



Netherlands Forensic Institute
Ministry of Security and Justice

Evaluating drop-out probabilities

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Estimating the probability of dropout

Given the observed profiles, what is the probability that an allele at a given locus has dropped-out?

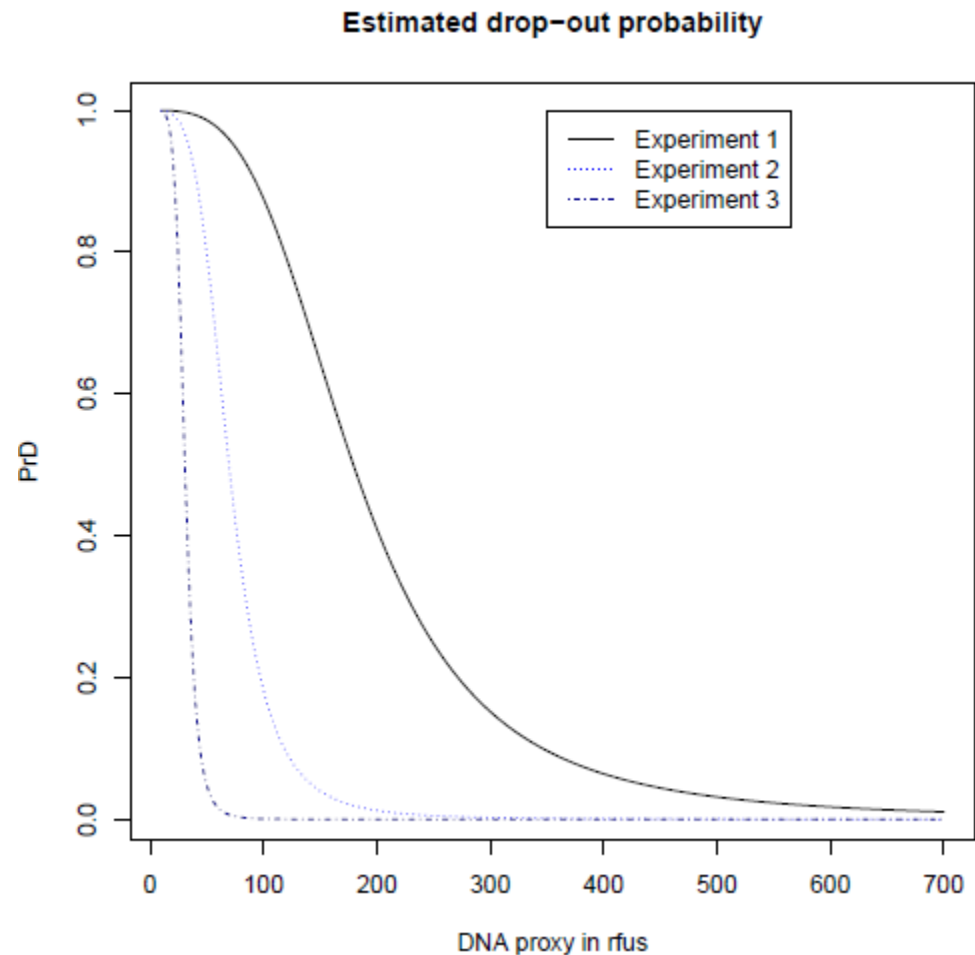
Recent contributions based on the logistic regression suggest predicting the probability of dropout from a measure of the quality of the DNA profiles, based on peak heights:

- DNA proxy model: based on an estimation of the amount of DNA contributed to the stain [Tvedebrink et al. FSIG, 2009](#)
- Surviving peak heights model: based on heterozygote dropout [Gill et al, FSIG, 2009](#)



