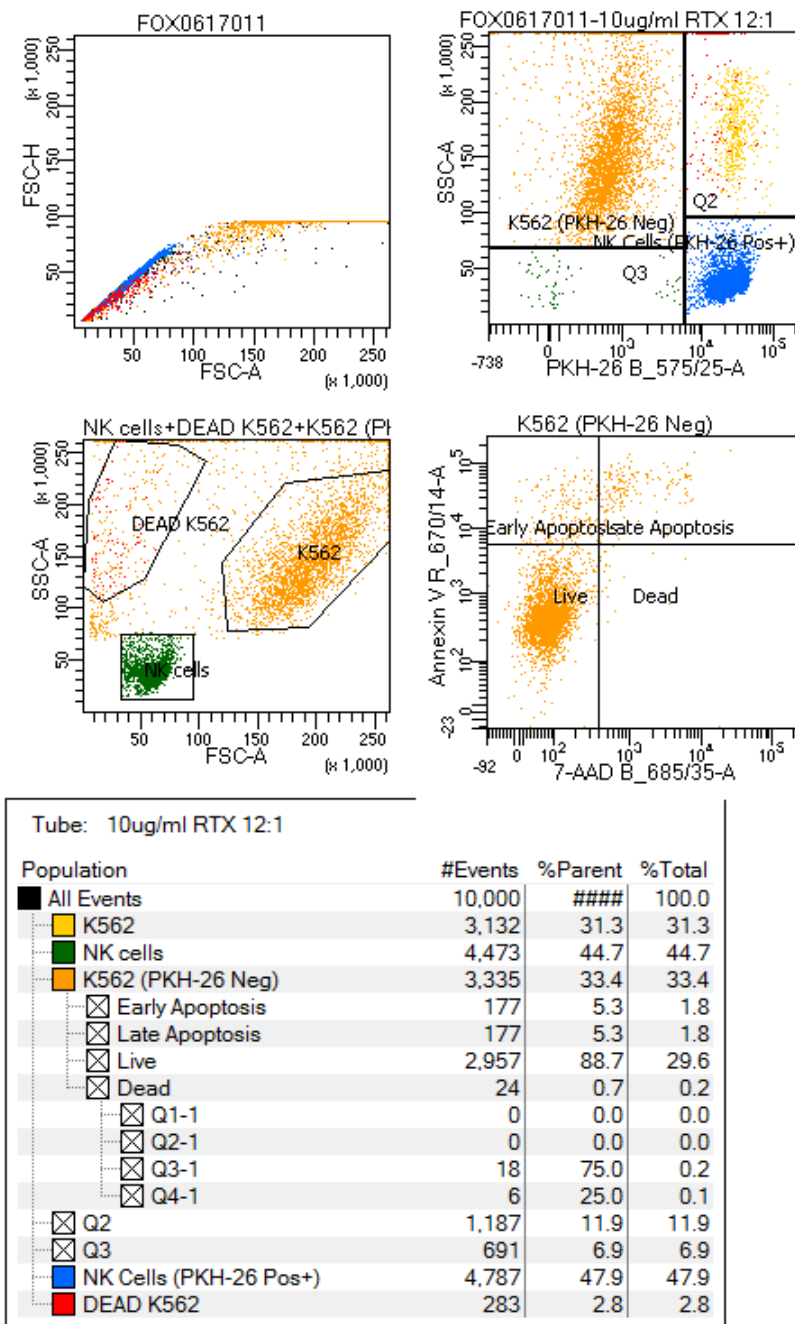
