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# LFTDI

LABORATORY FOR  
FORENSIC  
TECHNOLOGY  
DEVELOPMENT &  
INTEGRATION

Catherine M. Grgicak

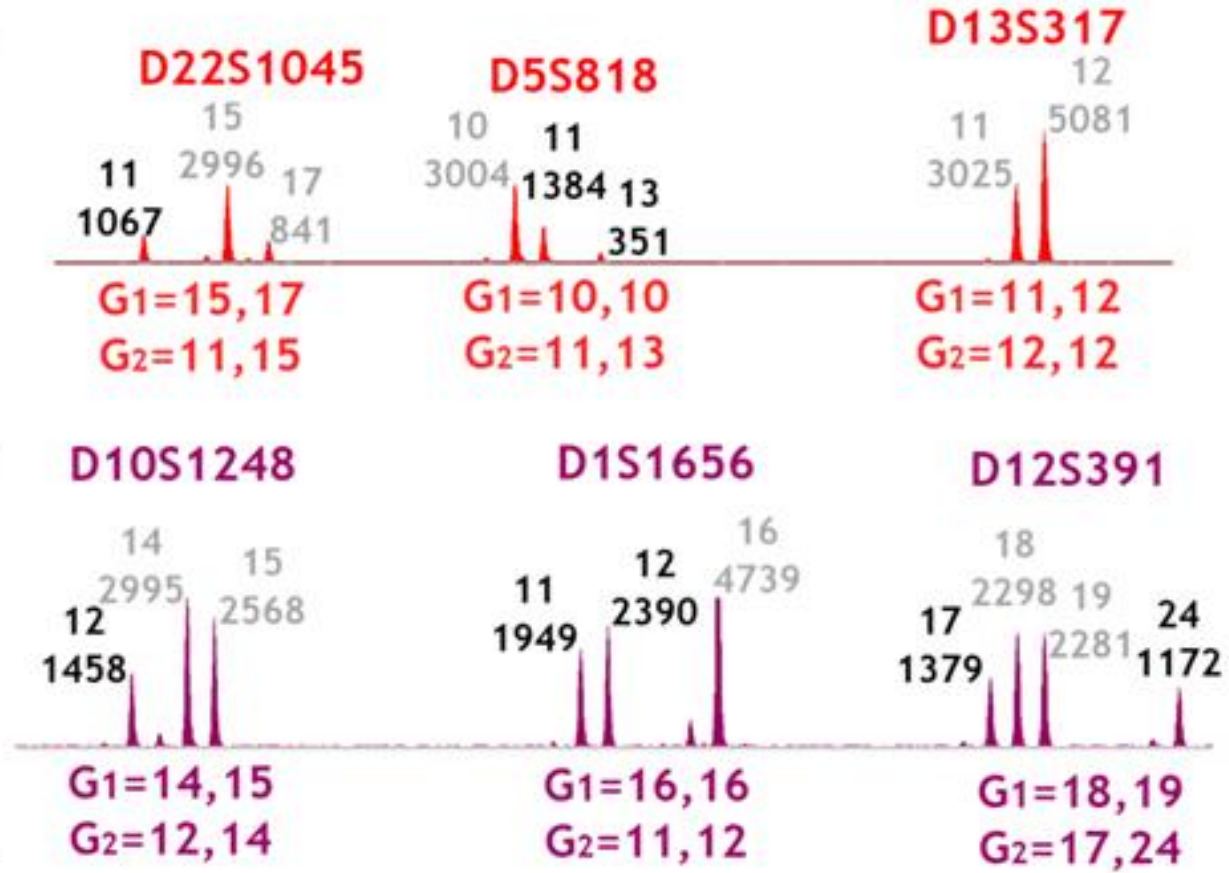
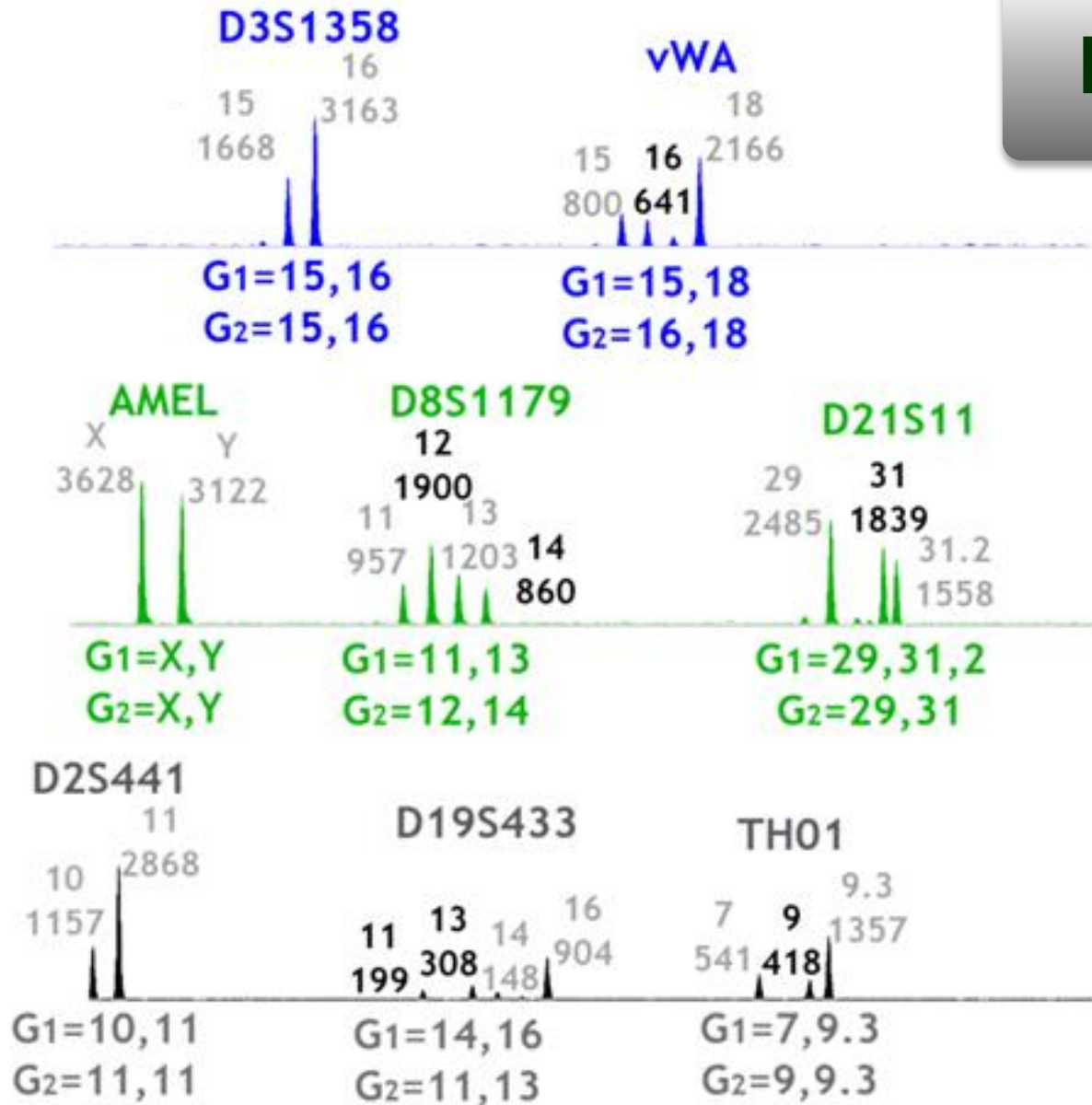