Logistic regression

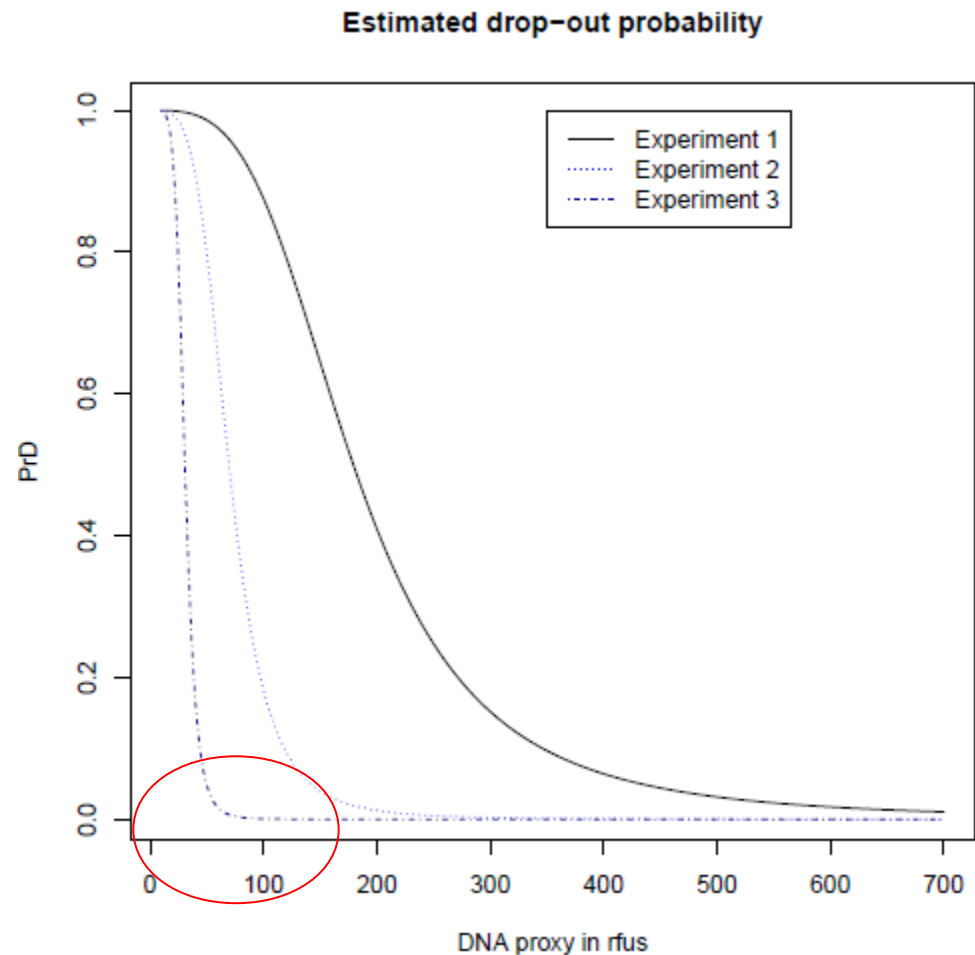
► Repeated experiments yield slightly different logistic curves describing the decay of dropout





