

Correction

Correction: Three-parameter lognormal distribution ubiquitously found in cDNA microarray data and its application to parametric data treatment

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Correction to formulae in methods section [1]

The lognormal distribution model and estimation of the parameters

The method assumes that the original intensity data, (r_i) for $i = 1, 2, \dots, n$, obey a lognormal distribution. The probability density function of the intensity data used was:

$$f(r_i) = [k / \{(2\pi)^{1/2} \sigma(r_i - \gamma)\}] \exp [-\{\log(r_i - \gamma) - \mu\}^2 / 2\sigma^2] \text{ for } r_i > \gamma,$$

where k is a compensation constant ($k = \log e = 0.4343$), σ and μ are the shape and scale parameters for $\log(r_i - \gamma)$, respectively.

The threshold parameter, γ , was found through trial and improvement calculation processes; in the trial, the distribution of $\log(r_i - \gamma)$ was checked by normal probability plotting, and the value that gave the best fit to the model was selected for γ . The fitness was evaluated by the sum of absolute differences between the model and $\log(r_i - \gamma)$, within the interquartile range of data. The parameter μ was found as the median of $\log(r_i - \gamma)$, and the parameter σ was found from the interquartile range of $\log(r_i - \gamma)$; these are known as robust alternatives for the arithmetic mean and standard deviation, respectively. Parameters μ and σ were found for each data grid, a group of data for DNA spots that were printed by an identical pin in order to avoid divergences caused by pin-based differences. Z-normalization was carried out for each datum as

$$Z_{ri} = \{\log(r_i - \gamma) - \mu\} / \sigma.$$

Intensity data (r_i) less than γ were treated as "data not detected", since such data might contain negative noise larger than the signal (see Results).

References

1. Tomokazu Konishi : **Three-parameter lognormal distribution ubiquitously found in cDNA microarray data and its application to parametric data treatment.** *BMC Bioinformatics* 2004, **5**:5.