# The probability of the number of contributors given forensic DNA data



**SOFTGENETICS®**

# DNA MIXTURES AND NUMBER OF CONTRIBUTORS (NoC)

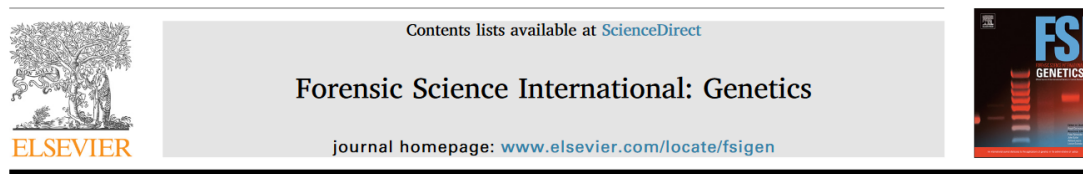


PROVEDIt: 31\_32-1;1-M2a-0.126GF

# NOCIT GIVES $P(N=n|H_D, I, E)$ FOR ALL $N$

## Why $P(N=n|H_d, I, E)$ ?

### The LR is the weighted average of the $n$ -specific LRs, for all $n$



Research paper

Contributors are a nuisance (parameter) for DNA mixture evidence evaluation



K. Slooten<sup>a,\*</sup>, A. Caliebe<sup>b</sup>

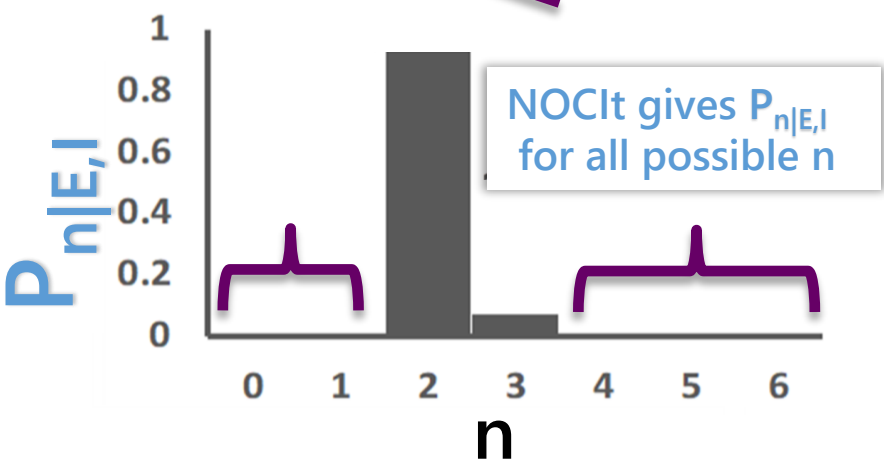
$$LR = \sum_n \left( \underbrace{\frac{P(E|H_p, N=n, I)}{P(E|H_d, N=n, I)}}_{LR^{E|n, I}} \underbrace{P(N=n|H_d, I, E)}_{P_{n|E, I}} \right)$$

$LR^{E|n, I}$

$P_{n|E, I}$



$$LR = LR^{E|n=1, I} P_{n=1|E, I} + LR^{E|n=2, I} P_{n=2|E, I} + LR^{E|n=3, I} P_{n=3|E, I} + LR^{E|n=4, I} P_{n=4|E, I} + LR^{E|n=5, I} P_{n=5|E, I} + \dots$$



NOCIt, therefore, meets 2 aims:

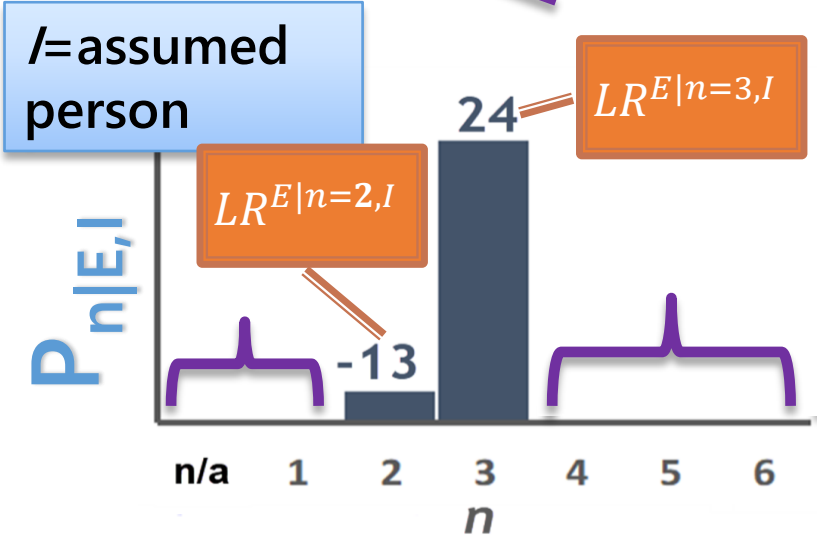
- Narrows  $n$  ranges by informing what  $n$  are associated with negligible  $P_{n|E, I}$
- Supports a process that does not apply default  $n$  or automatic  $P_{n|E, I}$

# NOCIT DETERMINES $P(N=n|H_D, I, E)$

Why  $P(N=n|H_D, I, E)$  aka  $P_{n|E, I}$ ?