Logistic regression

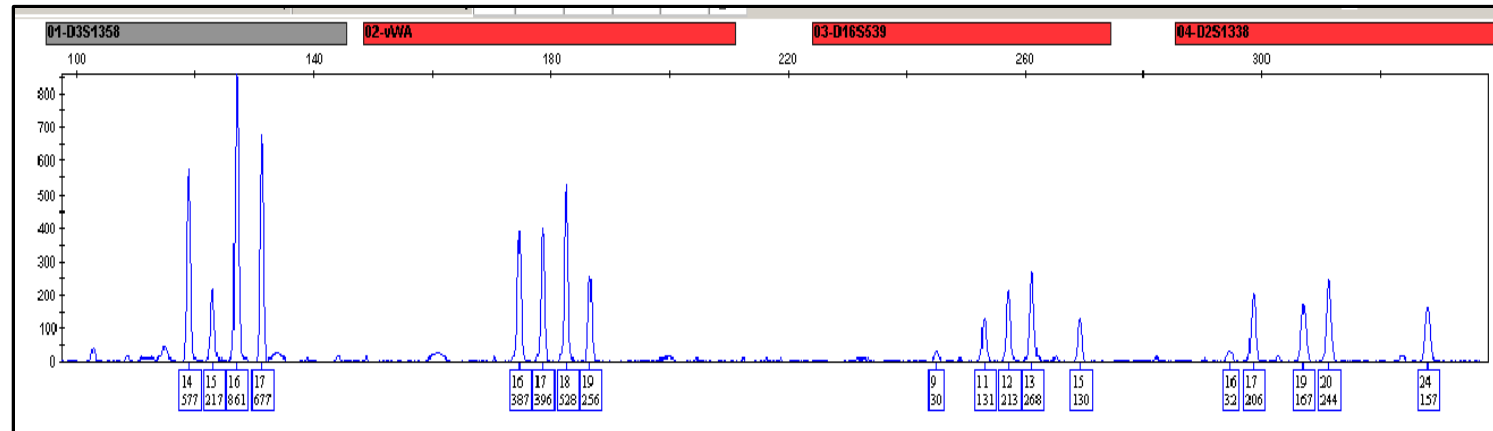
- Repeated experiments yield slightly different logistic curves describing the decay of dropout
- Critical range in LTDNA samples: 50-200 rfus (NGM system)





A three-person mixture

- sample: epithelial cells recovered from the victim of an assault
- two suspects are detained by the police





A three-person mixture

Hypotheses

- H_p : the victim, suspect 1 and one unknown contributed to the sample
- H_d : the victim and two unknowns contributed to the sample

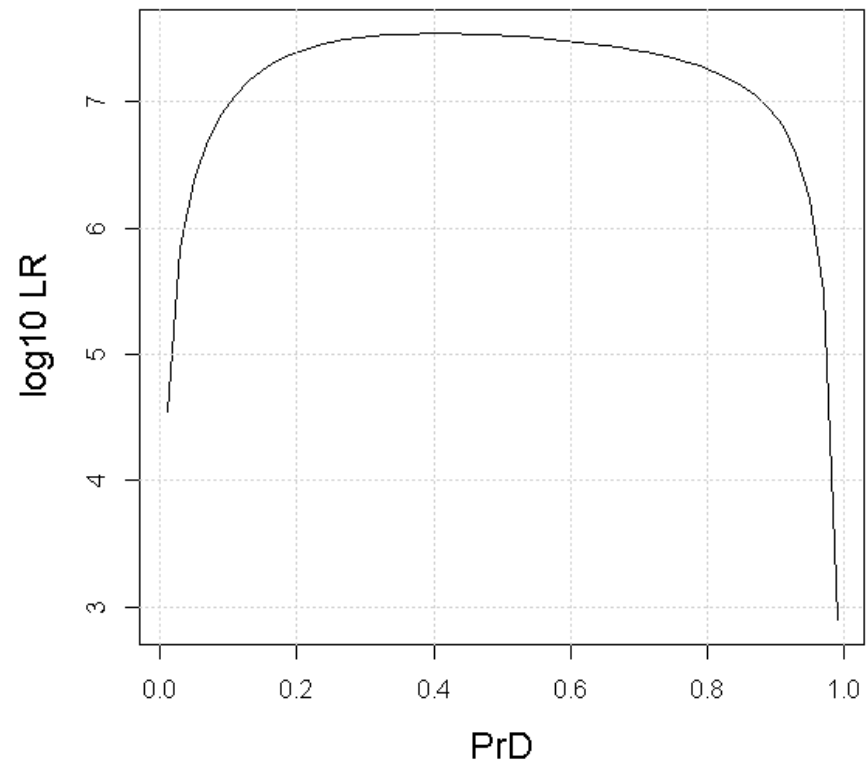
➔ Evaluation of the two hypotheses using likelihood ratios



