with models using Zero-Inflated Poisson and Zero-Inflated Negative Binomial conditional distributions. These add a separate “zero-inflation” submodel that predicts the logit of being an “extra” zero relative to what is expected given a Poisson or Negative Binomial conditional distribution. Here, we are fitting empty zero-inflation models that contain only an intercept for the probability of being an extra 0—if it’s small enough, we don’t need a zero-inflation submodel at all!

STATA ZIP: Model using a Log Link and a Zero-Inflated Poisson Conditional Distribution

```
display "STATA Zero-Inflated Poisson Model"
display "Only intercept in zero-inflation model (predict logit of extra zero)"
zip aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, inflate(_cons)
```

```
Zero-inflated Poisson regression              Number of obs   =          225
                                                Nonzero obs     =          124
                                                Zero obs        =          101
Inflation model = logit                     LR chi2(5)      =          64.17
Log likelihood = -489.5851                   Prob > chi2     =          0.0000
```

	aggr	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
aggr							
	IPV	-.0232234	.1536578	-0.15	0.880	-.3243872 .2779404	
	NvC	-.7697161	.2118299	-3.63	0.000	-1.184895 -.3545372	
	NvS	-.0763171	.1264847	-0.60	0.546	-.3242226 .1715883	
	c.IPV#c.NvC	-2.058436	.6469452	-3.18	0.001	-3.326425 -.7904467	
	c.IPV#c.NvS	.4808622	.2041397	2.36	0.018	.0807558 .8809686	
	_cons	1.399623	.0858963	16.29	0.000	1.231269 1.567977	
inflate							
	_cons	-.5453023	.1649153	-3.31	0.001	-.8685302 -.2220743	→ logit extra 0 ignore p-value

```
display "-2LL=" e(11)*-2 // Print -2LL for model
-2LL= 979.17018
```

I used "coeflegend" to get the name of zero-inflation submodel intercept

```
nlcom 1/(1+exp(-1*_b[inflate:_cons])) // Probability of extra 0
```

	aggr	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]
	_nl_1	.366955	.0383097	9.58	0.000	.2918695 .4420405

37% extra 0 values

R ZIP: Model using a Log Link and a Zero-Inflated Poisson Conditional Distribution

```
print("R Link=Log Dist=Zero-Inflated Poisson Model using pscl package")
print("Only intercept in zero-inflation model, predict logit of extra zero")
ModelZIP = zeroinfl(data=Example3, dist="poisson", link="logit",
                    formula=aggr~1+IPV+NvC+NvS +IPV:NvC +IPV:NvS | 1)
print("Print results with -2LL"); summary(ModelZIP)
```

Count model coefficients (poisson with log link):

	Estimate	Std. Error	z value	Pr(> z)
(Intercept)	1.399623	0.085896	16.2943	< 2.2e-16
IPV	-0.023224	0.153658	-0.1511	0.8798668
NvC	-0.769716	0.211830	-3.6337	0.0002794
NvS	-0.076317	0.126485	-0.6034	0.5462621
IPV:NvC	-2.058437	0.646945	-3.1818	0.0014637
IPV:NvS	0.480863	0.204140	2.3556	0.0184949

Zero-inflation model coefficients (binomial with logit link):

	Estimate	Std. Error	z value	Pr(> z)
(Intercept)	-0.54530	0.16492	-3.3066	0.0009445

→ logit extra 0 (ignore p-value)

```
-2*logLik(ModelZIP);
'log Lik.' 979.17018 (df=7)
```

```
print("Pearson Chi-Square / DF Fit")
sum(residuals(ModelZIP, type="pearson")^2)/(DDFk-1) # Not given in STATA
[1] 2.0623518
```

```
print("Get probability of being an extra 0")
ZIPprob=1/(1+exp(-1*ModelZIP$coefficients$zero)); ZIPprob
(Intercept)
0.36695499
```