Contexts between  $LR^{E|n, I}$  and  $P_{n|E, I}$  are consistent

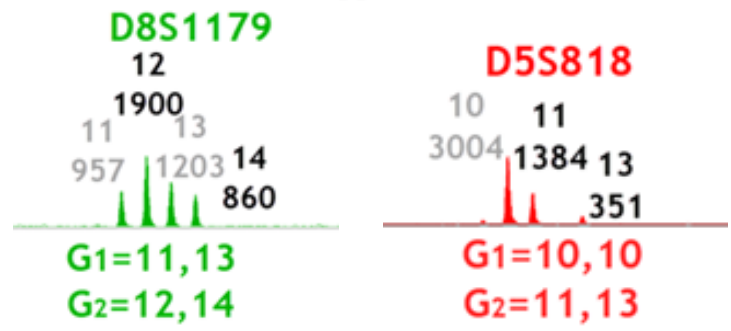
$$LR = LR^{E|n=1, I} P_{n=1|E, I} + LR^{E|n=2, I} P_{n=2|E, I} + LR^{E|n=3, I} P_{n=3|E, I} + LR^{E|n=4, I} P_{n=4|E, I} + LR^{E|n=5, I} P_{n=5|E, I} + \dots$$



$$LR = LR^{E|n=2, I} P_{n=2|E, I} + LR^{E|n=3, I} P_{n=3|E, I}$$

$$= 10^{-13} 0.1 + 10^{24} 0.9$$

$$= 10^{24}$$



NOCIt, therefore, meets a 3<sup>rd</sup> aim:

- Determines  $P_{n|E, I}$  using the *same context* as assigned to  $LR^{E|n, I}$

# NOCIT DETERMINES $P(N=n|H_D, I, E)$

## What is $P(N=n|H_d, I, E)$ ?

It is the posterior probability of  $n$  contributors given the data and a context

Probability of the data given  $n$  randomers

Probability that  $n$  people contributed (before seeing data - PRIOR)

$$P(N = n|H_d, I, E) = \frac{P(E|H_d, I, N = n) \cdot P(N = n|H_d, I)}{\sum_n P(E|H_d, I, N = n) \cdot P(N = n|H_d, I)}$$

Probability of  $n$  contributors after considering the data - POSTERIOR

For ease, drop  $I$  and  $H_d$  going forward

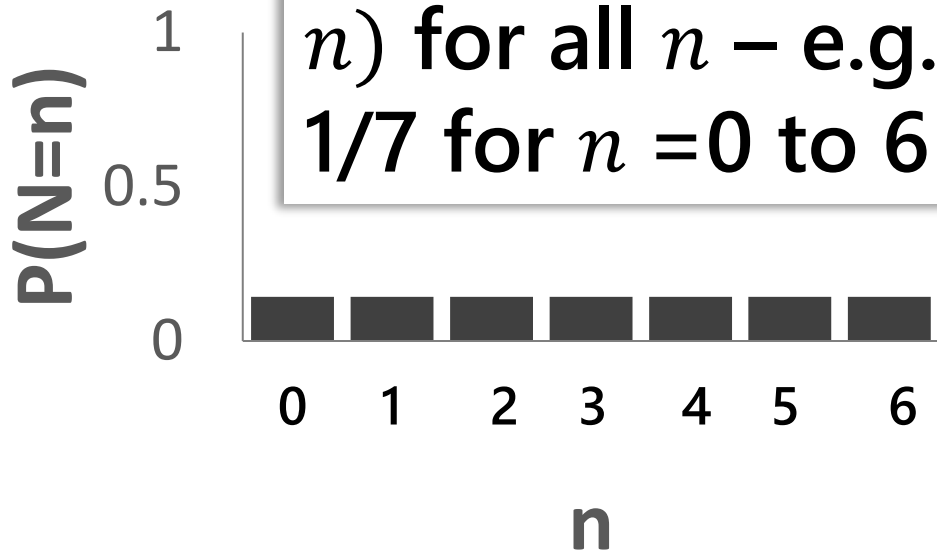
$$P(N = n|E) = \frac{P(E|N = n) \cdot P(N = n)}{\sum_n P(E|N = n) \cdot P(N = n)}$$

# NOCIT DETERMINES $P(N=n|H_D, I, E)$

What is  $P(N=n|E)$  graphically?