Sensitivity analysis - LRmix

LRs range from 10^3 to 10^7 :

- $10^3 \Rightarrow \text{PrD}=0.99$
- $10^7 \Rightarrow \text{PrD}=0.41$





Available methods

- Experimental mixtures (Perez et al, Coratian Med J, 2011)
 - the levels of drop-out, based on large sets of DNA mixtures obtained in different conditions
 - DNA quants
 - Number of contributors
 - Ratio of contribution

- Maximum likelihood principle LoComation software (Gill et al 2007)
 - Drop-out probabilities that maximize the probability of observing the questioned epg



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Methods derive estimates from empirical distributions



Qualitative approach to the estimation of PrD

Relies on:

- the number of alleles observed in the sample
- the genotypes of the hypothesized contributors under H

?

What are the probabilities of dropout that could have led to observing the same number of alleles recovered in the sample?



Qualitative approach to the estimation of PrD

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- the number of alleles observed in the sample
- the genotypes of the hypothesized contributors under H

?

What are the probabilities of dropout that could have led to observing the same number of alleles recovered in the sample?

➡ What is the distribution of the number of alleles for the questioned sample, conditioned on PrD?



Qualitative approach to the estimation of PrD

?

What are the probabilities of dropout that could have led to observing the same number of alleles recovered in the sample?

We don't know the probabilities of drop-out, but we can evaluate the drop-out probabilities that could have led to a mixture similar to the one we are investigating



Qualitative approach to the estimation of PrD

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What are the probabilities of dropout that could have led to observing the same number of alleles recovered in the sample?

We don't know the probabilities of drop-out, but we can evaluate the drop-out probabilities that could have led to a mixture similar to the one we are investigating



Build the empirical distributions of the numbers of alleles, conditioned on the probabilities of dropout ranging in $[0,1]$ using **Monte-Carlo simulations**



Monte Carlo method

Any method which solves a problem by generating suitable random numbers and observing that fraction of the numbers obeying some property or properties.



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Questioned sample properties:

- three-person mixture: SGM+
- **33** alleles observed in the epg
- profiles of **victim** and **suspect1** available



Monte Carlo method

Any method which solves a problem by generating suitable random numbers and observing that fraction of the numbers obeying some property or properties.

