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**DNA mixtures interpretation - A proof-of-concept multi-software comparison highlighting different probabilistic methods' performances on challenging samples**

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### Abstract

The present study investigated the capabilities and performances of semi-continuous and fully-continuous probabilistic approaches to DNA mixtures interpretation, particularly when dealing with Low-Template DNA mixtures. Five statistical interpretation software, such as Lab Retriever and LRmix Studio – involving semi-continuous algorithms – and DNA•VIEW®, EuroForMix and STRmix™ – employing fully-continuous formulas – were employed to calculate likelihood ratio, comparing the prosecution and the defense hypotheses relative to a series of on-purpose prepared DNA mixtures that respectively contained 2 and 3 known contributors. National Institute of Standards and Technologies (NIST) certified templates were used for samples set up, which contained different DNA amounts for each contributor. 2-person mixtures have been prepared with proportions equal to 1:1, 19:1 and 1:19 in terms of DNA concentration. Conversely, three person mixtures were constituted by proportions equal to 20:9:1, 8:1:1, 6:3:1 and 1:1:1 in terms of DNA concentration. Furthermore, 8 equally-proportioned 3-person mixtures were prepared by means of scalar dilutions starting from an overall amount of 0.500 ng, then ranging up to DNA samples with concentrations equal to 0.004 ng (i.e. Low-Template DNA). DNA mixtures were set up in triplicate and amplified with 7 DNA amplification kits (i.e. GlobalFiler PCR Amplification Kit, NGM SElect PCR Amplification Kit, MiniFiler PCR Amplification Kit, Power Plex Fusion, PowerPlex 6C Matrix System, Power Plex ESI 17 Fast and Power Plex ESX 17 Fast) in order to evaluate whether the selection of a certain kit might represent a bias factor, capable of altering the whole interpretation process. Multi-software approach helped us to highlight any trend in the likelihood ratio results provided by semi- and fully-continuous software. As a matter of fact, fully-continuous computations provided different results in terms of degrees of magnitude of the likelihood ratio values with respect to the ones from the semi-continuous approach, regardless of the amplification kit that was utilized.

<b>Keywords</b>	DNA mixture interpretation; Low-Template DNA; semi-continuous model; fully-continuous model; likelihood ratio.
<b>Taxonomy</b>	Criminal Casework, DNA Polymorphism
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## Submission Files Included in this PDF

### File Name [File Type]

Cover Letter.docx [Cover Letter]

Response to reviewers.docx [Response to Reviewers (without Author Details)]

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Figure 1.tif [Figure]

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## Research Data Related to this Submission

There are no linked research data sets for this submission. The following reason is given:  
Data will be made available on request



Torino, April 9<sup>th</sup>, 2018

**Prof. A. Carracedo**

Editor-in-Chief – **Forensic Science International: Genetics**

Dear Editor-in-Chief,

This letter accompanies submission to Forensic Science International: Genetics of a manuscript entitled: "*DNA mixtures interpretation – a proof-of-concept multi-software comparison highlighting different probabilistic methods' performances on challenging samples*".

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The present study investigated the capabilities and performances of semi-continuous and fully-continuous probabilistic approaches to DNA mixtures interpretation, particularly when dealing with Low-Template DNA mixtures. Five statistical interpretation software, such as Lab Retriever and LRmix Studio – involving semi-continuous algorithms – and DNA•VIEW<sup>®</sup>, EuroForMix and STRmix<sup>TM</sup> – employing fully-continuous formulas – were employed to calculate likelihood ratio, comparing the prosecution and the defense hypotheses relative to a series of *ad-hoc* prepared DNA mixtures that respectively contained 2 and 3 known contributors, in different proportions. Furthermore, eight equally-proportioned 3-person mixtures were prepared by means of scalar dilutions starting from an overall amount of 0.500 ng, then ranging up to DNA samples with concentrations equal to 0.004 ng (i.e. Low-Template DNA). All samples were performed in triplicate, then amplified by seven DNA amplification kits (i.e. GlobalFiler PCR Amplification Kit, NGM SElect PCR Amplification Kit, MiniFiler PCR Amplification Kit, Power Plex Fusion, PowerPlex 6C Matrix System, Power Plex ESI 17 Fast and Power Plex ESX 17 Fast) in order to evaluate whether the selection of a certain kit might represent a bias factor, capable of altering the whole interpretation process.

**Novelty statement:** This work is new and original and is not under consideration elsewhere. In comparison with the existing literature, the present study represents a proper interpretation approach that fulfil the extreme caution that is demanded in this forensic field, especially when Low-Template DNA and complex mixtures have to be interpreted. In particular, multi-software evaluations of a priori known 2-person and 3-person DNA mixtures prepared in our laboratory allowed us to compare the performance of both semi- and fully-continuous approaches. Log(LR) results provided by the tested fully-continuous software (i.e. DNA•VIEW<sup>®</sup>, EuroForMix and STRmix<sup>TM</sup>) turned always significantly higher than the ones calculated by the employed semi-continuous software (i.e. Lab Retriever and



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LRmix Studio). Obviously, the evaluations relative to these DNA mixtures just represent a proof-of-concept that fully-continuous model seems to be the most suitable bio-statistical methodology to be performed by analysts when Low-Template DNA mixtures have to be interpreted from a probabilistic point-of-view.

Thank you for considering the paper for Forensic Science International: Genetics.

Yours faithfully,

Eugenio Alladio, PhD

## Response to reviewers

### Reviewer #1:

- *This paper reports the interpretation of (I think) 15 mixtures that vary in mixture ratio and template, amplified in 7 kits, and I think with two strategies for setting AT. These are interpreted in five software packages. Think this is the largest intersoftware comparison yet reported. The work concentrates on the true donor tests and does no false donor tests. This is an omission.*

According to the comment made by the reviewer #1, the evaluation of a false donor was included within the text. Results are reported within the Supplementary Material within the Figures S6 and S8. New calculations were made by using a false donor DNA profile provided by NIST (i.e. NIST F reference samples). The results relative to such calculations have been added to the Supplementary Material. The following statement was added within the Materials and Method section *“The DNA profile from NIST F reference sample was used, conversely, as a known non contributor in order to perform false donor tests for all the tested software, too.”*

- *The conclusion seem supported EXCEPT this work really invalidates the “statistic consensus approach” which maximizes the false indication of exclusion rate. This paper gives very good evidence not to use it. The only way that I can see to support it would be to show a consequential reduction in the false inclusion rate. But this has not been investigated. It may be necessary to ask the authors to draw this conclusion.*

In our opinion, the use of a “statistic consensus approach” seems to be useful because experts in courtrooms might show LR results from different software involving different algorithms and probabilistic approaches. Therefore, we believe that a very conservative approach might be employed in order to avoid false inclusion that could lead to unwanted legal consequences. For this reason, due to the complexity and the relevance of the task, we believe that a statistic redundancy in the calculations is preferable at the current stage. The following paper was added as reference in order to support this concept: *“Randles M., Lamb D., Odat E., Taleb-Bendiab A., Distributed redundancy and robustness in complex systems, J. Comput. Syst. Sci. 77 (2011) 293–304. doi:10.1016/j.jcss.2010.01.008”*. Moreover, non-contributor tests were also performed, as suggested by reviewer #2 too, for the three employed reference samples (i.e. NIST A, NIST B and NIST C). These results are included into the Supplementary Material. In order to reply to the correct evaluation of the reviewers, the following statements have been added within the conclusions: *“Despite this fact, extreme caution has to be used when interpreting LT-DNA mixture, especially when different algorithms and probabilistic approaches are used and compared by forensic experts. Consequently, in our opinion, a conservative approach might be employed at the current stage in order to avoid false conclusions that could eventually lead to unwanted severe legal consequences. For this reason, due to the complexity and the relevance of the task, a statistic redundancy [51] in the calculations might be useful (i.e. like our adopted “statistic consensus approach”).*

- Line numbers, as required in authors' instructions, would have been helpful. "Formatting requirements ... Please ensure your paper has consecutive line numbering - this is an essential peer review requirement."

Line numbers have been added to the text, as suggested by reviewer #1.

- drop-out rate [6,9,18–23], stutters rate. fully-continuous approaches. The binary model[1] was the first one employed by the forensic community for DNA mixture interpretation but, over the time, it was gradually replaced. turned to be not suitable since it This method does not take into account neither the instrumental stochastic effects (i.e. drop-in and drop-out -evaluated by the semi-continuous models), nor the peak heights of the detected alleles (evaluated by the fully-continuous models).

had to deal with bigger larger amounts of data

traditional binary approach. For this reasons, semi-continuous [32,33] and fully continuous models [34,35] have largely replaced by far the binary one, and

The corrections suggested by reviewer #1 have been made.

- For this reason, in the previous years semi-continuous models (add LikeLTD [2, 3] reference to LRmix and Labretreiver please) have been largely employed by forensic analysts. since There are actually several open-source software available in literature such as, and methodologies in order to avoid misinterpretations or wrong applications of such algorithms As correctly remarked by reviewer #1, LikeLTD references were added to the manuscript.

- several mixtures were ad hoc (this is not a standard English usage.) prepared and composed by two and three known contributors mixed in different proportions. How about: mixtures of two and three known contributors mixed in different proportions were prepared.

As correctly remarked by reviewer #1, a correction has been made within the whole text.

- Thanks to This experimental plan, it had been possible to even allowed the assessment of whether the performances of the examined software turned to be was influenced or not by the adopted DNA amplification kits.

The correction suggested by reviewer #1 has been made.

- concentrations of DNA (i.e. up to (would this be down to?) proper LT-DNA samples). A real LT-DNA case is also discussed. Moreover, starting from such quantitative, An equally proportioned 1:1:1 mixture containing NIST samples A, B and C was serially diluted in order to obtain several further samples.

The corrections suggested by reviewer #1 have been made.

- and Scientific Working Group on DNA Analysis Methods (SWGDM) (what is the right reference for this? Maybe [4])requirements.

As suggested by reviewer #1 a reference (website) was added to the text.

- concentrations Lab Retriever and LRmix Studio are open-source and free of charge software performing a semi-continuous approach to DNA mixtures.  
 Markov Chain Monte Carlo (MCMC) approach for fully-continuous DNA mixtures interpretation [29,41,45], developed by Taylor, Bright and Buckleton.  
 All the described software were exploited with the aim of used to calculate LR values relative to each one of NIST samples used as known contributor and included into our prepared DNA mixtures.  
 All the corrections suggested by reviewer #1 have been made.
- A validated drop-in value equal to 0.05 (units please, maybe drop-in perlocus, if so this is VERY high).  
 A 0.05 global drop-in rate was observed and calculated during our validation studies. This values was also confirmed during our accreditation process and strvalidator software, too.  
 The term "global" was added within the text.
- Finally, NIST U.S. population dataset (note that this is the uncorrected database[5]) was adopted as reference database in all LR computations [46].  
 The database that was used in our study is the one that was revised in July 2017, according to <https://strbase.nist.gov/NISTpop.htm>. This note was added within its reference for clarification, as suggested by reviewer #1.
- As it can be seen, LR results provided by both semi-continuous models turned very were similar or identical.  
 results provided by fully-continuous models proved similar and convergent to one another, with slightly higher within-software differences (  
 Since log(LR) results turned convergent (convergent is not a standard English usage, you could define it early or maybe just use similar) among the tested approaches  
 be seen, LRFC results turned were always higher than the ones provided by fullysemi-continuous modelling (LRSC) for both F6C and GF  
 Similar increases in the log(LR) results from fully-continuous approach were observed, amplification kits. However, the FC results turned were always higher than the SC ones, regardless  
 All the corrections suggested by reviewer #1 have been made.
- In this case, probabilistic interpretation of 0.004 ng DNA mixture provided log(LR) values lower than zero for both the biostatistical models, May I ask for a bit more here. For example, how many unmasked alleles of the trace contributor were left above AT?  
 The number of unmasked alleles of the trace contributor left above the calculated AT for the different DNA amplification kits were around 5-20%. Different values were obtained according to the DNA amplification kit under evaluation. As a consequence, inconclusive values were "correctly" expected for these DNA mixtures. This comment has been added to the text, too.



- with the exception of FC  $\log(LR)$  result relative to NIST B known contributor equal to 1.9. Moreover, a  $\log(LR)$  value of -1.47 was observed for NIST C contributor in the mixture containing 0.008 ng of DNA. Once again, these observations proved a better sensitivity of the fully-continuous models in case of LT-DNA.

In the present paragraph section  $\log_{10}(LR)$  results relative to a forensic real casework that was evaluated in our laboratory will be discussed. In details, a Caucasian individual was charged as an alleged suspect (POI) for a series of robberies and thefts. A cap was collected on from a crime scene by police forces during their investigating activities. Since it was supposed to belong to the alleged suspect, flocked swabs Nylon® 4N6 (purchased by COPAN ITALIA S.P.A., Brescia, Italy) were applied on the visor of the cap aiming to detect biological evidences. The genetic material was recovered on different spots on the visor, then extracted from the swabs, amplified and analyzed. Before performing our “statistic consensus approach” over acquired data, differential analytical thresholds were calculated by means of ArmedXpert™ for each dye channel, following our

mixtures obtained were biostatistically interpreted. A summary of  $\log(LR)$  results are reported in results with respect to the ones obtained by adopting SD3 formula. More in details, uncoherent Categoricaly different Results differing on which side of  $\log(LR) = 0$  were observed between semi- and fully-continuous interpretations when evaluating the biological evidence as a 2-person mixture (i.e.  $H(p) = POI + 1$  unknown individual;  $H(d) = 2$ ).

All the corrections suggested by reviewer #1 have been made.

- However, even though fully-continuous software showed quite high  $\log(LR)$  values, semi- and fully-continuous models proved uncoherent one another gave results differing on which side of  $\log(LR) = 0$ . Consequently, according to our adopted “statistic consensus approach”, response expressed at the end of our interpretation process was inconclusive. This approach maximizes the false indication of exclusion rate. This paper gives very good evidence not to use it. The only way that I can see to support it would be to show a consequential reduction in the false inclusion rate. But this has not been investigated.

As remarked by reviewer #1 and showed by the results reported in our study, fully-continuous software provide higher absolute  $\log(LR)$  values with respect to the ones provided by the semi-continuous models. However, when dealing with real caseworks, our opinion is that the forensic expert has to be very conservative, especially when interpreting LT-DNA samples. Consequently, our lab developed, validated and accredited (via ISO17025 requirements) the cited “statistic consensus approach” in order to cross-validate the results provided by the different software and provide a trustful and robust interpretation for real caseworks and, particularly, for complex LT-DNA mixtures. The authors of this manuscript are aware that this is not the best interpretation process but, according to Italian rules, this approach seemed very conservative to us (and to ACCREDIA, the Italian accreditation body, too) in order to hel the expert to provide a conclusion “beyond any reasonable doubt”. Further studies are already under development by our group in order to combine the likelihood ratio results provided by different software and algorithms, using different interpretation weights, but their results are still preliminary and will not be reported in this study. A note has been added within the manuscript, as suggested by reviewer #1.