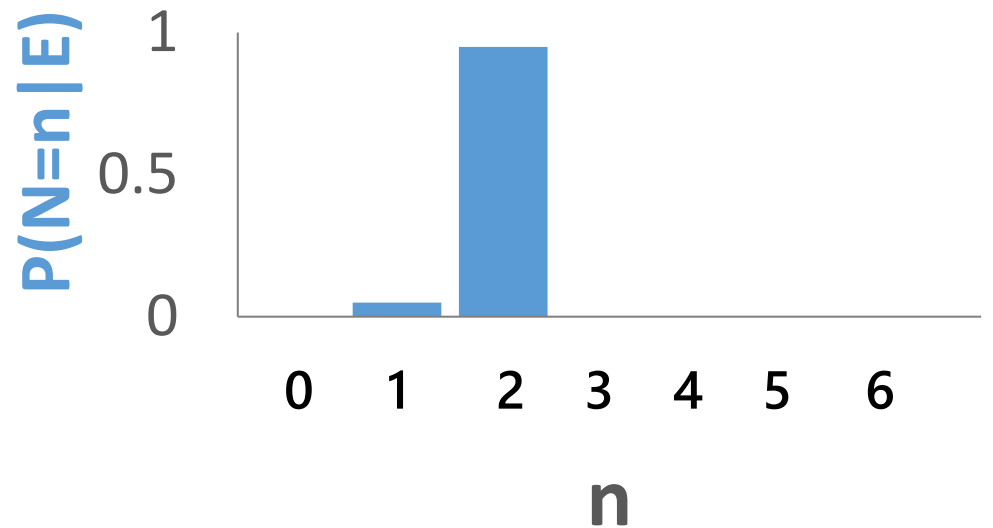
$$P(N = n|E) = \frac{P(E|N = n) \cdot P(N = n)}{\sum_n P(E|N = n) \cdot P(N = n)}$$

Before seeing data, set  $P(N = n)$  for all  $n$  – e.g.,  $1/7$  for  $n = 0$  to  $6$



Data, E

NOCIT,  
 $P(E|N=n)$



# What is $P(N=n | E)$ numerically?

$$P(N = n | E) = \frac{P(E | N = n) \cdot P(N = n)}{\sum_n P(E | N = n) \cdot P(N = n)}$$

Illustrative example:

$n$	$P(N = n)$	$P(E   N = n)$	$P(E   N = n) \cdot P(N = n)$	$P(N = n   E)$
0	1/7=0.143	0.00001	0.00000143	=0.0000014/0.0159 =0.00009
1	0.143	0.01	0.00143	0.09
2	0.143	0.1	0.0143	0.9
3	0.143	0.001	0.000143	0.009
4	0.143	0.000001	0.000000143	0.000009
5	0.143	0.0000001	1.43E-08	0.0000009
6	0.143	0.00000001	1.43E-09	0.000000009
			<b>Sum=0.0159</b>	<b>Sum=1</b>

# LARGE-SCALE VALIDATION WITH PUBLIC DATA

- Confirming NOCI meets **predetermined** expectations
  - Unimodal distributions
    - The distribution should be peaked in one location
  - Precision
    - The apex of the distribution is the same for > 95% of the samples across 3 runs
  - Accuracy
    - $P(N=TrueNOC|H_d, I, E) > 1\%$  for at least 90% samples
      - TrueNOC was confirmed by running all dilutions as single-source and confirming signal
      - Used ReSOLVIt to set up a lab pipeline with LOD=1
  - Comparison
    - NOCI outperformed current procedures
  - Robustness:
    - Resilient to sloping – i.e., degradation/inhibition effects
    - Works under different contexts



# PUBLIC MIXTURE DATA WITH LOD=1 ARE AVAILABLE

- 815 PROVEDIt samples ([www.lftdi.com](http://www.lftdi.com))
  - GlobalFiler samples (29 cycles; 25 sec 3500 Genetic Analyzer)

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## 815 GlobalFiler® samples used to validate NOCIt

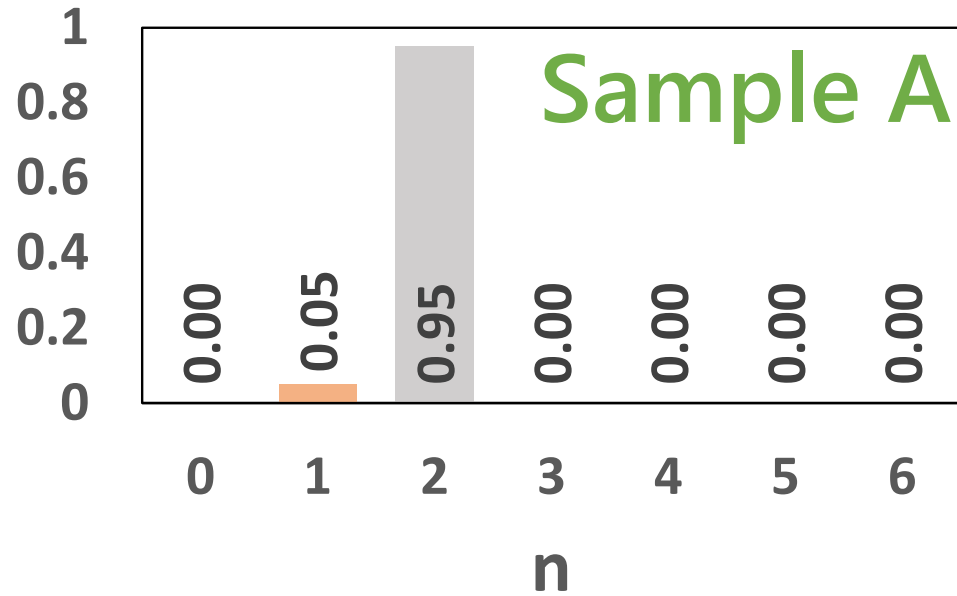
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NOC	1	2	3	4	5
No. Samples	100	193	170	186	166
Tot. Template Mass (ng)	0.5 – 0.0078	0.75 – 0.03	0.75 – 0.045	0.75 – 0.06	0.75 – 0.075
Contributor Ratio	N/A	1:1 – 1:9	1:1:1 – 1:9:9	1:1:1:1 – 1:9:9:1	1:1:1:1:1 – 1:9:9:9:1

