

Gate that cell!

Requirement for analyzing Flow Cytometry Data Reflection on H62 document

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Outline

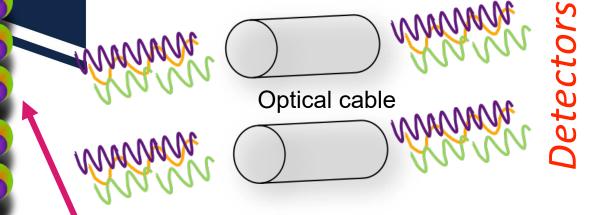
- Overview of how data generated
- Poisson distribution for flow
- Variability –accounting for planning
- Gating strategies and their usefulness

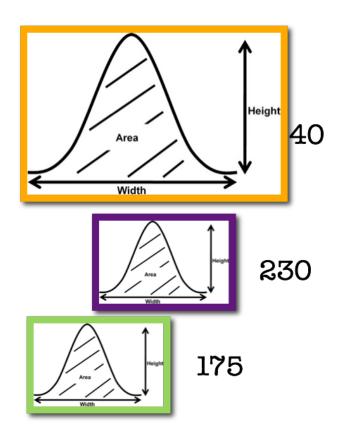


How flow works...

Brief overview of Flow Process

Lasers



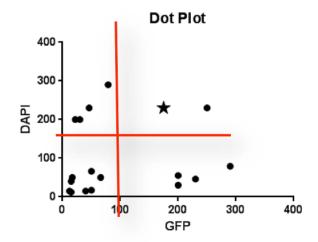


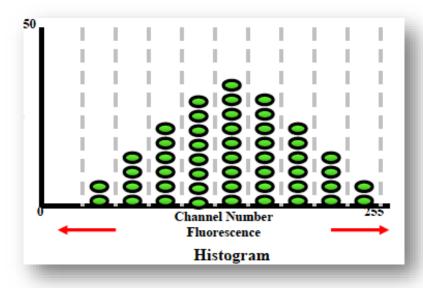


Brief overview of Flow Process



	Violet DAPI	Green GFP	Orange Propidi um Iodide	
Cell 1	200	30	30	
Cell 2	230	175	40	
Cell 3	250	50	50	
Cell 4	180	190	60	
Cell 5	200	46	70	
Cell 5	200	46	70	

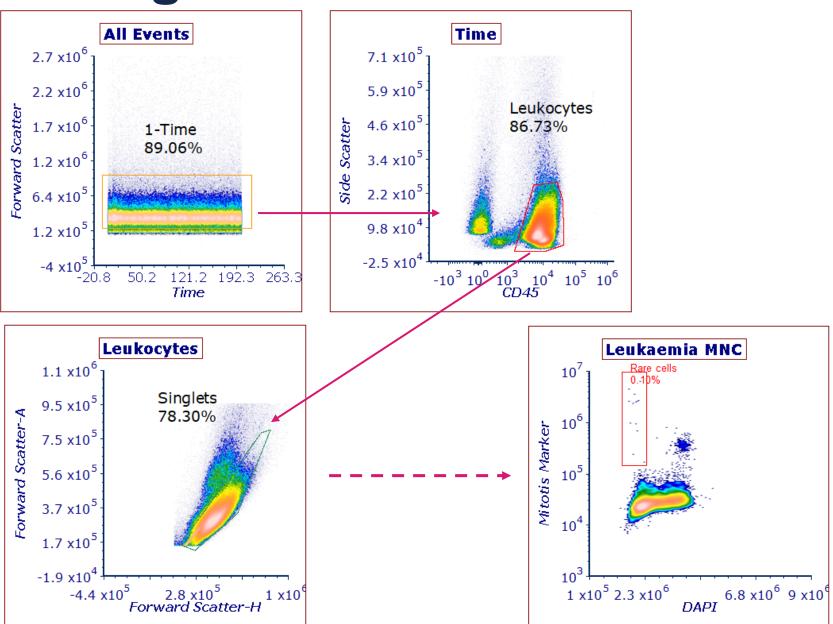






Gating is used to derive data, but it helps us clean up data.

Gating





Variability enemy of statistics

H62 and FDA references to rare cells

"To define a cell population [to gate it] and allow reproducible enumeration it has been recommended that 20-50 clustered events be present..." H62

Desired CV	1%	2.5%	5%	10%	20%	
Required Events for the Population of Interest	10 000	1600	400	100	25	
Population of Interest Frequency, %	Total Number of Acquired Events					
10	100 000	16 000	4000	1000	250	
1	1 000 000	160 000	40 000	10 000	2500	
0.1	10 000 000	1 600 000	400 000	100 000	25 000	
0.01	100 000 000	16 000 000	4 000 000	1 000 000	250 000	
0.001	1 000 000 000	160 000 000	40 000 000	10 000 000	2 500 000	

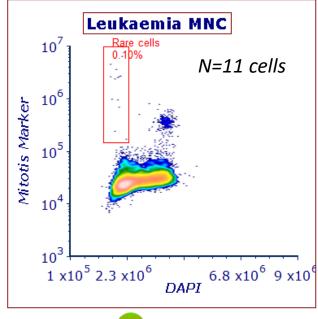


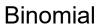
Poisson distribution How many cells we need?!



$$CV$$
, $\% = \frac{\sqrt{N}}{N}$

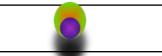
$$CV$$
, % = $\frac{\sqrt{N}}{N}$
 $CV = \frac{\sqrt{11}}{11} = 30.2\%$