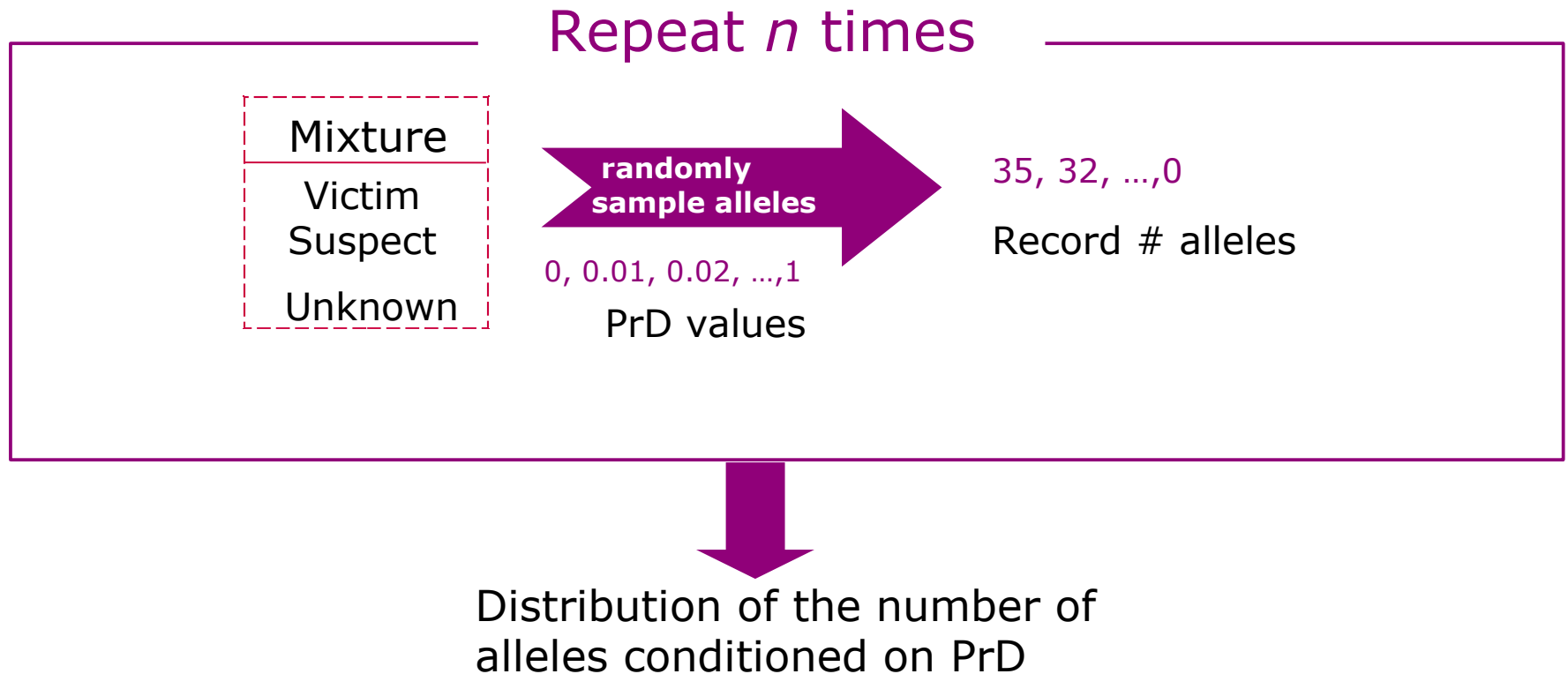
Questioned sample properties:

- three-person mixture: SGM+
- **33** alleles observed in the epg
- profiles of **victim** and **suspect1** available

➡ Simulate a large number of mixtures that have these properties



Monte-Carlo simulation procedure





Important note!

The hypothesized contributors change under H_p and under H_d :

- Derive distribution of the numbers of alleles under H_p and under H_d separately
- Yields two distributions, one under H_p and one under H_d