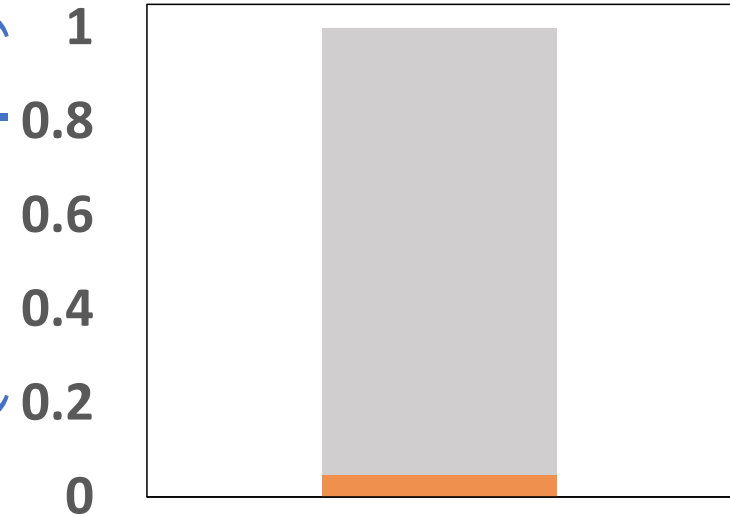
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# VISUAL EXPLORATION IS AIDED BY STACKED PLOTS