- *H(d) = 3 unknown individuals). In particular, semi-continuous approaches delivered a moderately strong support to H(p), while fully-continuous models delivered an extremely strong support such hypothesis. In the present case, response emitted the result reported at the end of our interpretation process supported the prosecution hypothesis, charging the POI as an effective contributor to the biological evidence collected on the visor of the cap that was recovered on the crime scene.*

*The correction suggested by reviewer #1 has been made.*

- *Please consider excising this part: At the end of the trial, suspect was convicted as guilty since further evidences incriminated him. We should have no interest in the outcome or the “other evidence.” In our opinion, results reported in Table 2 represent once again a proof to the concept that fully continuous models might be more sensitive than semi-continuous ones in case of LT-DNA mixtures interpretation.*

*In the present casework, fully-continuous log(LR) values always supported the prosecution hypothesis, which was later properly verified by the investigating authorities. Please consider excising: which was later properly verified by the investigating authorities. I cannot see how investigating authorities can verify an LR.*

*The cited sentences have been removed from the text, as correctly remarked by reviewer #1.*

- *related to the different semi- and fully-continuous algorithms. In particular, fully-continuous software takes into account a larger amount of data and information (i.e. detected alleles plus their relative peak heights), so that higher LR values can be obtained when significant matches are observed between the investigated biological samples. Similar trends in log (LR) values were observed when several serially scalarly-diluted 3-person mixtures were investigated, too. These mixtures were ad hoc prepared in order to contain the same DNA amount for each one of the included known contributors. Nevertheless, log (LR) values provided by semi-continuous software gave false indications of exclusion turned uncoherent I cannot translate this word. In the authors mean that SC gave LRs <1 then maybe define this early as something like: false exclusionary indications to the composition of the DNA mixtures, especially in case of Low-Template DNA (i.e. mixtures showing an overall DNA concentration of 0.004 ng, 0.008 ng and 0.016 ng). Furthermore, probabilistic software behaved in a similar way, regardless of the DNA amplification kits that were employed. Even though this outcome was expected, further analyses might indicate which DNA amplification kits would turned out to be the most useful in cases of Low-Template DNA.*

*All the corrections suggested by reviewer #1 have been made.*

- *Obviously, the These evaluations relative to these DNA mixtures just represent a proof-of-concept some evidence that the fully continuous model seems to be the most suitable bio-statistical methodology to be performed by analysts when Low-Template DNA mixtures have to be interpreted from a probabilistic point-of-view.*

*Further experiments (i.e. 4- and 5-person DNA mixtures[6, 7]) and their relative interpretation processes need to be performed, but in our opinion these results open the pathway towards the possibility of “weighting” LR results provided by semi- and fully-continuous models, particularly in case of LT-DNA mixtures interpretation.*

All the corrections suggested by reviewer #1 have been made.

- Table 1 and elsewhere

DNA Typing Kit Reference Material

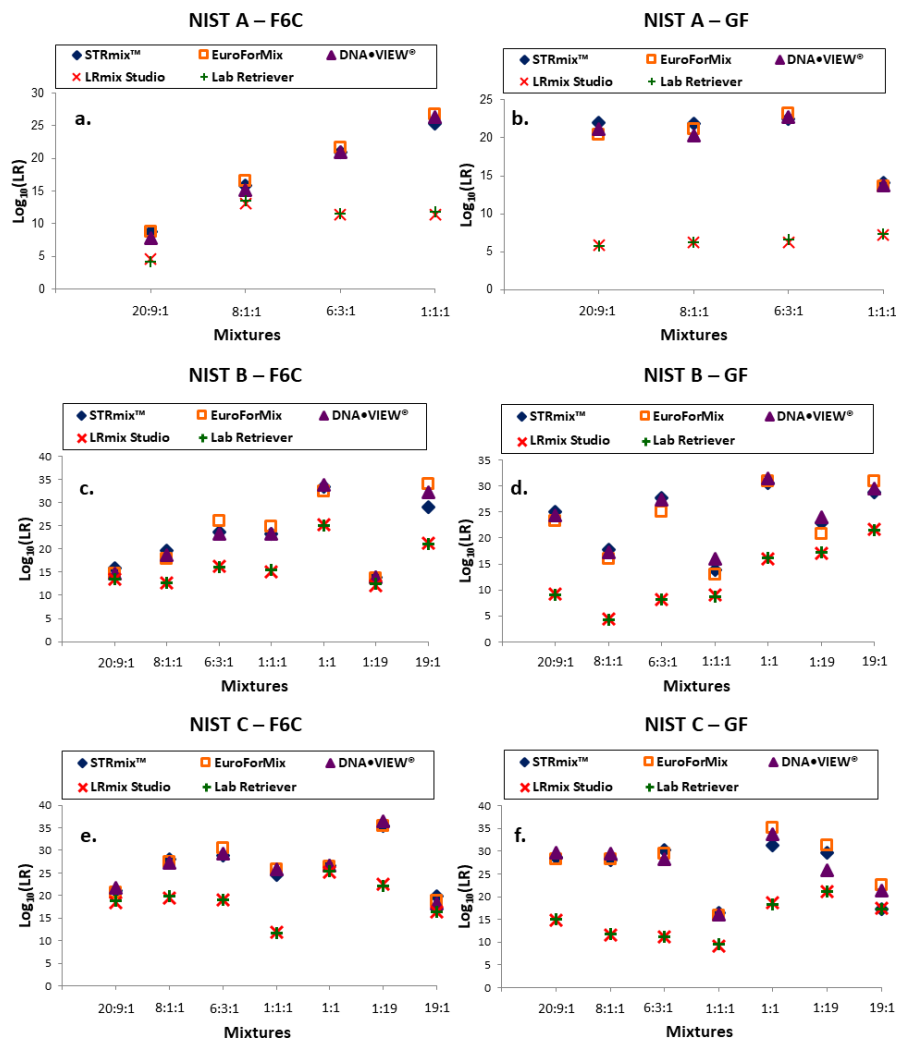
Mixtures proportion I think this is mixture ratios  
(0.500 ng)

Table 2 Log(LR) results relative to the interpretation process performed on the DNA mixture collected on a cap recovered on a crime scene. 3SD and Min-Max represent the algorithms that were employed to evaluate the differential analytical thresholds. POI represents the suspect (i.e. the person of interest), while U stands for unknown(s) individual(s) extracted from the allele frequencies reference dataset [46].

All the corrections suggested by reviewer #1 have been made.

- Figure 1. Top left. I cannot see the STRmix™ results. Can we try open symbols or any other method to show them. This also applies to some of the supplementary materials.

All the cited figures have been modified as suggested by reviewer #1. As an example, the modified Figure 1 is reported, as follows:



- Can I tell which component of the mixture ratio is NIST A, B or C for the three person mixtures. My guess is that it is C, B, A in that order. Please consider making the y-axis the same scale for all graphs. This also applies to some of the supplementary materials. As reported in Table 1, the main contributor of the 3-person mixture is NIST A, followed by NIST B and, then, NIST C. The authors opted to leave the y-axis as it is in order not to reduce the different figures and diminish their readability. However, the y-axis can be easily modified if required by the editor, too.
- Figure 2 Mean  $\log(LR)$  values provided by semi-continuous and fully-continuous models for F6C (Figure 2a) and GF (Figure 2b) amplification kits. The dashed line represents an hypothetical situation where all the  $\log(LR)$  results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms. Figure 3 Histograms displaying average  $\log(LR)$  values provided by semi-continuous (SC) and fully continuous (FC) models for 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified by GF (a) and F6C (b) amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. Mean  $\log(LR)$  values relative to NIST materials composing the mixtures are represented by yellow (SC) and orange (FC) histograms for NIST A contributor, light green (SC) and dark green (FC) histograms for NIST B contributor, pink (SC) and red (FC) histograms for NIST C contributor. All the corrections suggested by reviewer #1 have been made.
- References:  
All the corrections suggested by reviewer #1 have been made.

## Reviewer #2:

- The paper is interesting and it is useful to carry out comparative studies like this. However, the style of the paper and grammar are poor and needs improvement. It is quite difficult to follow. There are no line numbers or page numbers which makes referencing difficult for reviewers. According to the comment of reviewer #2 and the suggestions of reviewer #1, the grammar has been riedited and, hopefully, improved. Line and page numbers have been added, too.
- No model is “fully continuous” – because not all features are taken account of by any model. Alternative terms “qualitative” and “quantitative” models are adopted by ENFSI BPM, but terms like discrete are also acceptable. <http://enfsi.eu/wp-content/uploads/2017/09/Best-Practice-Manual-for-the-internal-validation-of-probabilistic-software-to-undertake-DNA-mixture-interpretation-v1.docx.pdf>. Due to the fact that the terms “fully continuous” are largely reported in literature (e.g. from the paper “H. Kelly, J. Bright, J. Buckleton, J. Curran, A comparison of statistical models for the analysis of complex forensic DNA profiles, Sci. Justice. 54 (2014) 66–70.

doi:10.1016/j.scijus.2013.07.003.”), the authors preferred to use this way of describing the difference between the employed probabilistic approaches for DNA mixture analysis. In case also the editor would suggest to modify it, the terms “fully continuous” will be promptly replaced by a more rigorous “quantitative” model, as correctly remarked by reviewer #2.

- *Use Hp Hd rather than H(p) H(d)*  
The correction suggested by reviewer #2 has been made.
- *“several open-source software available in literature” Euroformix is open source.*  
No corrections have been made due to the fact that, in our opinion, further open source software exist like, for instance, LikeLTD and DNAmixtures. However, in case also the editor would suggest to modify this sentence, it will be promptly modified as suggested by reviewer #2.
- *“it gradually turned to be not suitable since it does not take into account neither the instrumental stochastic effects” Reference ISFG DNA commission docs.*  
No further references have been added due to the fact that, in our opinion, several papers have been already cited. However, in case also the editor would suggest to modify this reference, it will be promptly added as suggested by reviewer #2.
- *“fully-continuous approaches, that properly include peak heights’ values into their algorithms, are supposed to be the most powerful methods” Who supposes this – not a scientific statement??*  
In order to fulfil the correct suggestion made by reviewer #2, the previous sentence was modified, as follows: “... are supposed to be more complex methods.”
- *Note 19:1 mixtures are quite extreme and minor contributors approach the limits of interpretation.*  
19:1 mixtures have been analysed to evaluate the behaviour of the different probabilistic approach when dealing with complex and unbalanced DNA mixtures, as remarked by reviewer #2, too.
- *“slightly higher within-software differences” Do you mean between quantitative continuous software? 3 orders of magnitude? Is this a range between software across all results?*  
The different fully-continuous models provided different log(LR) values when compared one another, showing average differences of 3 orders of magnitude. However, these differences were not observed for all the tested samples and calculations, as “3 orders of magnitudes” is an indicative value. In order to better explain this concept, the sentence has been modified, as follows: “ (i.e. approximatively around 3-4 degrees of magnitude, on average)”.
- *You expect that the differences between software will be greater in terms of absolute magnitude when LRs are very higher. I would like to see these divergences compared to the average LR across quantitative and qualitative models. Also if all software return LRs > 1bn then there is no practical impact if the software are divergent. Divergence is more important*

when the LRs are low. In fig 1 legend it isn't clear what mixture proportion the NIST A,B,C refer to. For example does the 20:9:1 mixture refer to A:B:C?

The differences between software and the average LR across quantitative and qualitative models have been already shown in the different figures, including the ones reported in the Supplementary Material. In our opinion, the similar trends have been observed when comparing the SC and the FC models, since the  $\log(\text{LR})$  values provided by the FC software were always higher than the ones from the SC models, for all the tested mixtures. Similar results were observed when dealing with the serially scalarly-diluted 1:1:1 DNA mixtures but, as it is shown in both Figure 3 and S6, the average  $\log(\text{LR})$  values from the FC models were always higher than the ones from the SC probabilistic approach. No further graphs have been added but if also the editor would suggest to prepare them, they will be promptly added as suggested by reviewer #2 within the Supplementary Material. Finally, as it is already reported in Table 1, the main contributor of the 3-person DNA mixtures is always NIST A, followed by NIST B and, then, NIST C.

- *“Similar increase in the  $\log(\text{LR})$  results from fully-continuous approach is observed, ranging from approximately 1.4 times – for F6C – up to 1.8 times – for GF – in terms of  $\log(\text{LR})$  values, with respect to the ones provided by semi-continuous calculations.” I don't see how this follows from fig 2 – please make this clearer. Surely it depends on the magnitude of the LR? Figure 2 shows the average  $\log(\text{LR})$  values from both SC and FC models for different DNA amplification kits. The values 1.4 and 1.8 are indicative, since they represent the slopes of the regression lines obtained when comparing the average  $\log(\text{LR})$  values of the SC models with the ones from the FC calculations. The dashed line represents the situation where all  $\text{LR}_{\text{SC}}$  are equal to  $\text{LR}_{\text{FC}}$  according to the corresponding couple of hypotheses (not the real situation). Conversely, the solid line (with intercept equal to zero) represents the average difference (or trend) between the calculated average log-likelihood ratio results provided by the two models. As a consequence, the slope is only indicative but it shows that FC models provided higher values than the SC models.*
- *“In practice, SC results turned equal to -3.74 compared to its corresponding FC  $\log(\text{LR})$  value of 1.98 for NIST A and -4.73 (compared to its corresponding FC  $\log(\text{LR})$  value of 1.98) for NIST C.” The authors need to consider false positive rates for the models used. A  $\log_{10} \text{LR}=1.98$  would usually not be sufficient to report. I don't understand “Despite SC results delivered a very strong support to  $H(d)$ , the DNA mixture under examination properly included both the contributors, as identified by FC results providing a moderate strong support to  $H(p)$ .” What criteria are used to define strong support etc?  $\log_{10}\text{LR}=1.98$  is weak surely.*

The aim of this study is not to show the consistency or the prominence of the FC models on the SC ones, but the authors want to highlight the fact the discordant LR results can be obtained, especially in case of extreme LT-DNA mixtures. The authors are aware that false positive rates should be taken into account and that a  $\log_{10}(\text{LR})$  of 1.98 is not sufficient to be reported as an inclusion, but our aim was to show such different behaviour of the tested models. Furthermore, according to reviewer #2's comment, the sentence “providing a moderate strong support...” was correctly modified, as follows: “providing a moderate support...”.