Monte-Carlo simulation procedure

Under H_p

Repeat n times

Mixture

Victim

Suspect

Unknown

randomly
sample alleles

0, 0.01, 0.02, ..., 1

PrD values

35, 32, ..., 0

Record # alleles

Distribution of the number of
alleles conditioned on PrD



Monte-Carlo simulation procedure

Under H_d

Repeat n times

Mixture

Victim

Unknown 1

Unknown 2

randomly
sample alleles

0, 0.01, 0.02, ..., 1

PrD values

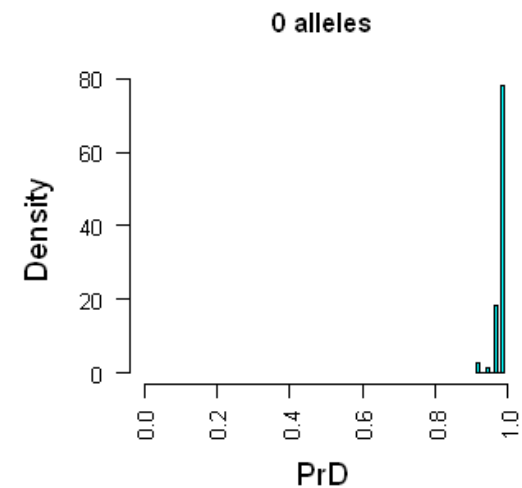
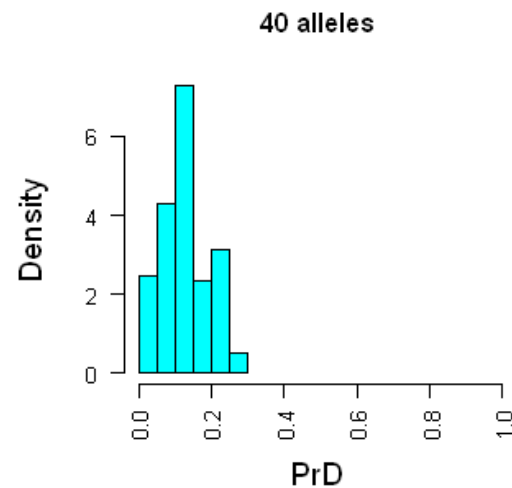
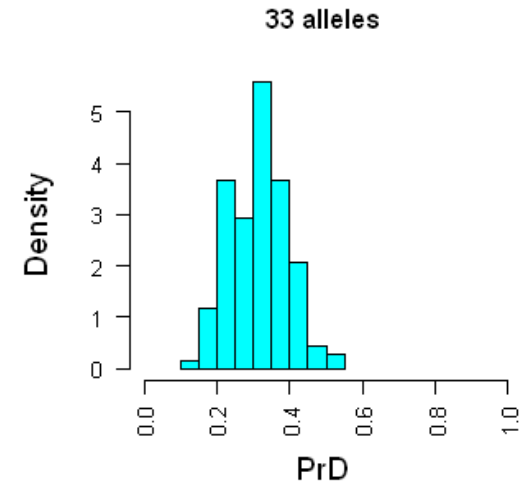
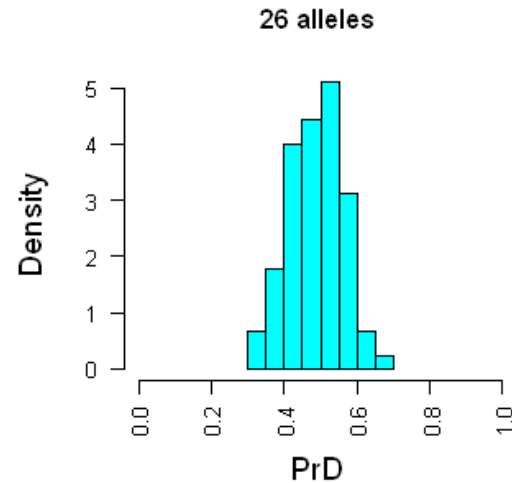
35, 32, ..., 0