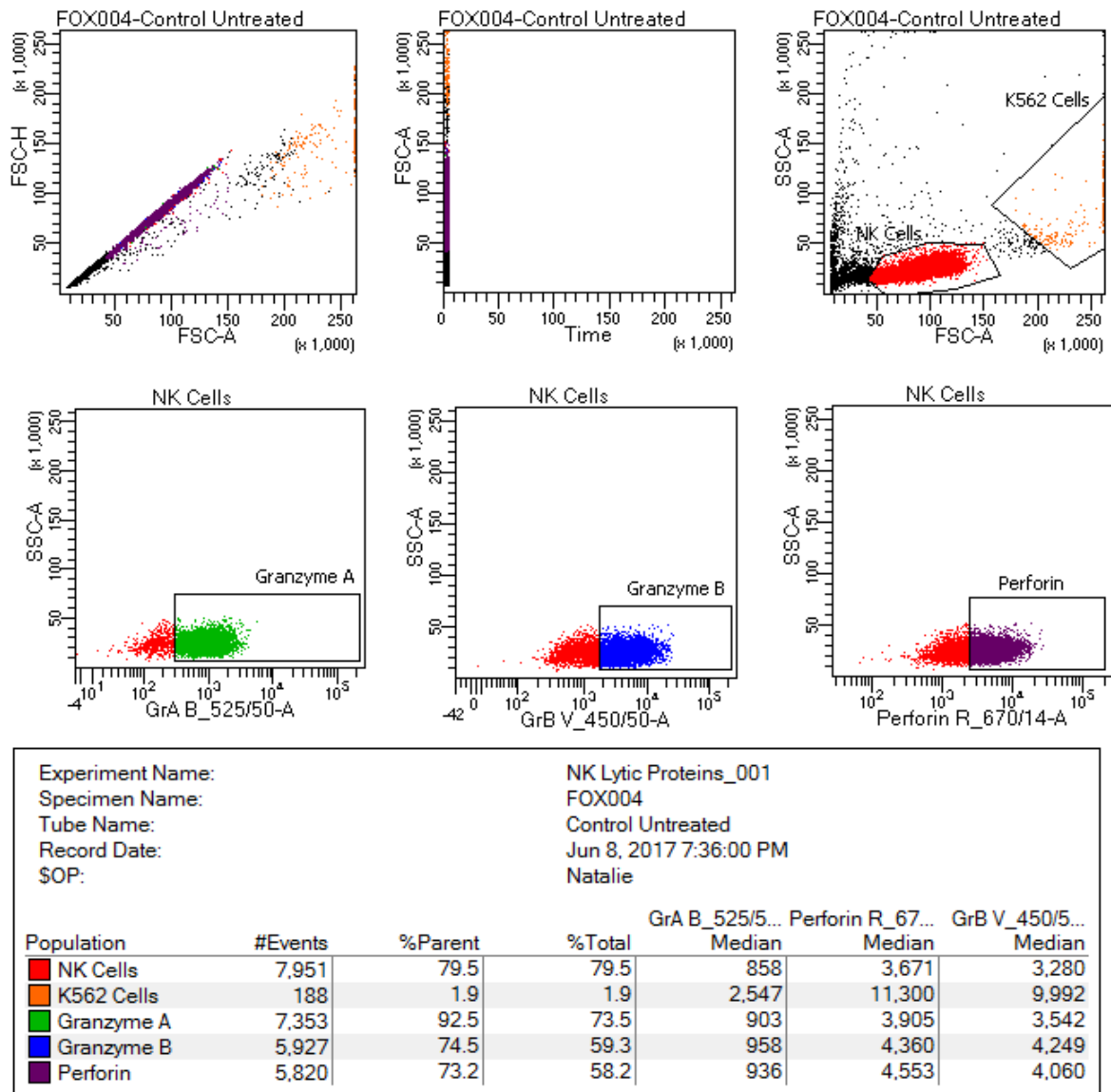