R ZINB: Model using a Log Link and a Zero-Inflated Negative Binomial Conditional Distribution (because STATA blew up!)

```
print("R Link=Log Dist=Zero-Inflated Negative Binomial Model using pscl package")
print("Only intercept in zero-inflation model, predict logit of extra zero")
ModelZINB = zeroinfl(data=Example3, dist="negbin", link="logit",
                    formula=aggr~1+IPV+NvC+NvS +IPV:NvC +IPV:NvS | 1)
print("Print results with -2LL"); summary(ModelZINB)
```

Count model coefficients (negbin with log link):

	Estimate	Std. Error	z value	Pr(> z)
(Intercept)	0.99953	0.19278	5.1849	0.0000002162
IPV	0.11485	0.35919	0.3198	0.7491508
NvC	-1.07798	0.29626	-3.6387	0.0002741
NvS	-0.14433	0.27327	-0.5282	0.5973929
IPV:NvC	-1.93361	0.77009	-2.5109	0.0120430
IPV:NvS	0.52476	0.49254	1.0654	0.2866862
Log(theta)	-0.47061	0.15180	-3.1003	0.0019334

Zero-inflation model coefficients (binomial with logit link):

	Estimate	Std. Error	z value	Pr(> z)
(Intercept)	-11.108	230.777	-0.0481	0.9616

Theta = 0.62462 → 1/k dispersion

```
-2*logLik(ModelZINB)
'log Lik.' 819.57674 (df=8) → Nearly identical to original negative binomial model
```

```
print("Pearson Chi-Square / DF Index of Fit")
sum(residuals(ModelZINB, type="pearson")^2)/(DFk-1)
[1] 1.1170425 → Very similar to original negative binomial

print("Scale factor in same k metric as SAS and STATA")
1/ModelZINB$theta
[1] 1.6009695 → Very similar to original negative binomial

print("Get probability of being an extra 0")
ZINBprob=1/(1+exp(-1*ModelZINB$coefficients$zero)); ZINBprob
0.000014994787
```

The logit of being an “extra 0” = -11.108 (with a crazy SE)! This is probability =.000014994787 of being an “extra” 0. So there are no extra 0 values in this distribution not already predicted by the negative binomial.

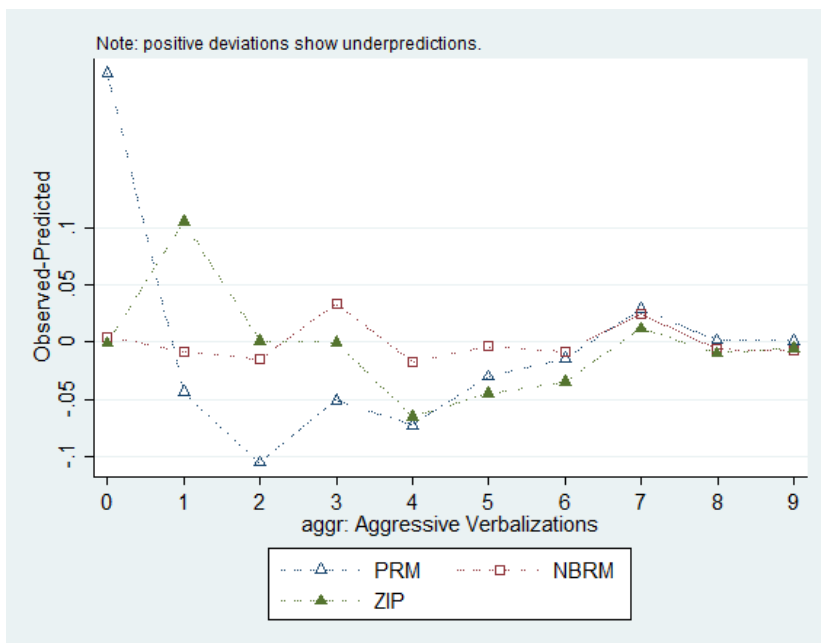
STATA user routine “countfit” can be used to compare count model conditional distributions:

```
// Run search below, then install from window that pops up
search countfit

display "STATA Countfit to Compare Fit of Alternative Count Model Distributions"
display "prm=Poisson, nbreg=Negative Binomial, zip=Zero-Inflated Poisson"
display "Results suggest NegBin fits best"
countfit aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, prm nbreg zip replace
graph export "$filesave\STATA Predicted Counts from Countfit.png", replace
```