- “These evaluations suggested that fully-continuous approaches should be adopted in case of LT-DNA interpretation.” No I don’t see the justification for this statement. Just because you get a bigger LR does not mean to say it is correct or preferable. This is because there is not evaluation of the effect of false positive results by carrying out non-contributor analysis which is recommended by most providers of software.

In our opinion, the comment made by reviewer #2 is undoubtedly correct. As a consequence, non-contributor tests were performed for all the prepared DNA mixtures. Non-contributor tests were performed for NIST A, NIST B and NIST C subjects with both LRmix Studio and EuroForMix software. The results obtained by LRmix Studio are reported in terms of boxplots in the Supplementary Material with the Figures S9a, S9b and S9c for the non-contributor tests of the subjects NIST A, NIST B and NIST C, respectively. The results provided by EuroForMix are not reported since they turned totally similar to the ones provided by LRmix Studio. Furthermore, the statement remarked by the reviewer #2 was modified, too, as follows: “These evaluations suggested that fully-continuous approaches might provide bigger LR values in case of LT-DNA interpretation”.

- “always supported the prosecution hypothesis, which was later properly verified by the investigating authorities.” With casework you can’t be sure of the ground truth so you have to be very cautious with statements like this.

The cited sentence has been removed as suggested by both the reviewers.

- “Although the authors seek to make a comparative study of qualitative versus quantitative models, they omit a previous study of Bleka, Øyvind, et al. Forensic Science International: Genetics 25 (2016): 85-96 which does the same thing. Here the authors conclude "However, the main benefit of EuroForMix was with the interpretation of major/minor mixtures where the minor was evidential. Here up to 11 allele dropouts for the POI in a three-person mixture could provide probative evidence, whilst LRmix may return a much lower LR or a false negative result. The two models are expected to return similar LR results when contributors have equal mixture proportions or for mixtures of higher order"

As it was correctly remarked by the reviewer #2, the cited reference was added within the text.

- The study is quite limited in that it only examines  $H_p: S+U$  vs  $H_d: U+U$ . What is the effect of conditioning with propositions like  $H_p: S+V$  vs  $H_d: V+U$ ?

In the present study, the effect of conditioning with propositions like  $H_p: S+V$  vs  $H_d: V+U$  was not evaluated. Further studies have to be performed in order to discuss such effect, too. However, the main aim of this study deals with the evaluation of the behaviour (in terms of proof-of-concept) of SC and FC models according to different 2-person and 3-person DNA mixtures.

- STRmix uses MCMC which means that there will be variation of LR between different runs. What is this variation and how would it impact fig 1 in comparison with the exact methods used?

Due to the fact that STRmix exploits MCMC methodologies, slight differences were observed in terms of LR values between consecutive STRmix runs, as correctly remarked by reviewer

#2. However, due to the fact that no significant differences were observed when STRmix calculations were randomly repeated (i.e. maximum difference lower than one order of magnitude), and due to the fact that a large number of probabilistic calculations were made, a non-comprehensive reply can be expressed for this comment. Nevertheless, the scope of the study is focused on the “proof-of-concept” relative to the evaluation of the behaviour (in terms of proof-of-concept) of SC and FC models according to different 2-person and 3-person DNA mixtures. More comprehensive validation studies dealing with STRmix approach already available in literature.

- *It isnt clear which Euroformix method is used? Bayesian? MLE? Conservative approach? All will give different results.*

The “Continuous LR” (i.e. MLE based) methodology was employed when using EuroForMix. This information has been added within the text.

- *Remember that when assessing whether a result is good or not, does not depend upon a model giving a bigger number. We are more interested in the number of times the model returns false positive and false negative results. The authors dont address this - it would be interesting to see results of non-contributor tests.*

As it was mentioned before, non-contributor tests were performed for all the prepared mixtures.

- *The problem with casework analysis is that you dont know the ground truth for certain - therefore you dont know if the model is behaving correctly. The grammar in the paper needs improvement.*

As it was mentioned before, the sentence dealing the final conclusion of the caseworks were removed and, hopefully, the grammar was edited.



- Semi- and fully-continuous models to DNA mixtures interpretation are investigated
- 2- and 3-person *ad hoc* DNA mixtures analysed by multiple STR amplification kits
- Lab Retriever, LRmix Studio, DNA•VIEW<sup>®</sup>, EuroForMix and STRmix<sup>™</sup> software were used
- LR values from fully-continuous software turned to be the highest

**Title: DNA mixtures interpretation – a proof-of-concept multi-software comparison highlighting different probabilistic methods’ performances on challenging samples.**

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1 **Title: DNA mixtures interpretation – a proof-of-concept multi-software**  
2 **comparison highlighting different probabilistic methods’ performances on**  
3 **challenging samples.**

4  
5  
6 **Abstract:**

7 The present study investigated the capabilities and performances of semi-continuous and fully-  
8 continuous probabilistic approaches to DNA mixtures interpretation, particularly when dealing with  
9 Low-Template DNA mixtures. Five statistical interpretation software, such as Lab Retriever and  
10 LRmix Studio – involving semi-continuous algorithms – and DNA•VIEW<sup>®</sup>, EuroForMix and STRmix<sup>™</sup>  
11 – employing fully-continuous formulas – were employed to calculate likelihood ratio, comparing the  
12 prosecution and the defense hypotheses relative to a series of on-purpose prepared DNA mixtures  
13 that respectively contained 2 and 3 known contributors. National Institute of Standards and  
14 Technologies (NIST) certified templates were used for samples set up, which contained different  
15 DNA amounts for each contributor. 2-person mixtures have been prepared with proportions equal  
16 to 1:1, 19:1 and 1:19 in terms of DNA concentration. Conversely, three person mixtures were  
17 constituted by proportions equal to 20:9:1, 8:1:1, 6:3:1 and 1:1:1 in terms of DNA concentration.  
18 Furthermore, 8 equally-proportioned 3-person mixtures were prepared by means of scalar dilutions  
19 starting from an overall amount of 0.500 ng, then ranging up to DNA samples with concentrations  
20 equal to 0.004 ng (i.e. Low-Template DNA). DNA mixtures were set up in triplicate and amplified  
21 with 7 DNA amplification kits (i.e. GlobalFiler PCR Amplification Kit, NGM SElect PCR Amplification  
22 Kit, MiniFiler PCR Amplification Kit, Power Plex Fusion, PowerPlex 6C Matrix System, Power Plex ES1  
23 17 Fast and Power Plex ESX 17 Fast) in order to evaluate whether the selection of a certain kit might  
24 represent a bias factor, capable of altering the whole interpretation process. Multi-software  
25 approach helped us to highlight any trend in the likelihood ratio results provided by semi- and fully-  
26 continuous software. As a matter of fact, fully-continuous computations provided different (higher)  
27 results in terms of degrees of magnitude of the likelihood ratio values with respect to the ones from  
28 the semi-continuous approach, regardless of the amplification kit that was utilized.

29  
30 **Keywords:** DNA mixture interpretation; Low-Template DNA; semi-continuous model; fully-  
31 continuous model; likelihood ratio.

32

33

## 34 1. Introduction

35 The probabilistic interpretation of DNA evidences recovered on crime scenes has been for many  
36 years a largely debated and investigated issue in the field of forensic biology [1–8], especially in case  
37 of Low-Template DNA (LT-DNA) samples. There are several questions and multiple choices, indeed,  
38 that the analysts have to face when bio-statistical interpretation processes on DNA mixtures take  
39 place, such as: (i) the calculation of analytical and stochastic thresholds [9–11], (ii) the evaluation of  
40 the most probable number of contributors composing the DNA samples [12–17], (iii) the  
41 establishment of parameters such as drop-in rate, drop-out rate [6,9,18–23], stutters rate [24,25]  
42 and co-ancestry coefficient ( $F_{st}$ ) [26,27], (iv) the selection of the appropriate allele frequency  
43 dataset [28]. Moreover, ~~the use of the appropriate model to be adopted on the acquired data to~~  
44 ~~perform bio-statistical interpretations is essential~~ bio-statistical interpretation, with the most  
45 accurate model, is essential. Briefly, there are three interpretative approaches that can be used in  
46 the sphere of DNA mixtures interpretation, which differ one another in terms of complexity,  
47 according to the input and the algorithms they take into account [29]. These methodologies are  
48 widely known as (i) binary, (ii) semi-continuous and (iii) fully-continuous approaches. The Bbinary  
49 model [30] was the first one employed by the forensic community for DNA mixture interpretation  
50 but, over the time, it was gradually ~~turned-replaced. This method to be not suitable since it~~ does  
51 not take into account neither the instrumental stochastic effects (i.e. drop-in and drop-out –  
52 evaluated by the semi-continuous models), nor the peak heights of the detected alleles (evaluated  
53 by the fully-continuous models). Due to developments of more sensitive instruments and high-  
54 throughput analyses ~~[30,31][31,32]~~, capable of evaluating very low concentrations of DNA too,  
55 analysts had to deal with bigger-larger amounts of data and parameters that could not be managed  
56 by the traditional binary approach. For this reasons, semi-continuous ~~[32,3333-35]~~ and fully  
57 continuous models ~~[34,35][36,37]~~ have largely replaced by far the binary one, and nowadays they  
58 represent the gold-standard methodologies to be adopted when DNA mixture interpretations occur.  
59 However, the selection of a specific interpretation approach is not trivial, but it can be related to  
60 several factors such as, for instance, the degree of expertise that the analysts are endowed with,  
61 together with the resources of the forensic laboratories themselves. In fact, several software  
62 performing fully-continuous models are neither free of charge nor open-source, and the  
63 laboratories necessarily have to buy a license in order to use them on their own data. For this reason,  
64 in the previous years semi-continuous models have been largely employed by forensic analysts.

65 ~~since~~ there are actually several open-source software available in literature such as, for instance,  
66 Lab Retriever ~~[33][34]~~, and LRmix Studio (previously, LRmix ~~[36][38]~~), which have been used for this  
67 study, or LikeLTD [35,39]. Another reason supporting the widespread of semi-continuous models is  
68 the fact that their algorithms and computations are more straightforward than the ones performed  
69 by fully-continuous approaches. Furthermore, their workings and results may be more easily shown  
70 and discussed in courtrooms. However, semi-continuous approach does not take into account the  
71 information regarding alleles' peak heights. As a consequence, fully-continuous approaches, that  
72 properly include peak heights' values into their algorithms, are supposed to be ~~the most~~  
73 powerful more complex methods since they exploit the whole available information included within  
74 the acquired data. Then, fully-continuous approaches might represent a desirable solution in case  
75 of complex DNA mixture involving multiple contributors and LT-DNA. However, analysts should be  
76 well-trained before employing these methodologies in order to avoid misinterpretations or wrong  
77 applications of such algorithms. Due to these problems, our laboratory adopted a "*statistic*  
78 *consensus approach*" ~~[37][40]~~, which seems to solve several issues that traditionally arise in case of  
79 LT-DNA mixture interpretation. This approach simply compares likelihood ratio (LR) results provided  
80 by different probabilistic software, reporting only the most conservative LR value (and its correlated  
81 verbal statement ~~[38][41]~~) if coherence among the tested models is observed. Otherwise,  
82 inconclusive decisions is taken into account. Even though the behaviours of semi- and fully-  
83 continuous models have been already compared in other studies ~~[29,39-41][29,42-44]~~,  
84 nevertheless no comprehensive guidelines have been drafted yet, describing an overall-accepted  
85 approach to be employed when dealing with complex DNA mixtures, especially in case of LT-DNA.  
86 In order to evaluate the different performances and outputs of semi- and fully-continuous models,  
87 together with the need of furtherly validating our developed "*statistic consensus approach*", several  
88 mixtures DNA mixtures of two and three known contributors mixed in different proportions were  
89 prepared, were ad hoc prepared and composed by two and three known contributors mixed in  
90 different proportions, then analyzed using 7 DNA amplification kits. 2 semi-continuous (Lab  
91 Retriever, LRmix Studio) and 3 fully-continuous (DNA•VIEW® ~~[42][45]~~, EuroForMix ~~[7,35][7,37]~~,  
92 STRmix™ ~~[24,43][24,46]~~) software were employed. The provided LR results were compared, with  
93 respect to the utilized DNA amplification kit, aiming to perform a wide comparison of the cited semi-  
94 continuous and the fully-continuous software. ~~Thanks to t~~ This experimental plan, ~~it had been~~  
95 possible to even evaluate allowed the assessment of whether ~~whether~~ the performances of the  
96 examined software turned to be influenced or not by the adopted DNA amplification kits.

97 Furthermore, several scalar dilutions in terms of DNA amount were prepared involving a known and  
98 equally-proportioned 3-person mixture. Once again, such diluted mixtures were ~~analysed~~analyzed  
99 with several DNA amplification kits aiming to evaluate the questioned probabilistic models with  
100 respect to decreasing concentrations of DNA (i.e. ~~up~~down to proper LT-DNA samples). An LT-DNA  
101 real casework is also discussed, ~~too~~ to test our approach.

102

## 103 2. Materials and methods

### 104 2.1. Sample preparation and analysis

105 DNA samples were all set up with Standard Reference Material® 2391c, primarily intended for use  
106 in the standardization of forensic QA (Quality Assurance) and paternity test procedures for PCR-  
107 based genetic testing. ~~NIST A, NIST B and NIST C~~ PCR-based DNA profiling standard NIST® SRM®  
108 reference samples 2391c (NIST) were selected for this experiment, and used as known contributors  
109 for 2-person and 3-person mixtures, as shown in Table 1. The DNA profile from NIST F reference  
110 sample was used, conversely, as a known non contributor in order to perform false donor tests for  
111 all the tested software, too. Different DNA proportions of NIST A, NIST B and NIST C were evaluated  
112 for both the ~~ad hoc~~-prepared 2- and 3-person mixtures containing known NIST contributors (mixture  
113 ratios are reported in Table 1). In particular, 2-person mixtures were prepared with contributors'  
114 proportion ratio of 19:1, 1:1 and 1:19, while 3-person mixtures included NIST samples mixed with  
115 the proportions ratios 20:9:1, 8:1:1, 6:3:1, 1:1:1. All the prepared DNA mixtures had an approximate  
116 concentration of 0.500 ng. ~~Moreover, starting from such quantitative, a~~ An equally-proportioned  
117 1:1:1 mixture (0.500 ng) containing NIST samples A, B and C was scalarly diluted in order to obtain  
118 several further samples (i.e. with lower concentration levels), such as 0.250 ng, 0.125 ng, 0.063 ng,  
119 0.031 ng, 0.016 ng, 0.008 ng and 0.004 ng. All the samples were repeatedly amplified using the  
120 following 7 DNA amplification kits: GlobalFiler™ PCR Amplification Kit (GF), AmpFISTR® NGM Select™  
121 Amplification Kit (NGM), AmpFISTR® MiniFiler™ PCR Amplification Kit (MF) - from Thermo-Scientific,  
122 (Waltham, MA, USA) - and PowerPlex® Fusion System (F), PowerPlex® Fusion 6C System (F6C),  
123 PowerPlex® ESI 17 Fast System (ESI), PowerPlex® ESX 17 Fast System (ESX) - from Promega  
124 Corporation (Madison, WI, USA). Allele detection was performed by capillary electrophoresis (CE)  
125 on Applied Biosystems® 3500 Series Genetic Analyzer with a 36 cm 3500 Genetic Analyzer Capillary  
126 Array and POP-4™ Polymer 3500 Genetic Analyzer (Thermo Fisher Scientific) together with an  
127 Injection standard protocol 1.2 kV/15 sec.