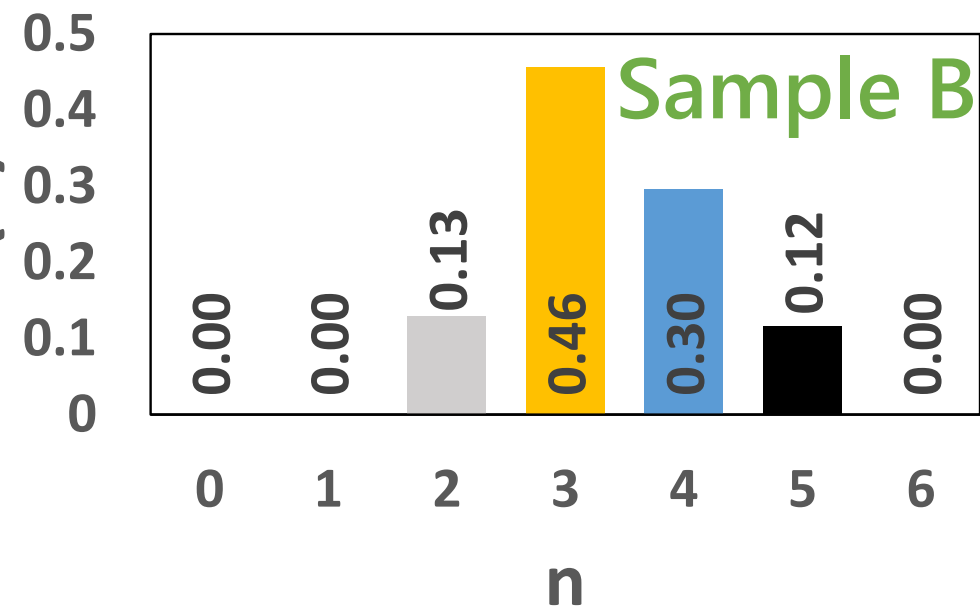
$P(N = n|E)$



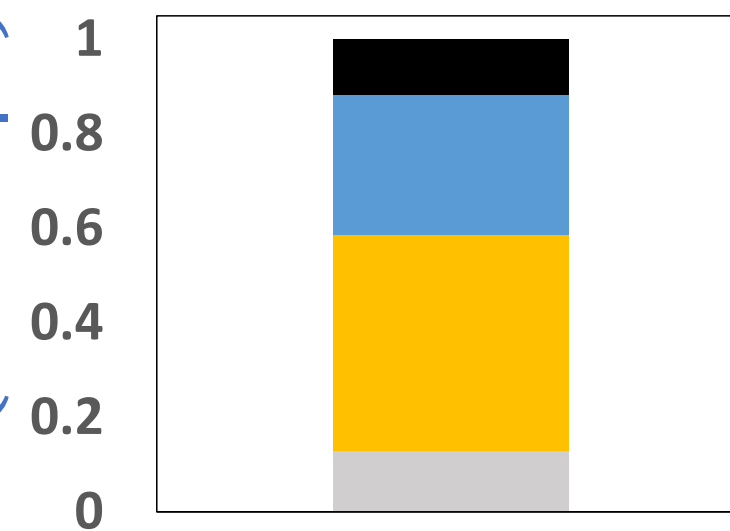
$P(N = n|E)$



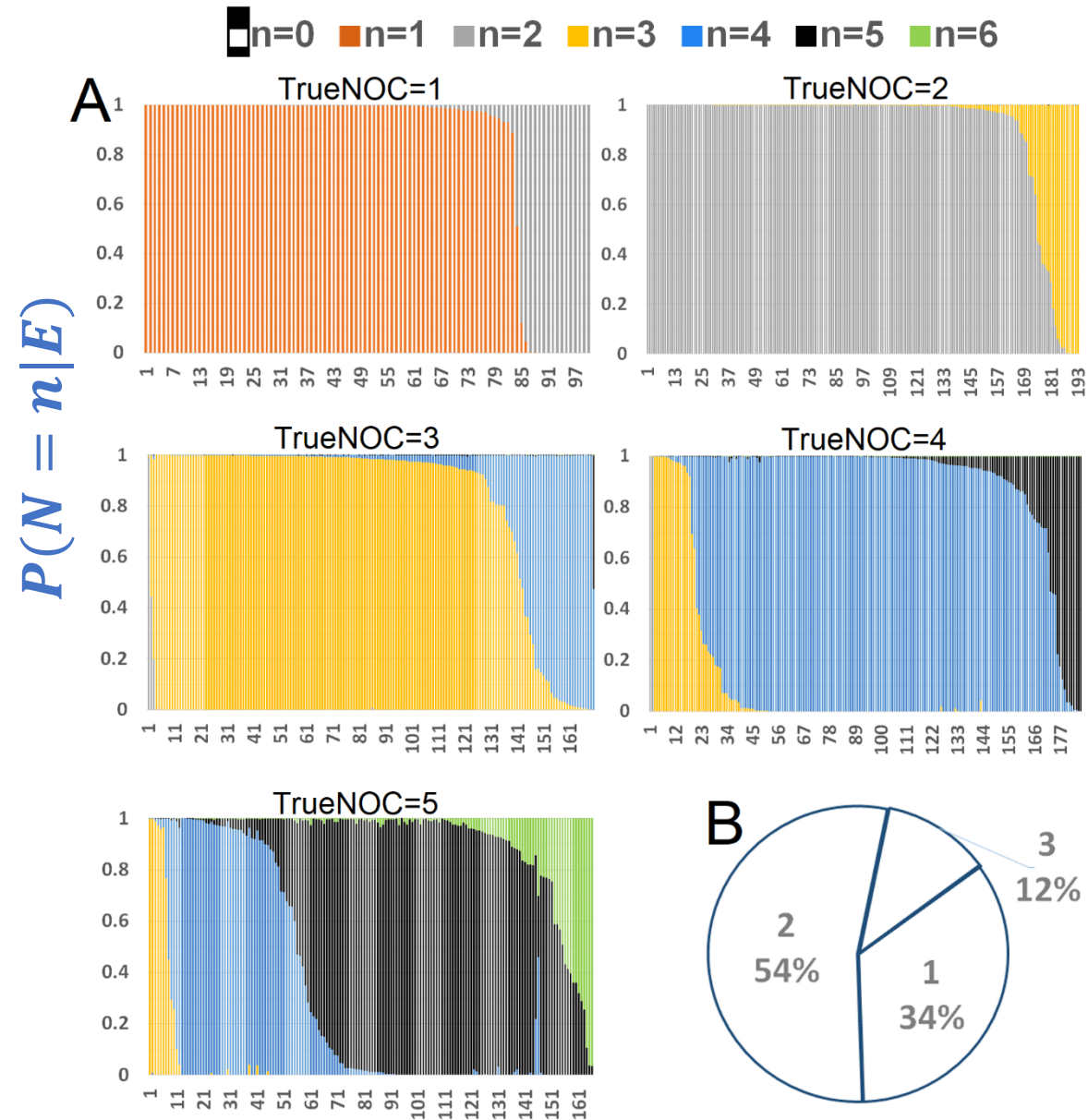
$P(N = n|E)$



$P(N = n|E)$



# NOCIT GIVES UNIMODAL DISTRIBUTIONS



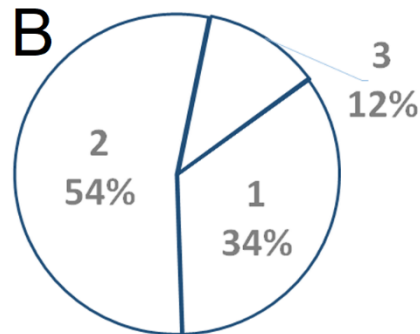
(A) Stacked Plots of APP( $n$ ) using *Condition 1* and the APP for  $n=$  (white bar)0; (orange)1; (grey)2; (yellow)3; (blue)4; and (black) 5; and (green) 6. X-axis is sample number.

(B) Pie Chart depicting percentage of samples resulting in one, two or three APP( $n$ )  $\geq 0.001$ .

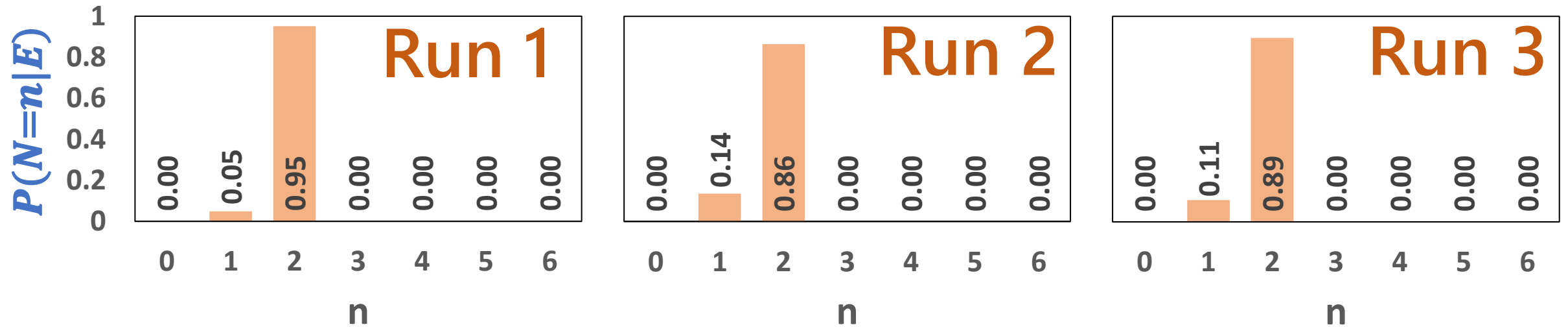
**Criterion:** The distribution should be peaked in one location (at one  $n$ )

**Results:** Distribution was always unimodal in that there was no instance where the probability was high for small values of  $n$ , low for medium  $n$  values and then high for large  $n$ .

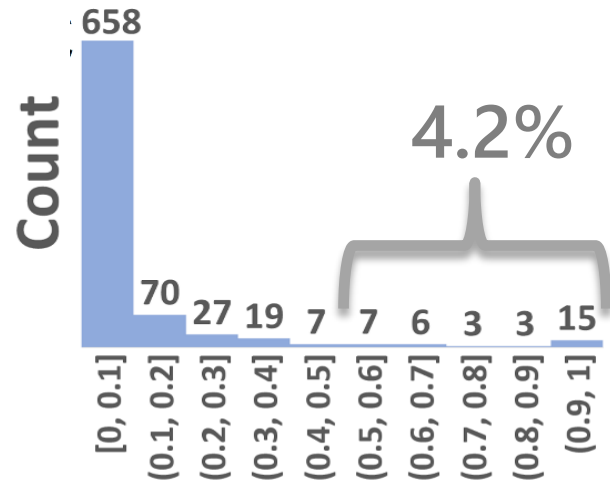
No sample gave more than 3  $P(N = n|E) \geq 0.001$



# NOCIT IS PRECISE ACROSS RUNS



**APP Range.** Record  $\max P(n)$  from Run 1 and determine max delta at that  $n$ .  $P(2)_{R1} = 0.95$ ,  $R2 = 0.86$ ,  $R3 = 0.89$ .  $\text{Range} = (0.95 - 0.86) = 0.09$ . Tells us if  $\max P(n)$  is changing.