Tests and Fit Statistics

PRM (Poisson)	BIC=	1188.260	AIC=	1167.764	Prefer	Over	Evidence
vs NBRM	BIC=	857.489	dif=	330.771	NBRM	PRM	Very strong
	AIC=	833.577	dif=	334.187	NBRM	PRM	
	LRX2=	336.187	prob=	0.000	NBRM	PRM	p=0.000
vs ZIP	BIC=	1031.774	dif=	156.487	ZIP	PRM	Very strong
	AIC=	990.780	dif=	176.983	ZIP	PRM	
	Vuong=	.	prob=	.	ZIP	PRM	p=.
NBRM (NegBin)	BIC=	857.489	AIC=	833.577	Prefer	Over	Evidence
vs ZIP	BIC=	1031.774	dif=	-174.284	NBRM	ZIP	Very strong
	AIC=	990.780	dif=	-157.204	NBRM	ZIP	
ZIP	BIC=	1031.774	AIC=	990.780	Prefer	Over	Evidence



Left: This plot from STATA countfit shows the match between the model-predicted counts and the actual counts for three models: PRM=Poisson, NBRM=Negative Binomial, and ZIP=zero-inflated Poisson. I did not add ZINB given that it blew up.

The NBRM has the closest match to the observed counts (smallest deviations), consistent with our Pearson χ^2 / DF results.

One last model—does the log of the negative binomial scale parameter need to differ by the same linear predictor used for the log of the expected count? Let's see—this is called a Heterogeneous Negative Binomial model. It is not directly available in SAS, but it can be programmed in NLMIXED, as I found [here](#) by Robin High. I searched but did not find it in R (although I'm sure it's in there somewhere).

```
display "STATA Heterogeneous Negative Binomial Model"
display "lnalpha gives linear model to predict log of scale factor"
gnbreg aggr c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS, ///
      lnalpha(c.IPV c.NvC c.NvS c.IPV#c.NvC c.IPV#c.NvS)
display "-2LL=" e(11)*-2 // Print -2LL for model
```

STATA Output for Heterogeneous Neg Bin:

```
Generalized negative binomial regression      Number of obs      =          225
                                             LR chi2(5)         =          22.98
                                             Prob > chi2        =          0.0003
Log likelihood = -406.66217                 Pseudo R2          =          0.0275
-----+-----
```

	aggr	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]
aggr						
	IPV	.1148393	.295082	0.39	0.697	-.4635109 .6931894
	NvC	-1.077993	.3395887	-3.17	0.002	-1.743575 -.4124115
	NvS	-.1443183	.2765458	-0.52	0.602	-.6863381 .3977014
	c.IPV#c.NvC	-1.933485	.7057295	-2.74	0.006	-3.316689 -.5502803
	c.IPV#c.NvS	.5247327	.4394642	1.19	0.232	-.3366012 1.386067
	_cons	.9995214	.193111	5.18	0.000	.6210309 1.378012
lnalpha						
	IPV	-.8075255	.5892574	-1.37	0.171	-1.962449 .3473977
	NvC	.6411665	.4645302	1.38	0.168	-.2692959 1.551629
	NvS	.0499735	.3927454	0.13	0.899	-.7197934 .8197403
	c.IPV#c.NvC	-12.20546	23.22818	-0.53	0.599	-57.73185 33.32094 → Uh oh..
	c.IPV#c.NvS	.6048024	.7515379	0.80	0.421	-.8681849 2.07779
	_cons	.4752475	.2696956	1.76	0.078	-.0533463 1.003841

```
-----+-----
```

After re-estimating the model removing the interaction terms in predicting the scale factor, none of the effects predicting different scale factors by condition or IPV are significant and the information criteria are higher (worse). This indicates the original negative binomial with a constant scale factor is likely to be sufficient.