128 The whole analytical methodology was internally validated following UNI CEI EN ISO/IEC 17025 and  
129 Scientific Working Group on DNA Analysis Methods (SWGAM - <http://www.swgdam.org/>)  
130 requirements. Parameters such as accuracy, linearity, quantification accuracy, limit of detection  
131 (analytical threshold), limit of quantitation, mixtures deconvolution, repeatability, concordance and  
132 repeatability limit, robustness, sensitivity, decision threshold, direct amplification inhibition,  
133 stochastic threshold, specificity, species specificity, uncertainty, stutters, drop-in and drop-out were  
134 validated proving satisfactory results (not reported in the study) for all the employed DNA  
135 amplification kits. Analytical methodologies, together with the mixture interpretation approach  
136 reported in [37][40], were verified and accredited by ACCREDIA, the Italian body appointed for the  
137 accreditation of analytical protocols and methodologies in laboratories.

138  
139

## 140 **2.2. Software and LR calculations**

141 GeneMapper® ID-X v1.4 from Thermo Fisher Scientific (Waltham, MA, USA), OSIRIS v2.7 (from  
142 <http://www.ncbi.nlm.nih.gov/projects/SNP/osiris/>) and ArmedXpert™ v3.0.7.999 from NicheVision  
143 Forensics LLC (Akron, OH, USA) were employed to manage the acquired raw data and filter them  
144 by applying the differential analytical thresholds calculated (i.e. specific validated analytical  
145 thresholds were observed for each dye channel). Then, data were modified on MS Excel in order to  
146 obtain suitable input formats for each bio-statistical software employed in this study, which were  
147 as follows: Lab Retriever, LRmix Studio, DNA•VIEW®, EuroForMix and STRmix™.

148 Lab Retriever ~~and LRmix Studio are-is-an~~ open-source and free of charge software performing a  
149 semi-continuous approach to DNA mixtures, ~~as well as LRmix Studio~~. The first (Lab Retriever, version  
150 2.2.1) was downloaded from the Scientific Collaboration, Innovation and Education (SCIEG) website  
151 ([http://scieg.org/lab\\_retriever.html](http://scieg.org/lab_retriever.html)), developed by K. Inman, K. Lohmueller and N. Rudin. The  
152 second, LRmix Studio (version 2.1.3), is available on the website <http://www.lrmixstudio.org/>,  
153 developed by H. Haned and P. Gill. On the other hand, DNA•VIEW® is a commercial software  
154 involving a fully-continuous algorithm that takes into account peak height (in terms of Relative  
155 Fluorescent Units, RFU) of each detected allele. This software was developed by C.H. Brenner  
156 (<http://dna-view.com/>) and version 37.17 was employed to perform LR calculations based mainly  
157 on stochastic variation, incorporating dropout, drop-in, stutter and allelic stacking naturally, without  
158 the use of Markov chain Monte Carlo (MCMC) methods. Moreover, EuroForMix is an open-source  
159 and free of charge software involving a fully-continuous approach for DNA mixtures. It is one of the



160 first fully-continuous open-source software to be available on internet (version 1.9.3 was  
161 employed), programmed by Ø. Bleka and working in R [47] [44] environment (package “euroformix”,  
162 R version 3.4.3 was used). In particular, it involves maximization (frequentistic) or integration  
163 (Bayesian) approaches over the likelihood function of a gamma peak height model for STR/SNP DNA  
164 data (as remarked on <http://www.euroformix.com/>). The “Continuous LR” mode (i.e. involving a  
165 Maximum Likelihood Estimation - MLE - methodology) was used. Finally, STRmix™  
166 (<http://strmix.esr.cri.nz/>) is a commercial software involving Markov Chain Monte Carlo (MCMC)  
167 approach for fully-continuous DNA mixtures interpretation [29,41,45][29,44,48], developed by D.  
168 Taylor, J.A. Bright and J. Buckleton. J. Buckleton and his research group.

169 A free-trial version (version 2.3.06) of STRmix™ was employed in this study.

170 All the described software were exploited with the aim of calculating used LR values relative to each  
171 one of NIST samples used as known contributor and included into our prepared DNA mixtures.

172 In case of 2-person mixtures, LR calculations were performed using the following hypotheses:

- 173 • Prosecution hypothesis  $H_p = \text{Subject } X_i + 1 \text{ unknown individual}$ ;
- 174 • Defence hypothesis  $H(d)d = 2 \text{ unknown individuals}$ .

175 where  $X_i$  stands for the  $i$ -th contributor (NIST A, NIST B or NIST C) included into the mixtures under  
176 exam.

177 Conversely, dealing with 3-person mixtures, the following hypotheses were evaluated for the LR  
178 calculation:

- 179 • Prosecution hypothesis  $H(p)p = \text{Subject } X_i + 2 \text{ unknown individuals}$ ;
- 180 • Defence hypothesis  $H(d)d = 3 \text{ unknown individuals}$ .

181 A validated and accredited overall drop-in value equal to 0.05, together with a  $F_{st}$  (theta) value of  
182 0.01, were set in all probabilistic software. Different drop-out values, according to estimation of  
183 drop-out probability range performed by LRmix Studio [36] [38], were set into both Lab Retriever  
184 and LRmix Studio, but only the most conservative LR value was recorded [21]. Finally, NIST U.S.  
185 population dataset was adopted as reference database in all LR computations [46][49].  
186

### 187 3. Results and discussion

#### 188 3.1. LR comparison of 2-person and 3-person mixtures

189 LR results relative to 2-person and 3-person mixtures amplified with Fusion 6C (F6C) and GlobalFiler  
190 (GF) amplification kits are described in this section. For these two amplification kits, 2-person DNA

191 mixtures included individuals/certified materials labelled as NIST B and NIST C, which were mixed  
192 together in different proportions (i.e. 1:1, 1:19 and 19:1). Then, NIST A, NIST B and NIST C were used  
193 to compose four different 3-person mixtures (i.e. 20:9:1, 8:1:1, 6:3:1 and 1:1:1). In this case, NIST C  
194 was used as major contributor of DNA mixtures with proportions-ratios of 20:9:1, 8:1:1 and 6:3:1.  
195 Moreover, NIST B certified material represented the second major contributor in the DNA mixtures  
196 with the proportions 20:9:1 and 6:3:1, while NIST A material simulated the minor contributor.

197 LR results of the different mixtures were evaluated comparing the prosecution hypothesis  $H(p)p$ ,  
198 which included the person of interest – POI, i.e. NIST A, NIST B or NIST –, versus the defence  
199 hypothesis  $H(d)d$ , that did not include the POI, but contained unknown individuals only. Results are  
200 graphically summarized in Figure 1. In particular, base-10 logarithms were applied so that  $\log(LR)$   
201 results are reported on the y axis, while codes indicating the DNA mixtures and their relative  
202 proportions are displayed on the x axis (NIST A contributor is shown in Figure 1a-b, NIST B in Figure  
203 1c-d and NIST C in Figure 1e-f). As it can be seen, LR results provided by both semi-continuous  
204 models turned-were very similar or identical. This is due to the fact that Lab Retriever and LRmix  
205 Studio software utilizes similar algorithms, with slightly divergent formulas—[40][43,44].  
206 Furthermore,  $\log(LR)$  results provided by fully-continuous models proved similar and convergent to  
207 one another, with slightly higher within-software differences (i.e. up-approximately around to-3-4  
208 degrees of magnitude, on average), if compared to semi-continuous software.

209 Since  $\log(LR)$  results turned convergent-similar among the tested approaches, mean  $\log(LR)$  values  
210 were calculated for both semi-continuous ( $LR_{SC}$ ) and fully-continuous ( $LR_{FC}$ ) algorithms.  $LR_{SC}$  and  $LR_{FC}$   
211 results are reported in Figure 2 for F6C (Figure 2a) and GF (Figure 2b) amplification kits. As it can be  
212 seen,  $LR_{FC}$  results turned-were always higher than the ones provided by fully-continuous modelling  
213 ( $LR_{SC}$ ) for both F6C and GF. The dashed line represents ana hypothetical situation where all  $LR_{SC}$  are  
214 equal to  $LR_{FC}$  according to the corresponding couple of hypotheses. Conversely, the solid line (with  
215 intercept equal to zero) represents the average difference (or trend) between the calculated  
216 average log-likelihood ratio results provided by the two models. Similar increases in the  $\log(LR)$   
217 results from fully-continuous approach is-were observed, ranging from approximately 1.4 times –  
218 for F6C – up to 1.8 times – for GF – in terms of  $\log(LR)$  values, with respect to the ones provided by  
219 semi-continuous calculations.

220 As well as F6C and GF kits, similar trends of  $\log(LR)$  results were observed for all the other validated  
221 DNA amplification kits (i.e. NGM, MF, F, ESI and ESX). Results relative to these kits, together with  
222 their average  $\log(LR)$  values (i.e.  $LR_{SC}$  and  $LR_{FC}$ ), are reported in Figures S1-S5 in the Supplementary

223 Material. Furthermore, false donor test were calculated for all the tested mixtures and software by  
224 involving NIST F reference profiles. The obtained results are reported in Figure S6 in the  
225 Supplementary Material. As it can be seen, the log(LR) values provided by semi-continuous software  
226 were always lower (in terms of absolute values) than the ones provided by fully-continuous  
227 approaches.

228

### 229 3.2. LR variation ~~with respect to total DNA concentration~~ related to DNA 230 quantity

231 Eight equally-proportioned 3-person mixtures were prepared starting from an overall DNA  
232 concentration of 0.500 ng. DNA samples showing a concentration of 0.500ng, 0.250 ng, 0.125 ng,  
233 0.063 ng, 0.031 ng, 0.016 ng, 0.008 ng and 0.004 ng were prepared including NIST A, B and C certified  
234 materials as known contributors. Once again, prosecution hypotheses  $H(p)_p$  including the POI were  
235 compared to defence hypotheses  $H(d)_d$  involving only unknown individuals. Base-10 logarithms  
236 were evaluated for all contributors by means of the same panel of software. Then, average log(LR)  
237 values were calculated aiming to compare the results provided by semi-continuous software (SC)  
238 with the ones provided by fully-continuous approaches (FC). As it can be seen in Figure 3a-b for GF  
239 and F6C amplification kits, respectively, log(LR) data are reported on y axis, according to employed  
240 algorithm (SC or FC), while the codes indicating the concentrations of the 1:1:1 DNA mixtures are  
241 indicated on the x axis. In particular, log(LR) results seemed to increase in correlation with the DNA  
242 concentrations of the scalarly-diluted mixtures under examination. Similar behaviours were  
243 observed for all known NIST contributors and both GF and F6C amplification kits. However, the FC  
244 results turned always higher than the SC ones, regardless of the DNA amplification kit that was  
245 adopted. This observation proved undoubtedly remarkable in case of the DNA mixtures that present  
246 very low DNA concentrations (i.e. 0.004 ng, 0.008 ng and 0.016 ng – LT-DNA). As it can be seen in  
247 Figure 3a, 0.004 ng mixture amplified using GF kit showed log(LR) values lower than zero – thus  
248 supporting the exclusionary hypothesis – when SC approach was performed involving NIST A or NIST  
249 C as POI. Otherwise, FC models provided log (LR) results higher than zero – thus supporting the  
250 inclusionary hypothesis -. In practice, SC results turned equal to -3.74 (compared to its  
251 corresponding FC log(LR) value of 1.98) for NIST A and -4.73 (compared to its corresponding FC  
252 log(LR) value of 1.98) for NIST C. Despite SC results delivered a very strong support to  $H(d)_d$ , the  
253 DNA mixture under examination properly included both the contributors, as identified by FC results

254 providing a moderate ~~strong~~ support to  $H(p)p$ . These evaluations suggested that fully-continuous  
255 approaches ~~should be adopted~~ might provide bigger log(LR) values in case of LT-DNA interpretation.  
256 This statement was corroborated by further results, such as SC log(LR) value for NIST B contributor  
257 which turned lower (i.e. 0.84, scarcely supporting  $H(p)p$ ) than the one provided by FC approach (i.e.  
258 1.78, moderately supporting  $H(d)d$ ). Similar performances were observed in the cases of 0.008 ng  
259 and 0.016 ng DNA mixtures, where SC log(LR) values relative to NIST C contributor turned lower  
260 than zero, and SC log(LR) results relative to NIST A contributor turned lower (0.008 ng) or very close  
261 (0.016 ng) to zero. Moreover, log(LR) results for both SC and FC models proved always higher than  
262 zero starting from a DNA concentration of 0.031 ng, but FC values always showed higher results  
263 than those from SC approaches. Similar behaviours were observed evaluating the DNA mixtures  
264 amplified by F6C kit (Figure 3b), too. In this case, probabilistic interpretation of 0.004 ng DNA  
265 mixture provided log(LR) values lower than zero for both the biostatistical models, with the  
266 exception of FC log(LR) result relative to NIST B known contributor equal to 1.9. Moreover, a log(LR)  
267 value of -1.47 was observed for NIST C contributor in the mixture containing 0.008 ng of DNA. Once  
268 again, these observations proved a better sensitivity of the fully-continuous models in case of LT-  
269 DNA. However, the number of unmasked alleles of the trace contributors left above the calculated  
270 AT for the different DNA amplification kits were around 5-20%. Different percentage values were  
271 obtained according to the DNA amplification kit under evaluation and, therefore, inconclusive  
272 conclusion or inappropriate LR values were expected for these LT-DNA mixtures. As well as F6C and  
273 GF kits, similar performance were observed for all the DNA mixtures amplified by the other validated  
274 DNA amplification kits (i.e. NGM, MF, F, ESI and ESX). The results relative to these kits are depicted  
275 in Figure S67 in the Supplementary Material. Furthermore, false donor tests were calculated for all  
276 the tested mixtures and software by involving NIST F reference profiles. The obtained results are  
277 reported in Figure S8 in the Supplementary Material. As it can be seen, the log(LR) values provided  
278 by semi-continuous software were, on average, similar or even higher (in terms of absolute values)  
279 than the ones provided by fully-continuous approaches when dealing with the mixture containing  
280 0.004, 0.008, 0.016, 0.31 and 0.063 ng of DNA. Then, for DNA amounts equal to 0.125 ng or higher,  
281 the log(LR) values from the false donor calculations provided by semi-continuous software were  
282 lower (in terms of absolute values) than the ones provided by fully-continuous approaches. Non-  
283 contributor tests were also performed for all the prepared DNA mixtures. Non-contributor tests  
284 were performed for NIST A, NIST B and NIST C subjects with both LRmix Studio and EuroForMix  
285 software. The results obtained by LRmix Studio are reported in terms of boxplots in the