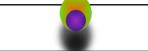


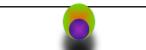




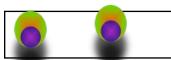




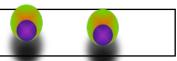




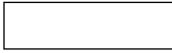
Poisson

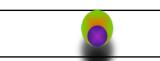














Variability enemy of statistics



- Population separation
- Blood Lysis vs. PBMC
 - Debris and Viability
- Analysts
- Instruments
- Temperature
- Centrifugation, xg force
- Type of assay
 - Intracellular
 - PhosphoFlow
- Logistics
- Sample type
- Analysis of data



Phosphorylated Biomarker MD

- Very low frequency
- Expressed in cell lines
- Mitosis biomarker
- Predictive in number of tumours as progression marker
- Matrix is rare
- Not every patient expresses biomarker of interest
- This was whole blood assay



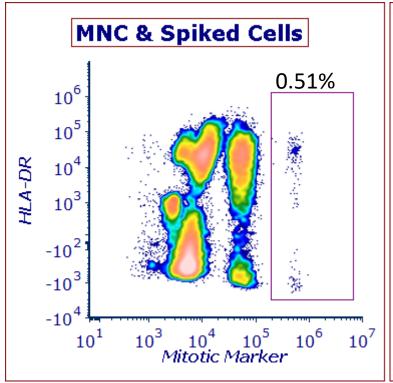
H62 Specificity and Gating

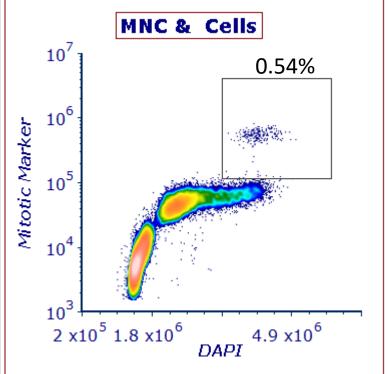
"...Once gating strategy is determined the results should be verified against reference or comparative method....."

"...final document should provide how gating was designed to ensure specificity"



Spiked healthy individuals



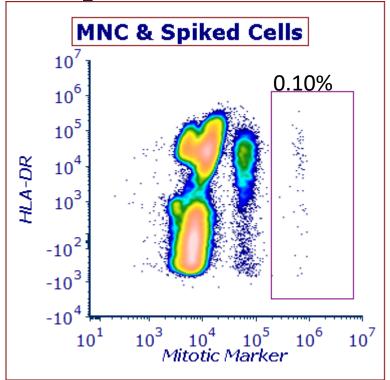


0.177% of total CV=4.04

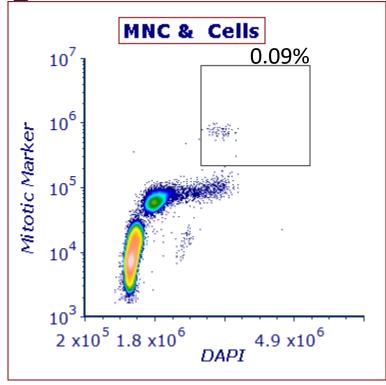
0.18% of total CV=2.71



Spiked healthy individuals



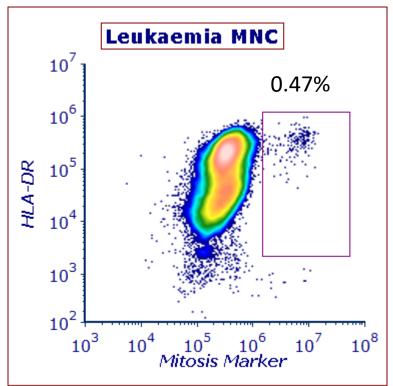
0.03% of total CV=15.71% of Blast gate

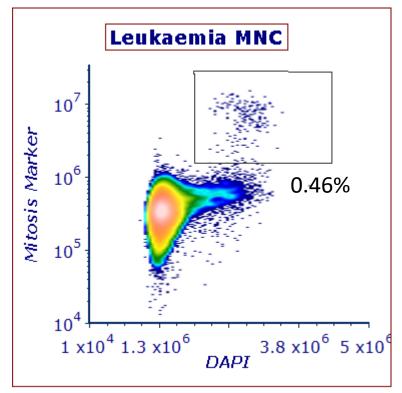


0.03% of Blast gate CV=8.32%



Disease Matrix

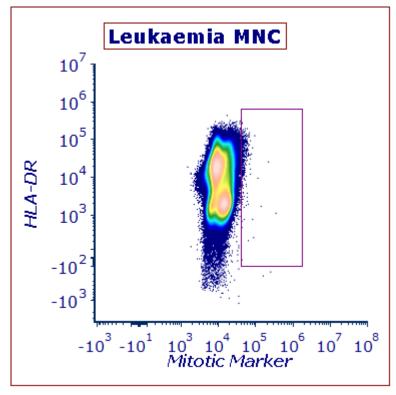




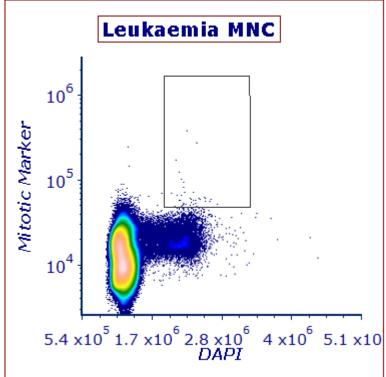
High Concordance!!!



Disease Matrix



0.1% of blasts CV=0%



0.01% of blasts CV=0%

Conclusion



- Plan should contain descriptive element of gating: Clustered events, Pattern
- Gating strategy comes first!
- Effective gating strategy for biomarker and disease
- Inclusion of further means to resolve population may improve specificity and sensitivity



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Additional Slide for question

Q: Were there any cases with Mitotic Marker expressed aberrantly?