One other idea—given the lack of quantitative predictors to be given linear slopes (that could create predicted counts below zero without a link function) we could also estimate a general linear model in which the residual variance is allowed to differ across the six conditions. This would also address the problem of non-constant residual variance by tying it to the linear predictor rather than the predicted mean per se. This type of heterogeneous variance model could be done in SAS MIXED or GLIMMIX, as well as STATA MIXED and R LME. But given our focus on generalized linear models, I will leave that idea for another example...

Sample results section [notes what else should be included]:

We examined the extent to which how the count of aggressive verbalizations in the experimental condition differed across three strategy conditions (none, cognitive reappraisal, or suppression) as a function of whether participants had a history of intimate partner violence (IPV; no, yes). We estimated generalized linear models using maximum likelihood without denominator degrees of freedom. Effect sizes are provided using incident-rate ratios (IRR), which are exponentiated slope coefficients interpreted similarly to odds ratios. IRR values between 0 and 1 indicate negative effects, an IRR value of 1 indicates no effect, and IRR values > 1 indicate positive effects.

Before examining the results, we first examined the fit of the conditional distribution to the model residuals. As expected given the highly skewed observed count distribution, a model specifying an identity link function and normal residuals (i.e., a standard analysis of variance) resulted in confidence intervals for the cell means that included negative (impossible) count values. An alternative model specifying a log link function and Poisson conditional distribution (in which the conditional mean and variance are the same) did not appear to fit the observed distribution (Pearson $\chi^2/DF = 4.70$). This is because the conditional variance significantly exceeded the conditional mean, as indicated by a significant likelihood ratio test for a model specifying a negative binomial distribution instead (i.e., that included a scale factor to allow over-dispersion as a quadratic function of the mean, NB2), $-2\Delta LL(1) = 336.19, p < .0001$. Adding a zero-inflation parameter did not improve model fit, $-2\Delta LL(1) = 0$, indicating that the observed 0 values were adequately captured within the negative binomial distribution (Pearson $\chi^2/DF = 1.12$). Finally, we examined the potential for group differences in the log of the dispersion scale factor using the same linear predictor as for the log count, but no main effects or interactions were significant, suggesting the original negative binomial with a single scale factor is likely to be sufficient.

The overall model explained a significant amount of variance in aggressive verbalizations, $\chi^2(5) = 41.22, p < .0001$. The correlation between the predicted and actual counts was $.336 (R^2 = .113)$; dispersion parameter = 1.601. As expected, there was a significant interaction between strategy condition and history of IPV, $\chi^2(2) = 10.47, p = .005$. [Figure 1 depicts the adjusted cell means for the log counts in panel A, and the expected counts in panel B. Table 1 provides simple slopes and slope differences within the interaction].

Let us first consider the pattern of the interaction with IPV as a moderator of the effect of strategy condition. The number of aggressive verbalizations was significantly lower when using a cognitive reappraisal strategy than when using no strategy (control) or a suppression strategy, and this was true for persons with or without a history of IPV. However, these benefits of a cognitive reappraisal strategy (relative to control or suppression) were significantly stronger in persons with a history of IPV than persons without a history of IPV.

Let us next consider the pattern of the interaction with strategy condition as a moderator of the effect of IPV. There were no significant IPV group differences when using no strategy (although aggressive verbalizations were marginally higher in persons with a history of IPV than without when using a suppression strategy). Surprisingly, the number of aggressive verbalizations when using a cognitive reappraisal strategy was significantly lower in persons with a history of IPV than without a history of IPV. This IPV group difference was significantly larger for participants using a cognitive reappraisal strategy than those using no strategy, and the IPV effect differed significantly between the cognitive reappraisal and suppression strategy conditions.