286 Supplementary Material with the Figures S9a, S9b and S9c for the non-contributor tests of the  
287 subjects NIST A, NIST B and NIST C, respectively. The results provided by EuroForMix are not  
288 reported since they turned totally similar to the ones provided by LRmix Studio.  
289

### 290 3.3. An intriguing real casework

291 In the present paragraph log<sub>10</sub>(LR) results relative to a forensic real casework that was evaluated in  
292 our laboratory will be discussed. In details, a Caucasian individual was charged as an alleged suspect  
293 (POI) for a series of robberies and thefts. A cap was collected on-from a crime scene by police forces  
294 during their investigating activities. Since it was supposed to belong to the alleged suspect, flocked  
295 swabs Nylon® 4N6 (purchased by COPAN ITALIA S.P.A., Brescia, Italy) were applied-used to collect  
296 material on the visor of the cap aiming to detect biological evidences. The genetic material was  
297 recovered on different spots on the visor, then extracted from the swabs, amplified and analyzed.  
298 Before performing our “*statistic consensus approach*” over acquired data, differential analytical  
299 thresholds were calculated by means of ArmedXpert™ for each dye channel, following our validated  
300 analytical protocol. Due to the fact that several algorithms might be employed in order to calculate  
301 analytical thresholds [9,10], two different series of analytical thresholds to be applied on raw data  
302 were provided by ArmedXpert™ Analytical Threshold tool. This application involves different  
303 formulas that are labelled as SD3 (i.e. taking into account 3 standard deviations of the peak heights  
304 in terms of RFU; results are displayed in the upper part of Figure 4) and Min-Max (i.e. taking into  
305 account minima and maxima peak heights in terms of RFU; results are displayed in the lower part  
306 of Figure 4). As it can be seen in Figure 4, analytical thresholds involving SD3 algorithm showed  
307 higher thresholds than the ones obtained after the application of Min-Max algorithm. Both the  
308 series of differential analytical thresholds were applied on raw data and the DNA mixtures obtained  
309 were biostatistically interpreted. A sSummary of log(LR) results are reported in Table 2. Same sets  
310 of H(ϕ)p and (Hd) hypotheses including 2 or 3 contributors were evaluated and compared according  
311 to the different algorithms used in terms of analytical thresholds (3SD or Min-Max). Then,  
312 interpretative responses were stated depending on the degree of coherence among log(LR) results  
313 or, even better, between semi- and fully-continuous models. As it can be observed in Table 2, the  
314 adoption of lower RFU values (i.e. using Min-Max algorithm) provided higher log(LR) results with  
315 respect to the ones obtained by adopting SD3 formula. More in details, uncoherent results were  
316 observedCategorically different results, i.e. differing on which side of log(LR) = 0, were observed

317 between semi- and fully-continuous interpretations when evaluating the biological evidence as a 2-  
318 person mixture (i.e.  $H(p)p = \text{POI} + 1$  unknown individual;  $H(d)d = 2$  unknown individuals). Regardless  
319 of the algorithm used to calculate analytical thresholds, both Lab Retriever and LRmix Studio  
320 provided log (LR) values lower than zero, even if very close to such value. Otherwise, DNA•VIEW®,  
321 EuroForMix and STRmix™ showed log(LR) results suggesting either moderately strong (in case of  
322 3SD algorithm) or strong (in case of Min-Max algorithm) support to be delivered to the prosecution  
323 hypothesis that indicated an inclusionary contribution to the mixture by the POI. However, even  
324 though fully-continuous software showed quite high log(LR) values, semi- and fully-continuous  
325 models proved uncoherent one another. Despite fully-continuous software provided higher  
326 absolute log(LR) values with respect to the ones from the semi-continuous models, our lab adopted,  
327 validated and accredited (via UNI CEI EN ISO/IEC 17025 requirements) the cited “statistic consensus  
328 approach” in order to cross-validate the results provided by the different software and provide a  
329 trustful and robust interpretation for real caseworks and, particularly, for complex LT-DNA mixtures.  
330 Consequently, according to our adopted “*statistic consensus approach*”, the response expressed at  
331 the end of our interpretation process was inconclusive. Furthermore On the other hand, coherent  
332 results were observed when evaluating the biological evidence as a 3-person mixture (i.e.  $H(p)p =$   
333  $\text{POI} + 2$  unknown individuals;  $H(d)d = 3$  unknown individuals). In particular, semi-continuous  
334 approaches delivered a moderately strong support to  $H(p)p$ , while fully-continuous models  
335 delivered an extremely strong support to such hypothesis. In the present case, ~~response emitted~~the  
336 result reported at the end of our interpretation process supported the prosecution hypothesis,  
337 charging the POI as an effective contributor to the biological evidence collected on the visor of the  
338 cap that was recovered on the crime scene. ~~At the end of the trial, suspect was convicted as guilty~~  
339 ~~since further evidences incriminated him.~~ In our opinion, results reported in Table 2 represent once  
340 again a proof to the concept that fully-continuous models might be more sensitive than semi-  
341 continuous ones in case of LT-DNA mixtures interpretation. In the present casework, fully-  
342 continuous log(LR) values always supported the prosecution hypothesis, ~~which was later properly~~  
343 ~~verified by the investigating authorities.~~

344



## 345 Conclusions

346 Thanks to the recent developments involving DNA extraction and typing, the interpretation of the  
347 biological evidence is gradually being more and more crucial in trials, strongly affecting the  
348 judgments expressed in courtrooms. As a consequence, extreme caution is demanded in this  
349 forensic field, especially when Low-Template DNA and complex mixtures have to be interpreted.  
350 Semi-continuous and fully-continuous approaches present different degrees of complexity in terms  
351 of comprehensibility, computation and interpretation of the outputs. Multi-software evaluations of  
352 *a priori* known 2-person and 3-person DNA mixtures prepared in our laboratory allowed us to  
353 compare the performance of both semi- and fully-continuous approaches. Log(LR) results provided  
354 by the tested fully-continuous software (i.e. DNA•VIEW<sup>®</sup>, EuroForMix and STRmix<sup>™</sup>) turned always  
355 significantly higher than the ones calculated by the employed semi-continuous software (i.e. Lab  
356 Retriever and LRmix Studio). A plausible explanation of this might be related to the different semi-  
357 and fully-continuous algorithms. In particular, fully-continuous software takes into account a larger  
358 amount of data and information (i.e. detected alleles plus their relative peak heights), so that higher  
359 LR values can be obtained when significant matches are observed between the investigated  
360 biological samples. Similar trends in log (LR) values were observed when several serially scalarly-  
361 diluted 3-person mixtures were investigated, too. These mixtures were ~~ad hoc~~ prepared in order to  
362 contain the same DNA amount for each one of the included known contributors. Nevertheless, log  
363 (LR) values provided by semi-continuous software gave false exclusionary indications to the  
364 composition of the DNA mixtures, especially in case of Low-Template DNA (i.e. mixtures showing an  
365 overall DNA concentration of 0.004 ng, 0.008 ng and 0.016 ng). ~~turned uncoherent to the~~  
366 ~~composition of the DNA mixtures, especially in case of Low-Template DNA (i.e. mixtures showing an~~  
367 ~~overall DNA concentration of 0.004 ng, 0.008 ng and 0.016 ng).~~ Furthermore, probabilistic software  
368 behaved in a similar way, regardless of the DNA amplification kits that were employed. Even though  
369 this outcome was expected, further analyses might indicate which DNA amplification kits would  
370 turned out to be the most useful in cases of Low-Template DNA. ~~Obviously, the~~ These evaluations  
371 relative to these DNA mixtures just represent ~~a proof-of-concept~~ some evidence that fully-  
372 continuous model seems to be the most suitable bio-statistical methodology to be performed by  
373 analysts when Low-Template DNA mixtures or major/minor mixtures, where the minor contributor  
374 is evidential [50], have to be interpreted from a probabilistic point-of-view. Furthermore, the fully-  
375 continuous model is more effective to determine the contributors' ratios. Despite this fact, extreme  
376 caution has to be used when interpreting LT-DNA mixture, especially when different algorithms and

377 probabilistic approaches are used and compared by forensic experts. Consequently, in our opinion,  
378 a conservative approach might be employed at the current stage in order to avoid false conclusions  
379 that could eventually lead to unwanted severe legal consequences. For this reason, due to the  
380 complexity and the relevance of the task, a statistic redundancy [51] in the calculations might be  
381 useful (i.e. like our adopted “*statistic consensus approach*”). Further studies will be performed in  
382 order to combine the likelihood ratio results provided by different software and algorithms, using  
383 different interpretation weights. Further-~~Moreover, further~~ experiments (i.e. 4- and 5-person DNA  
384 mixtures [52,53]) and their relative interpretation processes need to be performed, but in our  
385 opinion these results open the pathway towards the possibility of “weighting” LR results provided  
386 by semi- and fully-continuous models, particularly in case of LT-DNA mixtures interpretation.  
387

### 388 **Conflict of interest statement**

389 The authors of this manuscript certify that they have no affiliations with or involvement in any  
390 organization or entity with any financial interest or non-financial interest in the subject matter or  
391 materials discussed in this manuscript.  
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552

553 **Table 1** List of NIST samples used as known contributors for DNA mixtures set up. Different proportions were  
 554 evaluated both for 2- and 3-person mixtures. Referring to 2-person mixtures, different NIST reference  
 555 materials were used for samples preparation according to DNA amplification kits user guide.  
 556

**Known 2-person mixtures**

DNA Typing Kit	Reference Material	Mixtures <u>proportionratios</u> (0.500 ng)
ESI 17 Fast	B:C / male:male	
ESX 17 Fast	A:C / female:male	
Fusion	B:C / male:male	19:1
Fusion 6C	B:C / male:male	<u>8:1</u> 1:1
GlobalFiler	B:C / male:male	1:19
Mini Filer	A:C / female:male	
NGM SElect	A:C / female:male	

**Known 3-person mixtures**

DNA Typing Kit	Reference Material	Mixtures <u>proportionratios</u> (0.500 ng)
		1:1:1
All kits	A:B:C / Female:male:male	6:3:1 8:1:1 20:9:1

557

558

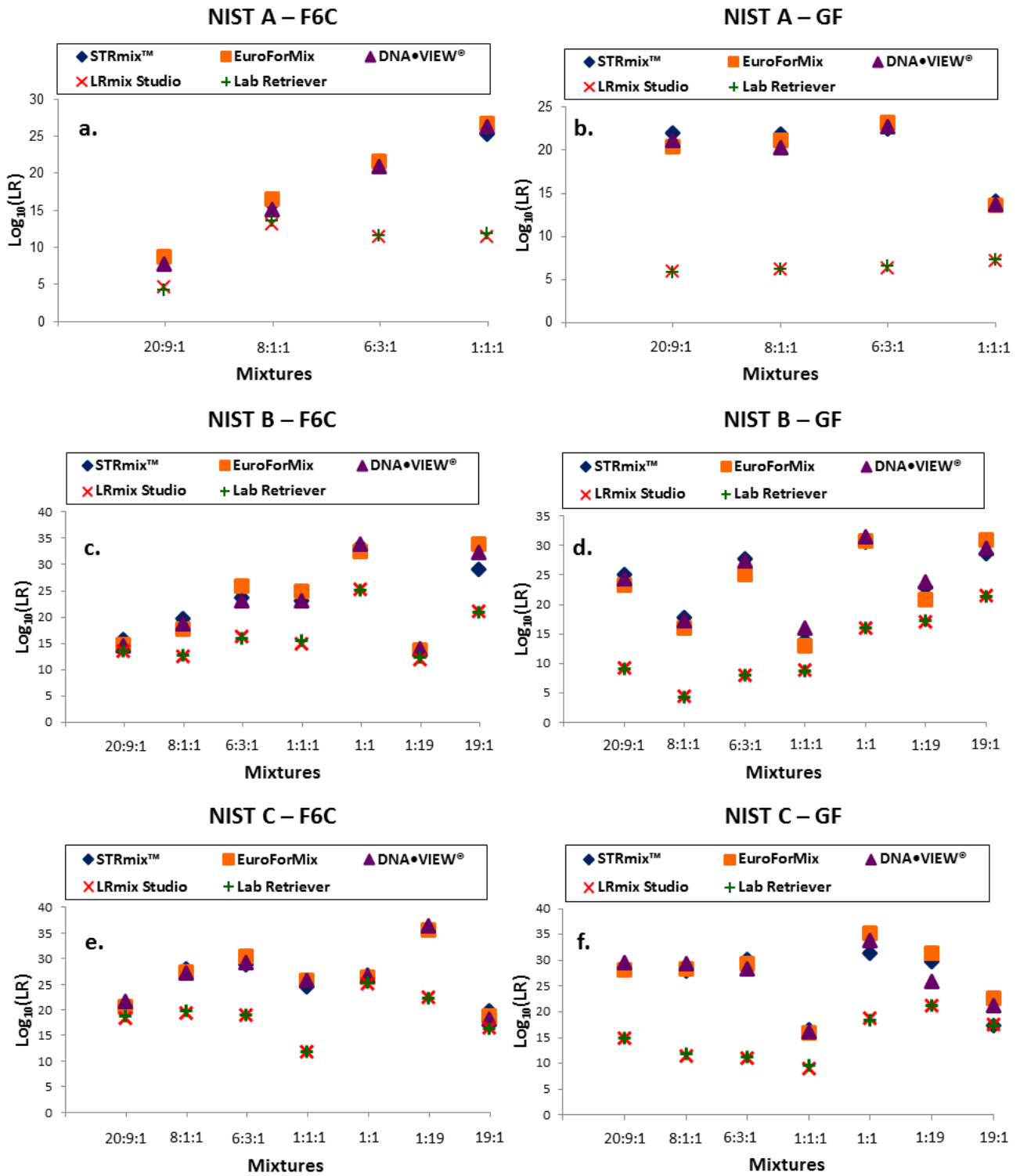
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560

561 **Table 2** Log(LR) results relative to the interpretation process performed on the DNA mixture collected on a  
 562 cap recovered on a crime scene. 3SD and Min-Max represent the algorithms that were employed to evaluate  
 563 the differential analytical thresholds. POI represents the suspect (i.e. the person of interest), while U stands  
 564 for unknown(s) individual(s) extracted from the allele frequencies reference dataset  
 565 to the interpretation process performed on the DNA mixture collected on a cap recovered on a crime scene.  
 566 3SD and Min-Max represent the algorithms that were employed to evaluate the differential analytical  
 567 thresholds. POI represents the suspect (i.e. the person of interest), while U stands for unknown(s)  
 568 individual(s) extracted from the allele frequencies reference dataset [467].