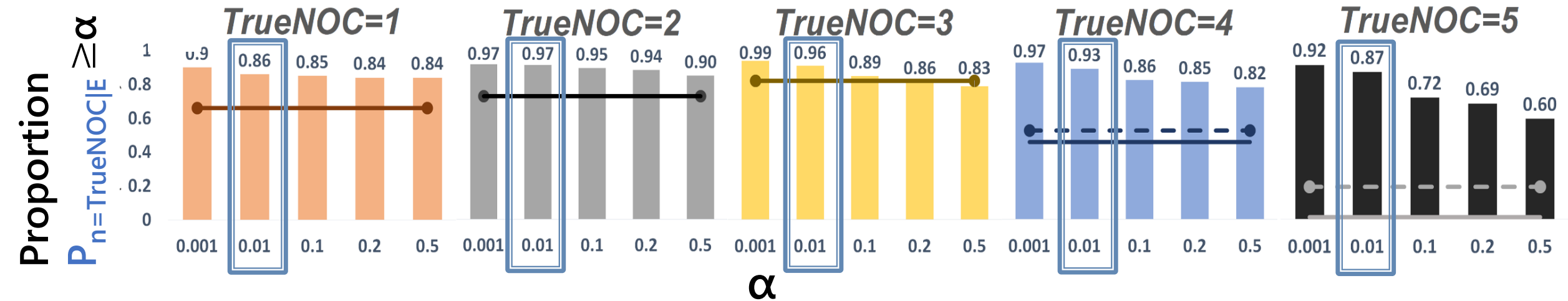
**Criterion:** The distribution's apex located at the same  $n$  for  $> 95\%$  of the samples across 3 runs

**Results:** 95.8% of the samples had repeatable distributions across 3 runs

$$\text{Range} = \max(\Delta P_{R1,R2}; \Delta P_{R1,R3}; \Delta P_{R2,R3})$$

# NOCIt Is ACCURATE, OUTPERFORMING CURRENT PROCEDURES

Proportion of samples giving  $P(N = TrueNOC|E) \geq \alpha$ , and proportion of samples for which (—) MAC and (- -) MLE (H. Haned, et al., J Forensic Sci 56(1) (2011)) estimates equaled TrueNOC



**Criterion:**  $P(N = TrueNOC|E) \geq 1\%$  for at least 90% samples

Results: 92.5% of the samples gave  $P(N = TrueNOC|H_d, I, E) \geq 1\%$

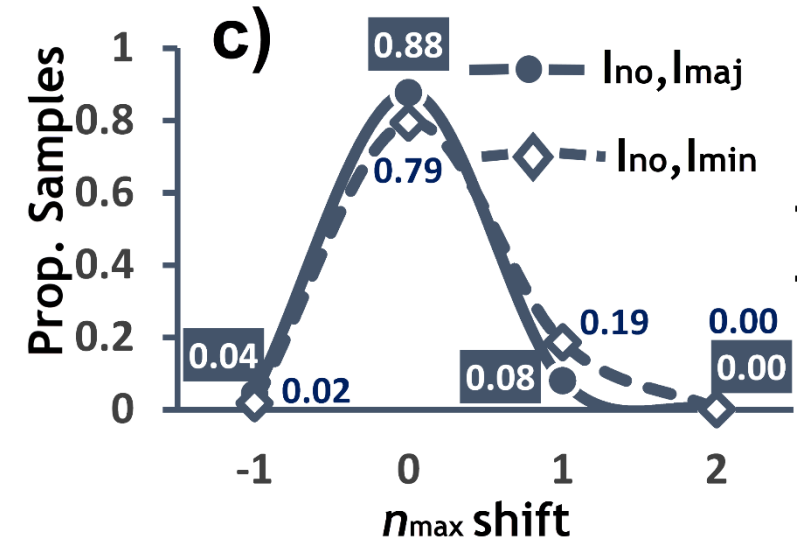
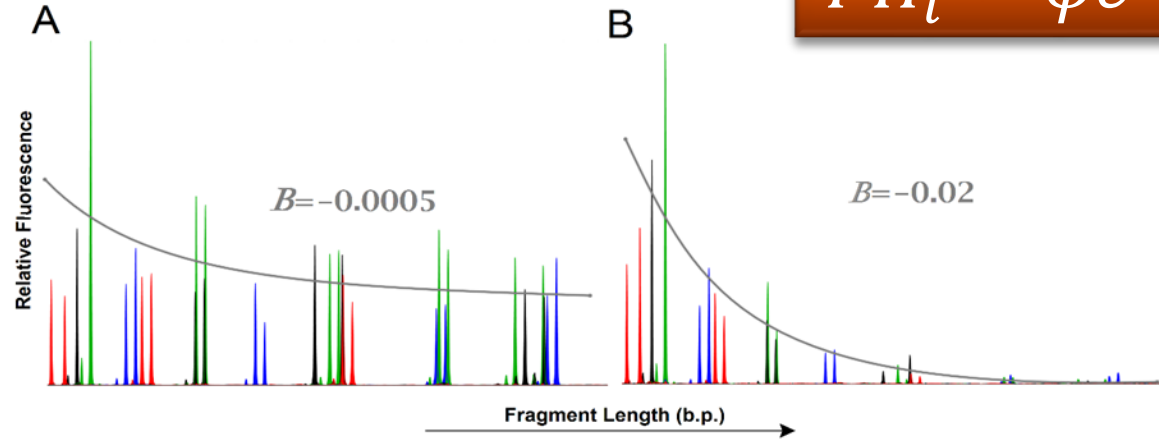
**Criterion:** NOCIt must outperform current procedures