Supplementary Figure 1:



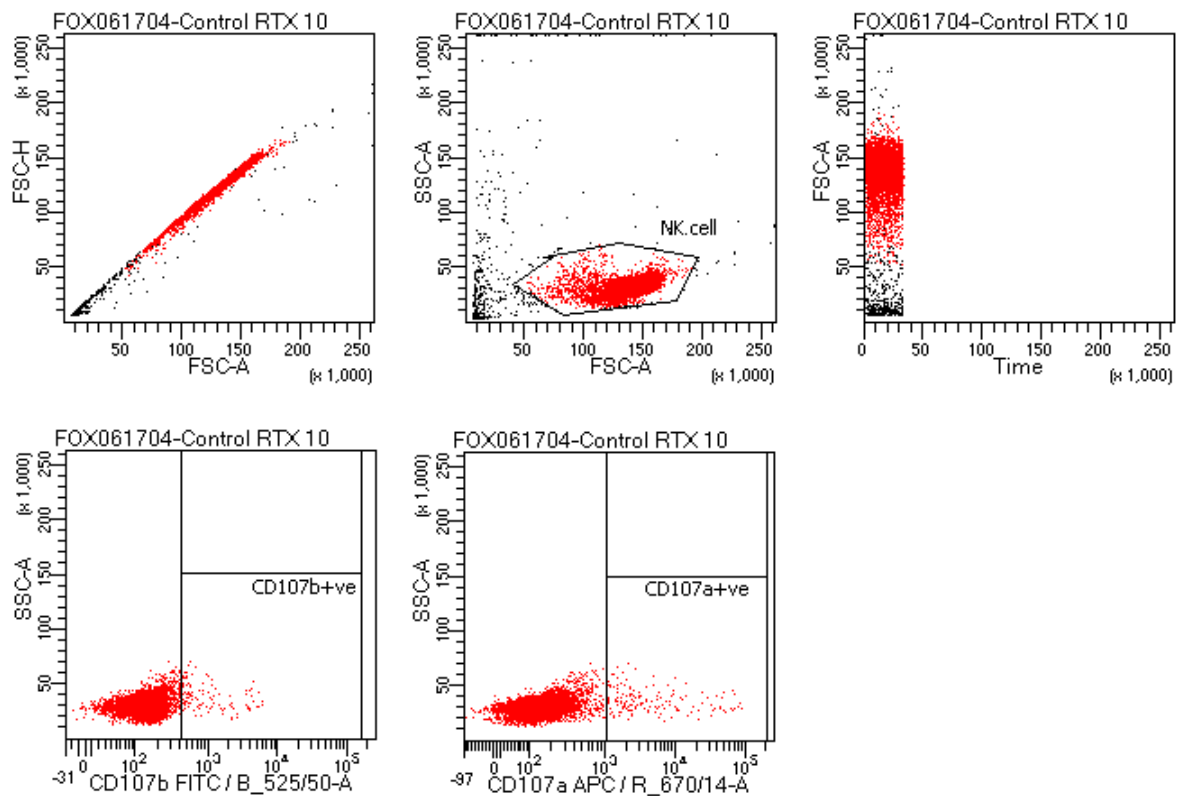
Supplementary Figure 1: Representative flow cytometric plot of NK cell cytotoxic activity. PKH-26 was used to gate NK cells. PKH-26 negative cells were identified as K562 cells and used to determine apoptosis. Annexin V was used to determine events undergoing apoptosis and 7-AAD was used to determine cells undergoing late apoptosis and cells that were dead.

Supplementary Figure 2:



Supplementary Figure 2: Representative flow cytometric plot of intracellular staining analysis of lytic proteins. NK cells were isolated and were used if isolation purity was $\geq 95\%$. NK cells were selectively gated on using flow cytometry and isotype controls were used to determine the positive population for Granzyme A and Granzyme B expression.

Supplementary Figure 3:



Experiment Name:	NK Cell Degranulation_001				
Specimen Name:	FOX061704				
Tube Name:	Control RTX 10				
Record Date:	Jun 8, 2017 6:10:51 PM				
SOP:	Natalie				
Population	#Events	%Parent	%Total	CD107b FITC / B... Median	CD107a APC / R... Median
All Events	10,130	####	100.0	131	169
NK cell	9,648	95.2	95.2	131	167
CD107b+ve	112	1.2	1.1	733	5,064
CD107a+ve	148	1.5	1.5	511	2,925

Tube: Control RTX 10			
Population	#Events	%Parent	%Total
All Events	10,130	####	100.0
NK cell	9,648	95.2	95.2
CD107b+ve	112	1.2	1.1
CD107a+ve	148	1.5	1.5

Supplementary Figure 3: Representative flow cytometric plots for degranulation markers. NK cells were isolated and were used if isolation purity was $\geq 95\%$. NK cells were gated and selected using flow cytometry to determine CD107a and CD107b expression. Isotype controls were used to determine the positive population.