Analytical Threshold	3SD	Min-Max	3SD	Min-Max
Hp	Susp + 1U	Susp + 1U	Susp + 2U	Susp + 2U
Hd	2U	2U	3U	3U
Software	Log(LR)			
Lab Retriever	-0.36	-0.16	2.37	2.49
LRmix Studio	-0.37	-0.16	2.36	2.48
DNA•VIEW®	2.29	3.47	6.79	7.06
EuroForMix	2.33	3.60	6.25	7.12
STRmix™	2.84	3.63	7.01	7.03
Interpretative decision	Inconclusive	Inconclusive	Support to H <sub>(p)</sub> <del>p</del>	Support to H <sub>(p)</sub> <del>p</del>

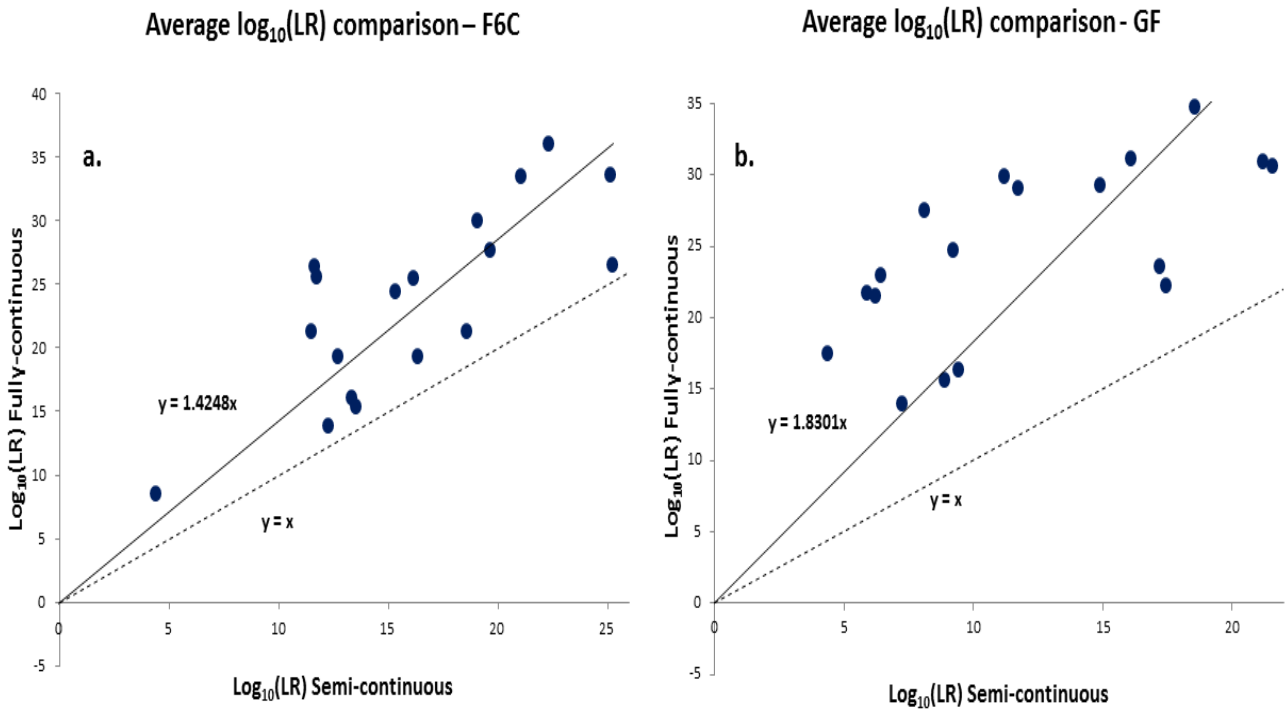
569



571

572 **Figure 1** Log(LR) results relative to 2-person and 3-person mixtures that were amplified with Fusion 6C (F6C,  
 573 figures 1a, 1c, 1e) and GlobalFiler (GF, figures 1b, 1d, 1f) DNA amplification kits. Log(LR) results of the known  
 574 contributor labelled as NIST A (a-b), NIST B (c-d) and NIST C (e-f) are shown. In particular, log(LR) values are  
 575 represented by: blue diamonds for STRmix™, orange squares for EuroForMix, purple triangles for  
 576 DNA•VIEW®, red addition marks for LRMix Studio and green crosses for Lab Retriever.

577

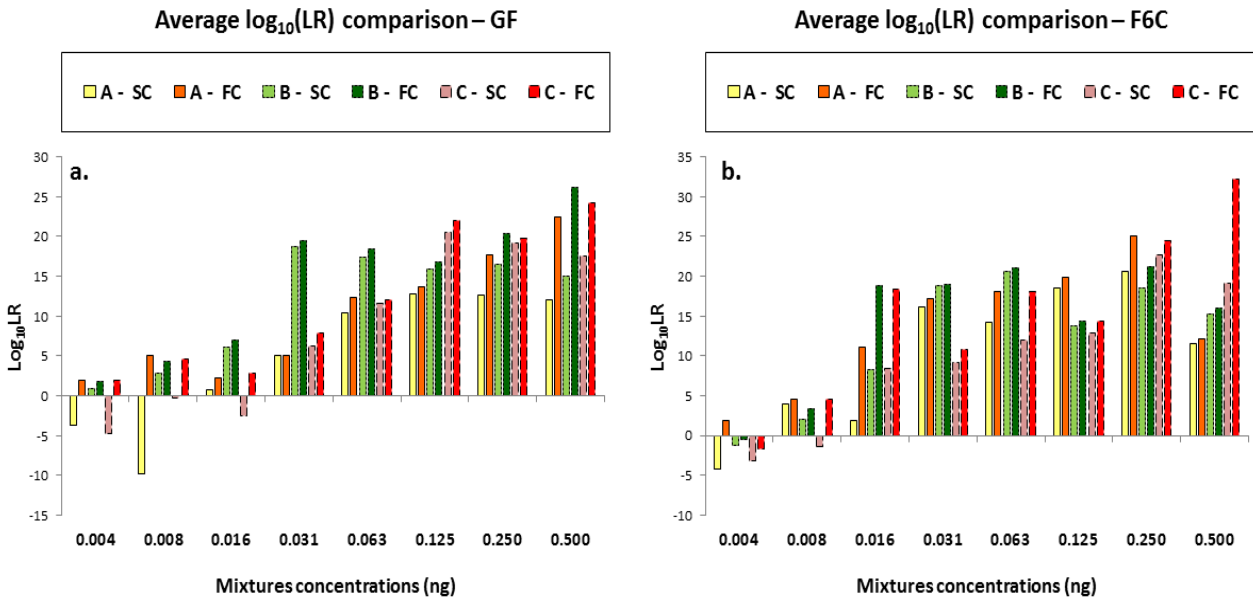


579

580 **Figure 2** Mean  $\log(\text{LR})$  values provided by semi-continuous and fully-continuous models for F6C (Figure 2a)  
 581 and GF (Figure 2b) amplification kits. The dashed line represents a hypothetical situation where all the  
 582  $\log(\text{LR})$  results are the same for both the investigated models, while the solid line (with intercept equal to  
 583 zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous  
 584 algorithms.

585





586

587 **Figure 3** Histograms displaying average  $\log(\text{LR})$  values provided by semi-continuous (SC) and fully-continuous  
 588 (FC) models for 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified by GF (a) and F6C  
 589 (b) amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. Mean  
 590  $\log(\text{LR})$  values relative to NIST materials composing the mixtures are represented by yellow (SC) and orange  
 591 (FC) histograms for NIST A contributor, light green (SC) and dark green (FC) histograms for NIST B contributor,  
 592 pink (SC) and red (FC) histograms for NIST C contributor.

593



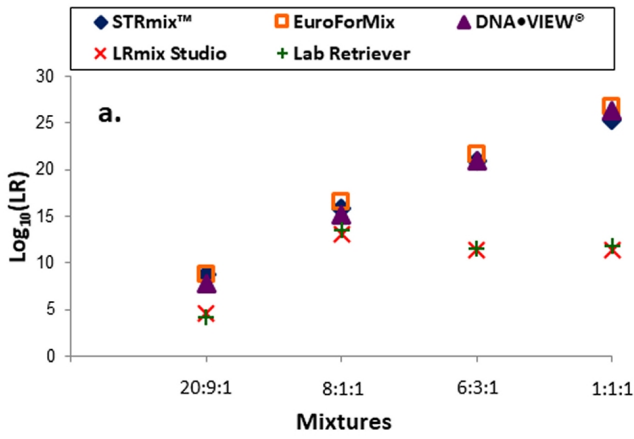
SD3 Channel Based Analytical Threshold					
Channel	DStdDev	Variance	Avg PBHW	Thresholds	SD3
Blue	5,002	25,022	10,000	47,455	48
Green	9,435	89,012	10,000	89,505	90
Yellow	4,977	24,766	10,000	47,212	48
Red	8,342	69,596	10,000	79,143	80
Purple	8,973	80,508	10,000	85,122	86
* Blue, Green, Yellow, Red, Purple channel(s) have no peaks.					
Channel Based RFU Extremums					
Channel	Max-Min Range	Avg Min RFU	Avg Max RFU	Avg RFU Range	Min-Max Fitted Lines Range Avg
Blue	60	-9,65	8,42	36,14	35,95
Green	132	-17,33	16,90	68,47	68,30
Yellow	76	-10,60	8,55	38,30	38,30
Red	96	-15,25	14,53	59,55	59,54
Purple	104	-16,60	16,84	66,88	66,86
Avg	93,60	-13,89	13,05	53,87	53,79

595

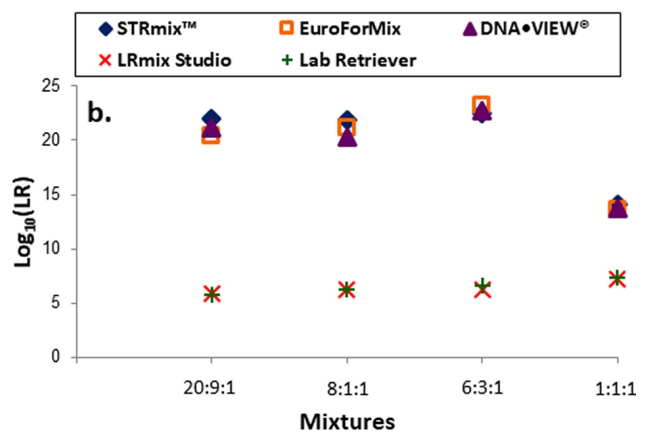
596 **Figure 4** ArmedXpert™ results relative to the algorithms performing the calculation of differential analytical  
597 thresholds (i.e. per dye channel).