Results: NOCIt performance equals or exceeds current procedures at all  $\alpha$

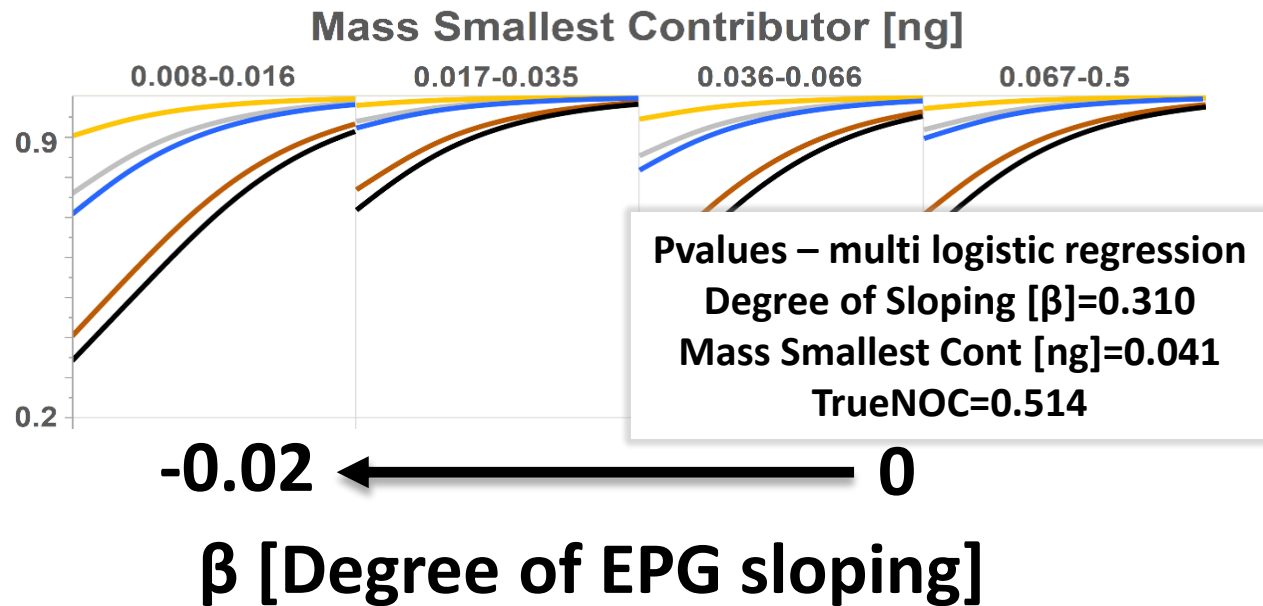
# NOCIT IS ROBUST ACROSS CONTEXTS AND DNA

## QUALITIES

$$PH_l = \varphi e^{\beta \bar{w}_l}$$



Prob  $P(N = \text{TrueNOC}|E) > 0.001$



**Criterion:** Resilient to sloping – i.e., degradation/inhibition effects

**Results:** Pvalue of 0.310 suggests sloping  $[\beta]$  does not significantly affect probability of including TrueNOC when  $\alpha=0.001$  (or 0.5)

**Criterion:** Works under different contexts

**Results:** Most apexes shifted when a known contributor was assumed and it was the minor

# SUMMARY

NOCIt reports  $P(N = n|E)$  – i.e., the posterior distribution for all  $n$  up to 6

**Supports pipelines that do not apply default or subjective  $n$ , or automatic  $P(N = n|E)$**

Helps target  $n$  with non-negligible  $P(N = n|E)$

**Engineered to use all data (even noise)**

Full descriptions in:



Research paper

A large-scale validation of NOCIt's *a posteriori* probability of the number of contributors and its integration into forensic interpretation pipelines

Catherine M. Grgicak<sup>a,b,\*</sup>, Slim Karkar<sup>b</sup>, Xia Yearwood-Garcia<sup>c</sup>, Lauren E. Alfonse<sup>c</sup>, Ken R. Duffy<sup>d</sup>, Desmond S. Lun<sup>b,e,f</sup>



Research paper

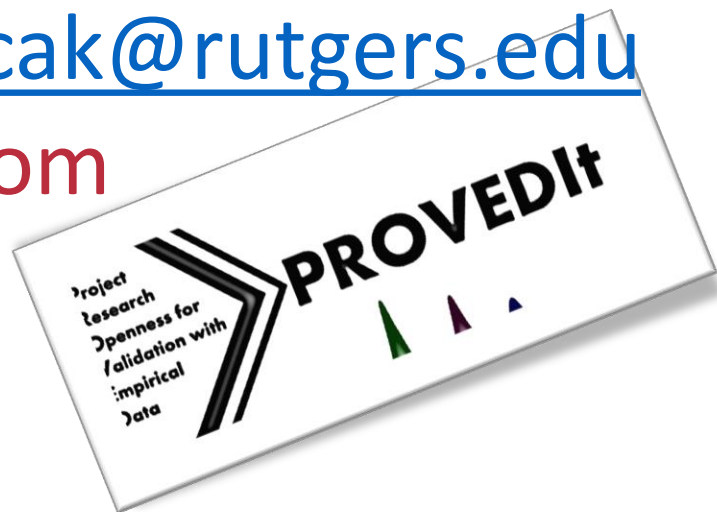
The *a posteriori* probability of the number of contributors when conditioned on an assumed contributor

# FUNDING, COLLABORATORS & STUDENTS

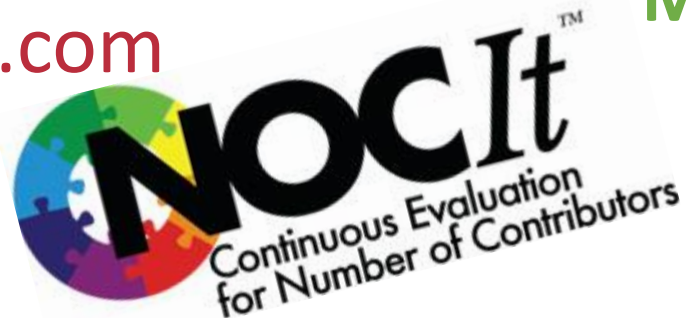
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