Record # alleles

Distribution of the number of
alleles conditioned on PrD

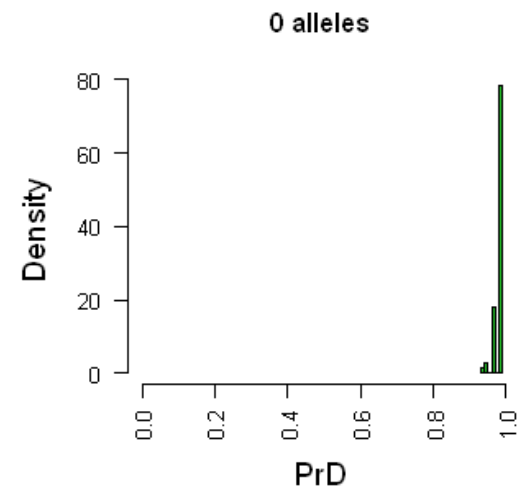
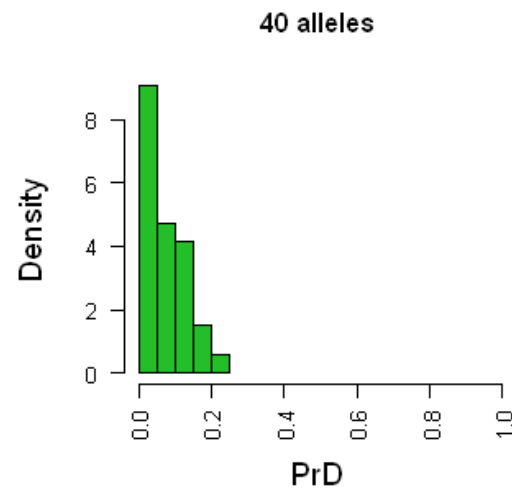
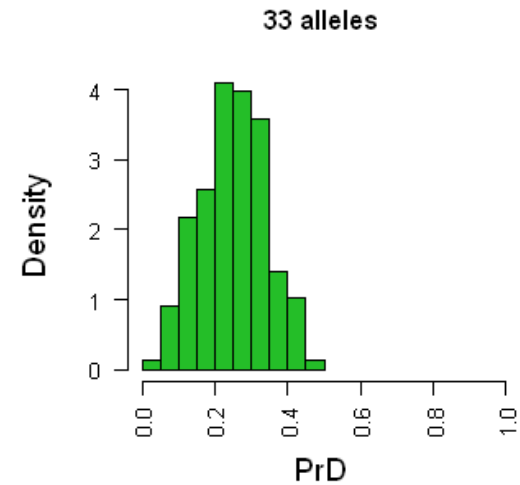
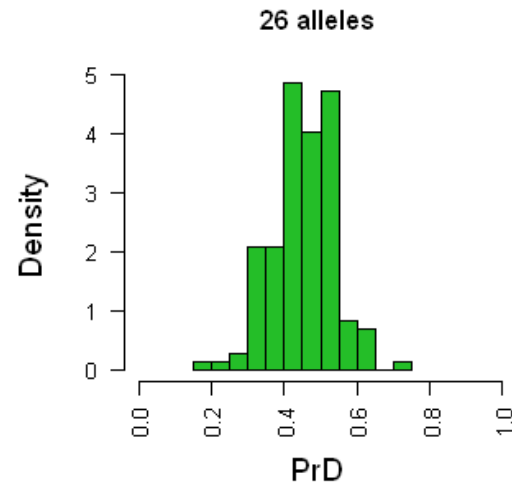