598

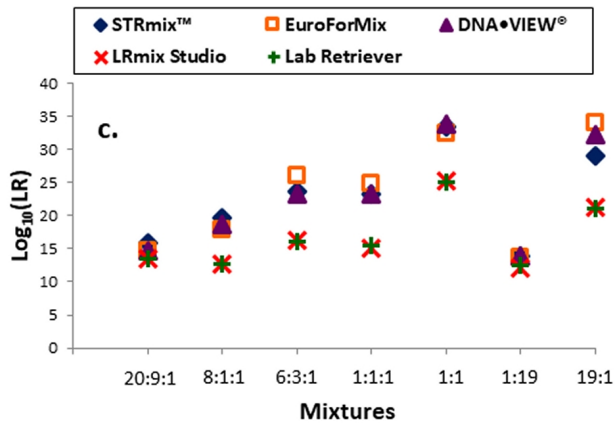
NIST A – F6C



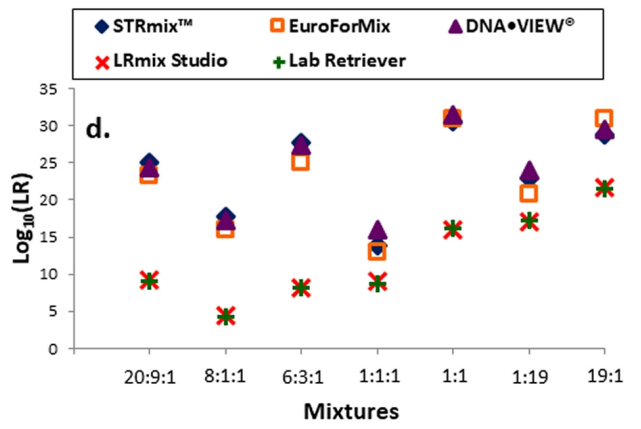
NIST A – GF



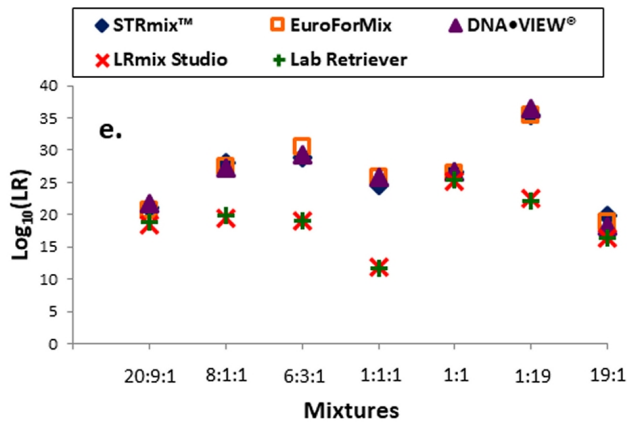
NIST B – F6C



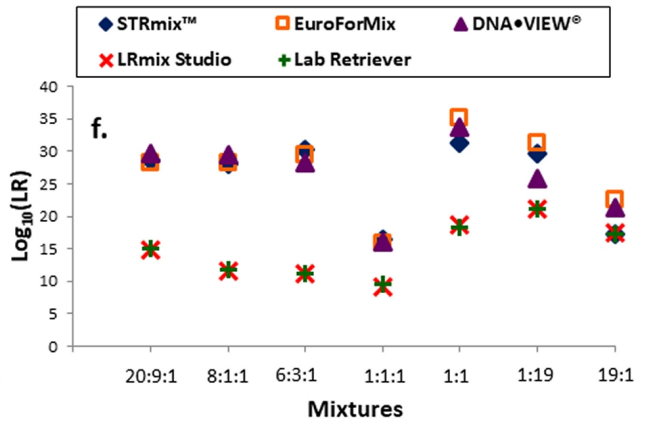
NIST B – GF



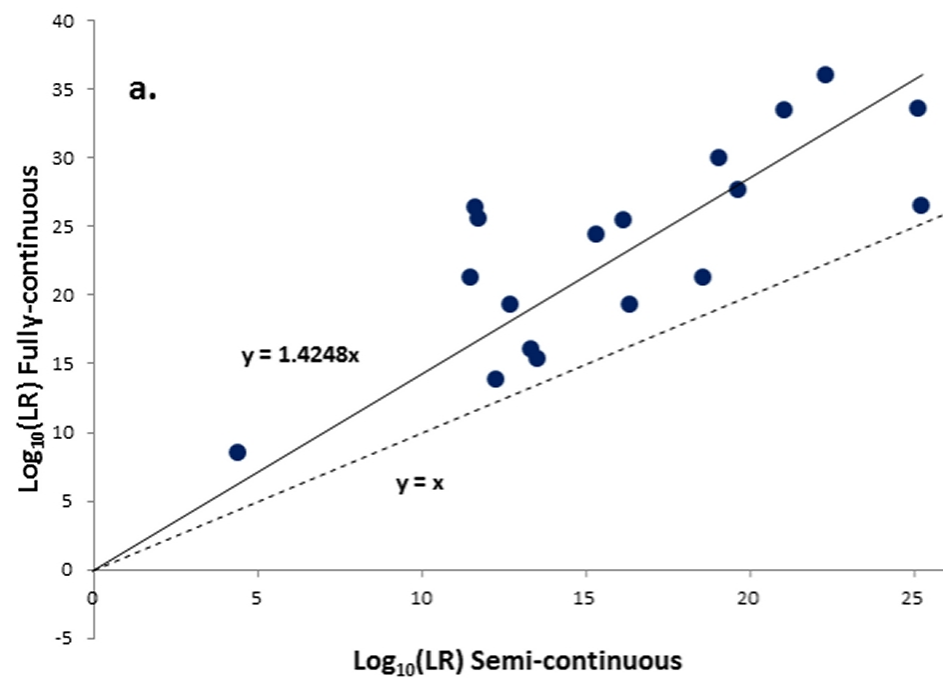
NIST C – F6C



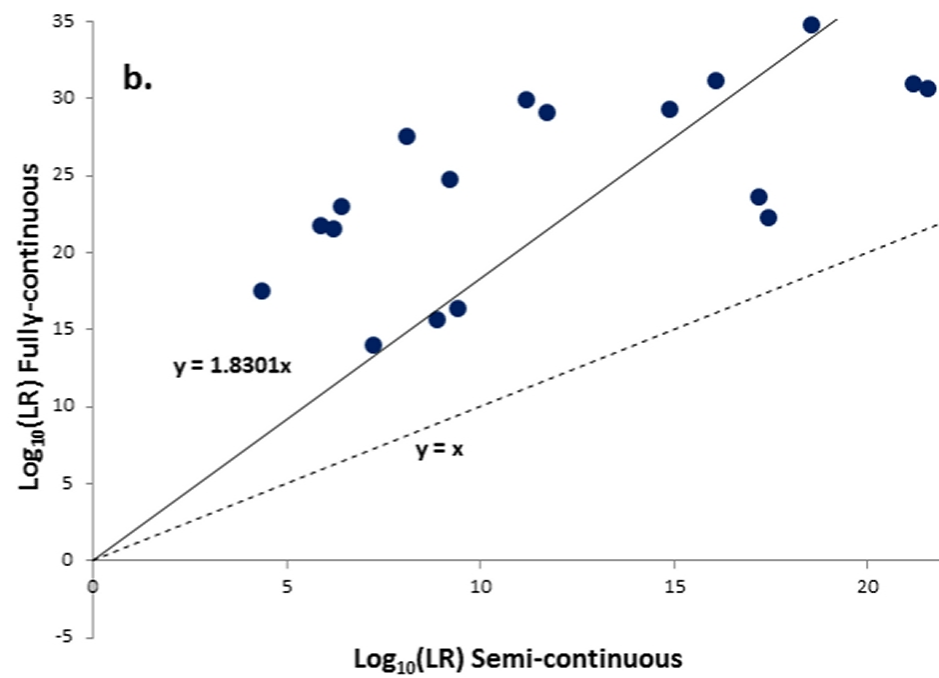
NIST C – GF



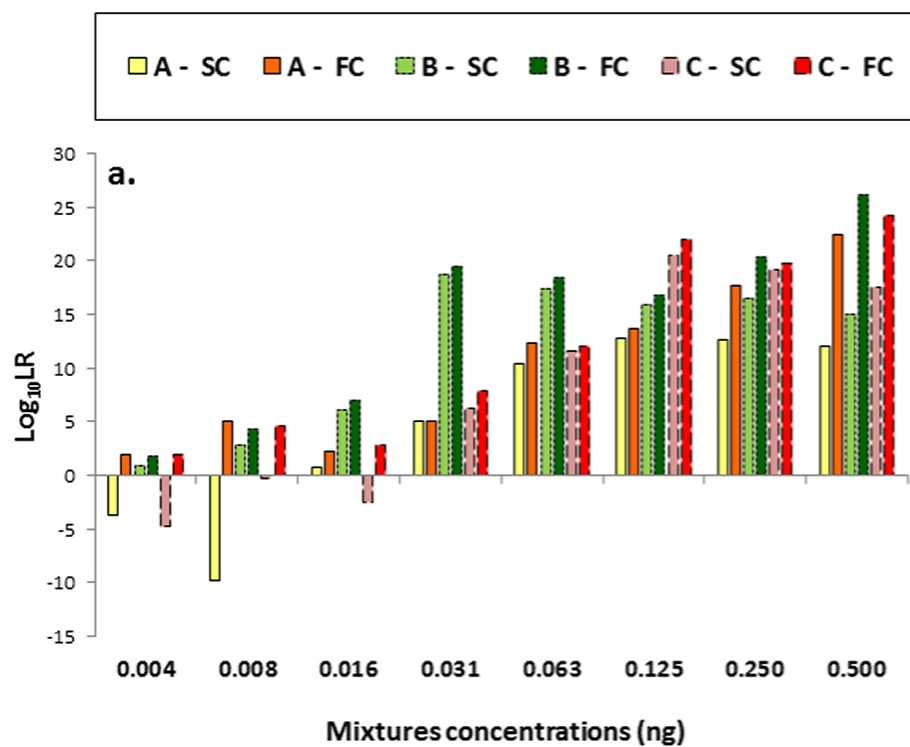
Average  $\log_{10}(\text{LR})$  comparison – F6C



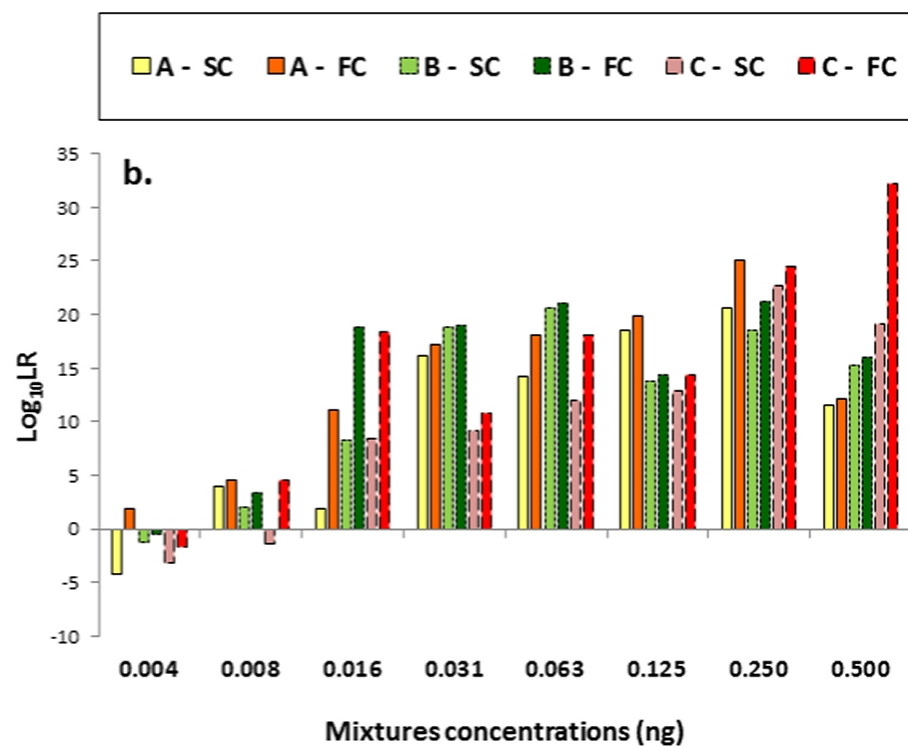
Average  $\log_{10}(\text{LR})$  comparison - GF



Average  $\log_{10}(\text{LR})$  comparison – GF



Average  $\log_{10}(\text{LR})$  comparison – F6C



## SD3 Channel Based Analytical Threshold

Channel	DStdDev	Variance	Avg PBHW	Thresholds	SD3
Blue	5,002	25,022	10,000	47,455	48
Green	9,435	89,012	10,000	89,505	90
Yellow	4,977	24,766	10,000	47,212	48
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Purple	8,973	80,508	10,000	85,122	86

\* Blue, Green, Yellow, Red, Purple channel(s) have no peaks.

## Channel Based RFU Extremums

Channel	Max-Min Range	Avg Min RFU	Avg Max RFU	Avg RFU Range	Min-Max Fitted Lines Range Avg
Blue	60	-9,65	8,42	36,14	35,95
Green	132	-17,33	16,90	68,47	68,30
Yellow	76	-10,60	8,55	38,30	38,30
Red	96	-15,25	14,53	59,55	59,54
Purple	104	-16,60	16,84	66,88	66,86
Avg	93,60	-13,89	13,05	53,87	53,79

**Table 1** List of NIST samples used as known contributors for the preparation of DNA mixtures. Different proportions were evaluated for both 2- and 3-person mixtures. Referring to 2-person mixtures, different NIST materials were employed for the preparation of the samples according to the utilised DNA amplification kit.

<b>Known 2-person mixtures</b>		
<b>DNA Typing Kit</b>	<b>Reference Material</b>	<b>Mixtures ratios (0.500 ng)</b>
ESI 17 Fast	B:C / male:male	
ESX 17 Fast	A:C / female:male	19:1
Fusion	B:C / male:male	8:1
Fusion 6C	B:C / male:male	1:1
GlobalFiler	B:C / male:male	1:19
Mini Filer	A:C / female:male	
NGM SElect	A:C / female:male	
<b>Known 3-person mixtures</b>		
<b>DNA Typing Kit</b>	<b>Reference Material</b>	<b>Mixtures ratios (0.500 ng)</b>
		1:1:1
All kits	A:B:C / Female:male:male	6:3:1
		8:1:1
		20:9:1



**Table 2** Log(LR) results relative to the interpretation process performed on the DNA mixture collected on a cap recovered on a crime scene. 3SD and Min-Max represent the algorithms that were employed to evaluate the differential analytical thresholds. POI represents the suspect (i.e. the person of interest), while U stands for unknown(s) individual(s) extracted from the allele frequencies reference dataset [47].

<b>Analytical Threshold</b>	<b>3SD</b>	<b>Min-Max</b>	<b>3SD</b>	<b>Min-Max</b>
<b>Hp</b>	S + 1U	S + 1U	S + 2U	S + 2U
<b>Hd</b>	2U	2U	3U	3U
<b>Software</b>	<b>Log(LR)</b>			
<b>LRmix Studio</b>	-0.37	-0.16	2.36	2.48
<b>Lab Retriever</b>	-0.36	-0.16	2.37	2.49
<b>DNA•VIEW®</b>	2.29	3.47	6.79	7.06
<b>EuroForMix</b>	2.33	3.60	6.25	7.12
<b>STRmix™</b>	2.84	3.63	7.01	7.03
<b>Interpretative decision</b>	<b>Inconclusive</b>	<b>Inconclusive</b>	<b>Support to H(p)</b>	<b>Support to H(p)</b>

## **Authors' contributions**

EA conceived the study, carried out the multivariate studies and drafted the manuscript.