Under H_p





Under Hd





5% - 95% percentiles of the distributions

Under Hp

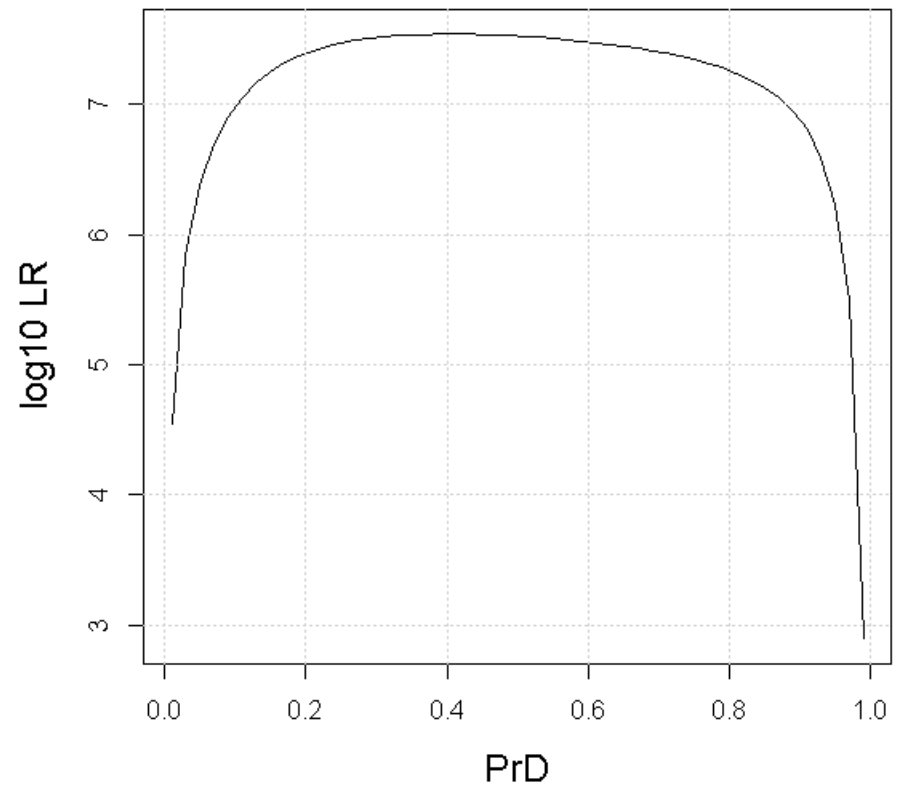
5%	0.19
95%	0.45

Under Hd

5%	0.09
95%	0.41



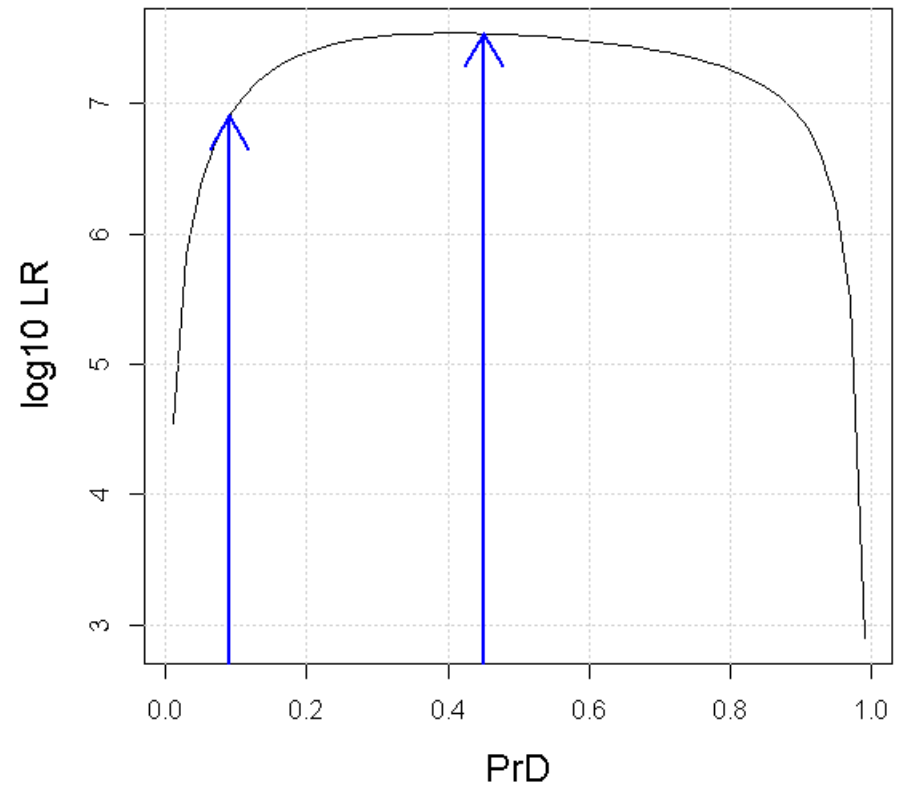
Sensitivity analysis





Sensitivity analysis vs. plausible ranges for PrD

$$LR \cong 10^7$$





Summary

- the ranges of the drop-out probability can be evaluated separately under H_p and H_d
- avoid reporting values of drop-out that are supported by one hypothesis but not by its alternative
- qualitative data only
- peaks slightly under threshold not taken into account

Assess the uncertainty the data!



LRmix module

Available in Forensim 3.1

Analyse the profiles

Hypotheses

Contributors under Hp

Suspect1

☒ Victim

Contributors under Hd

☒ Victim

Parameters

Unknown contributors

Under Hp

Under Hd

Pr(D), Pr(C), theta

Probability of Dropout Pr(D)

Probability of Contamination Pr(C)

Theta Correction (Fst)

Tippet plots

Choose suspect

Suspect1 ☒

number of iterations