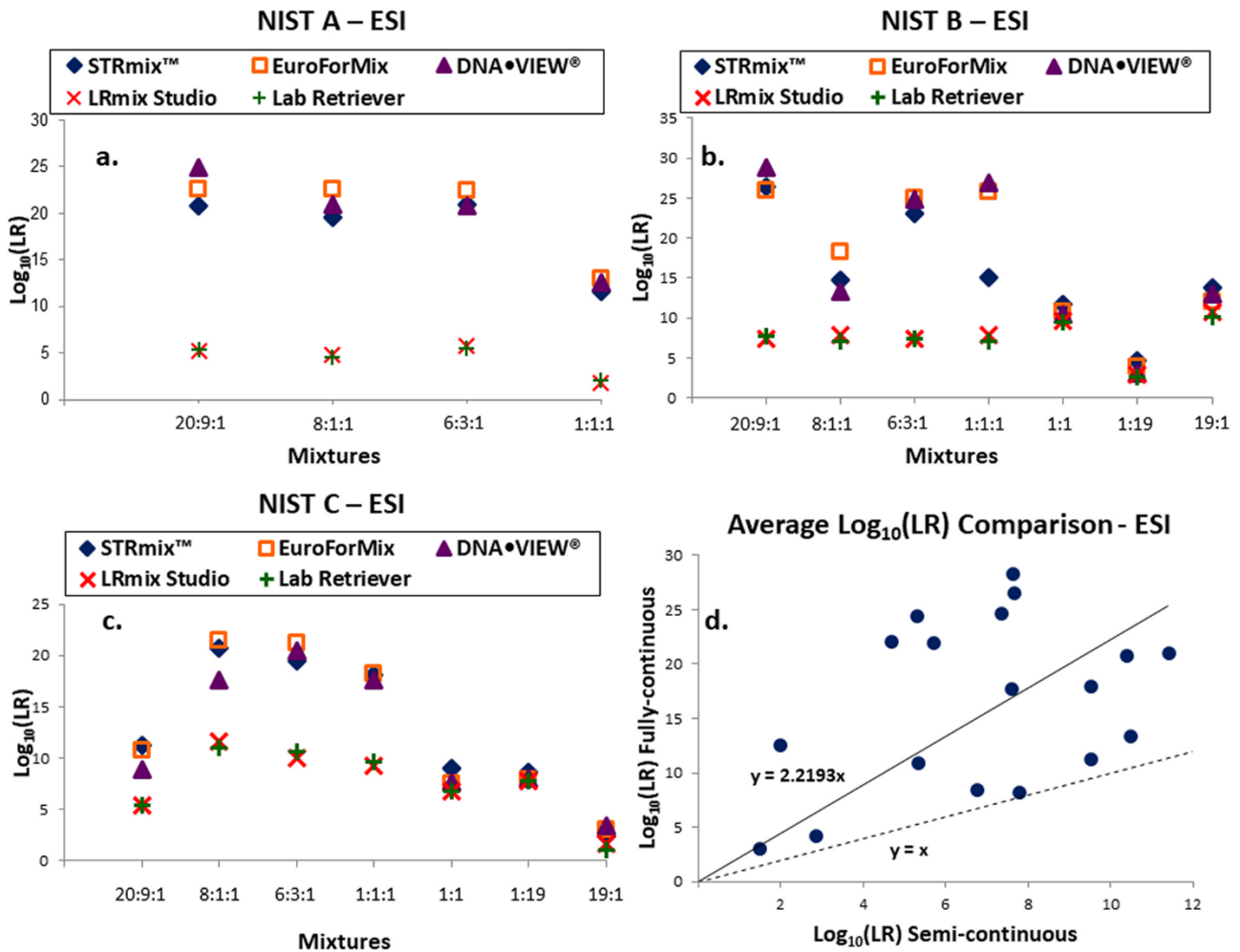
MO, GD, DC and PG participated in the design of the study, carried out the genotyping step and helped to draft the manuscript.

MV participated in the design of the study and helped to draft the manuscript.

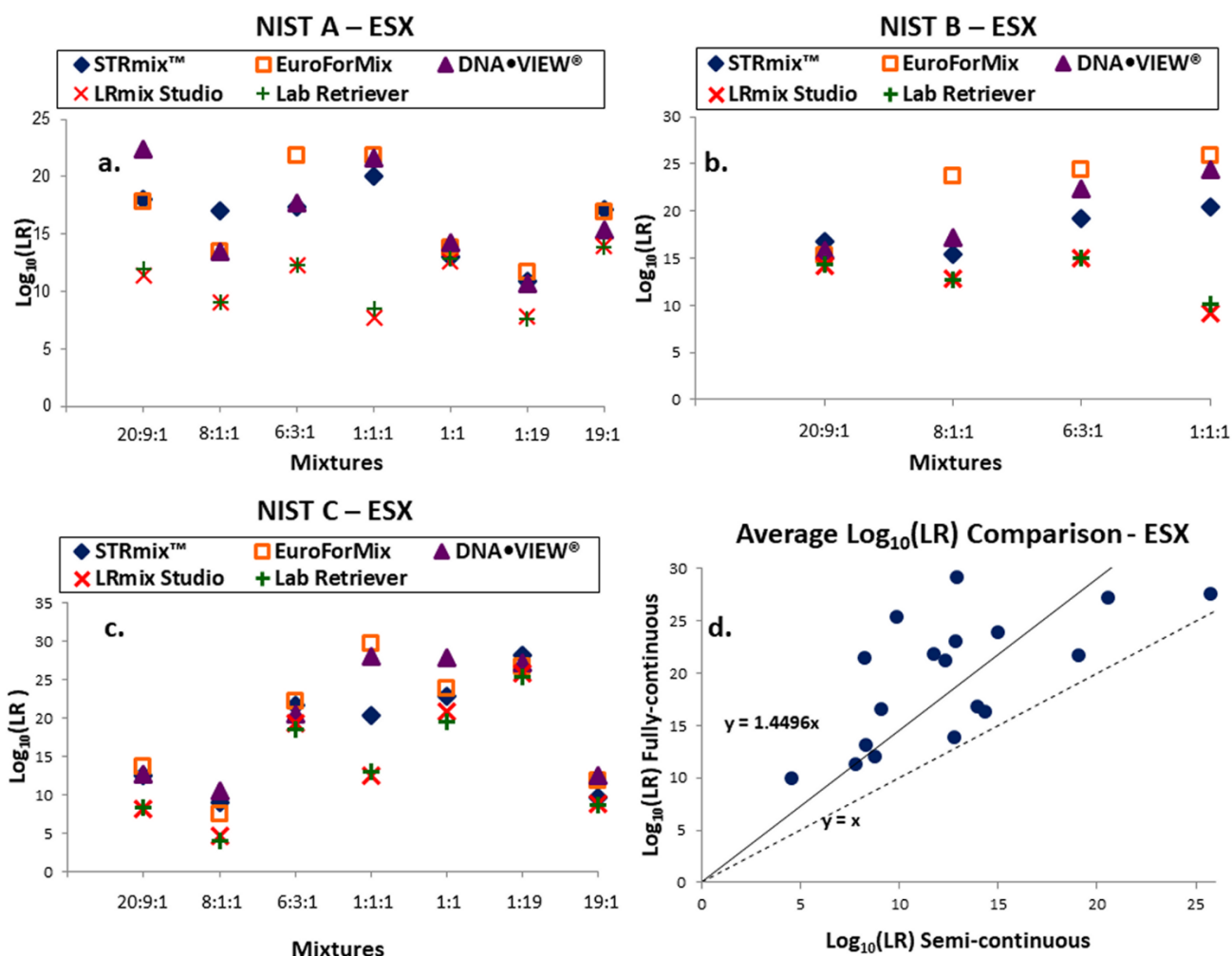
All authors read and approved the final manuscript.

## **Acknowledgements**

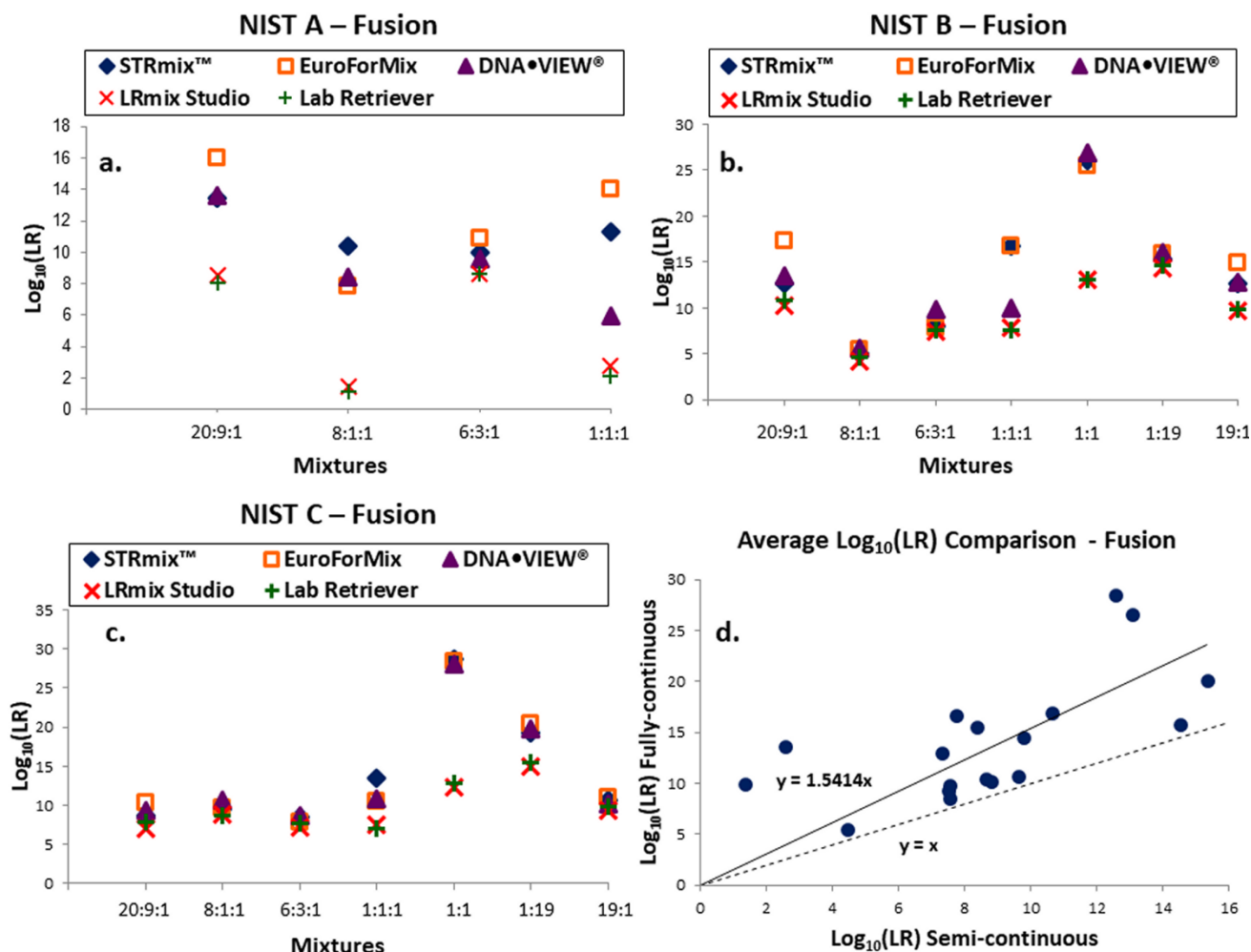
This study was partially founded by the Fondazione Giovanni Gorla, within the grant program named “Bando Talenti della Società Civile”. Continuous support from M.I.U.R. and Regione Piemonte is kindly acknowledged.



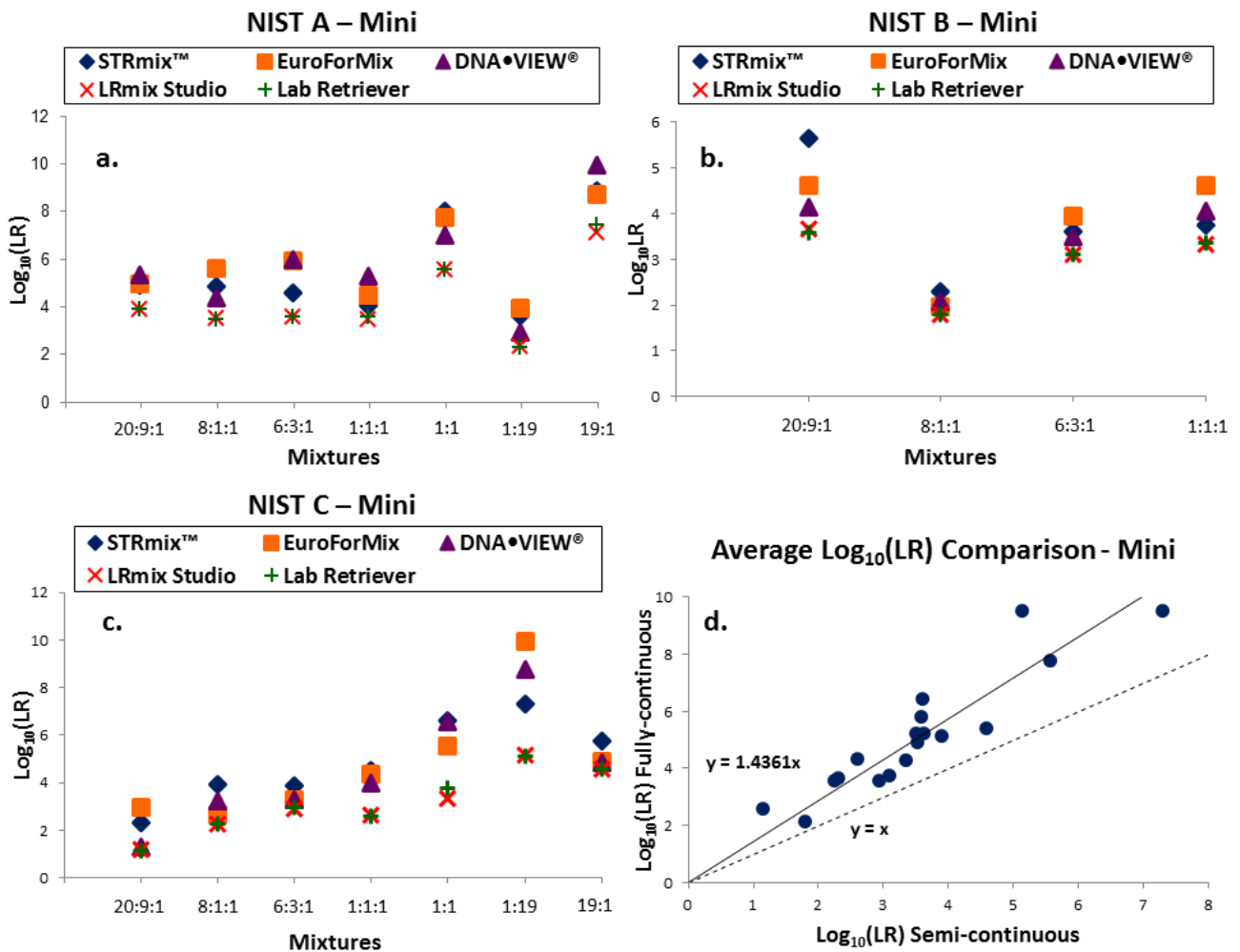
**Figure S1** Log(LR) results relative to 2-person and 3-person mixtures that were amplified with ESI DNA amplification kit. Log(LR) results of the known contributor labelled as NIST A (a), NIST B (b) and NIST C (c) are shown. In particular, log(LR) values are represented by: blue diamonds for STRmix™, orange squares for Euroformix, purple triangles for DNA•VIEW®, red addition marks for LRmix Studio and green crosses for Lab Retriever. Average log(LR) values provided by semi-continuous and fully-continuous models for ESI DNA amplification kit are remarked in (d). The dashed line represents an hypothetic situation where all the log(LR) results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms.



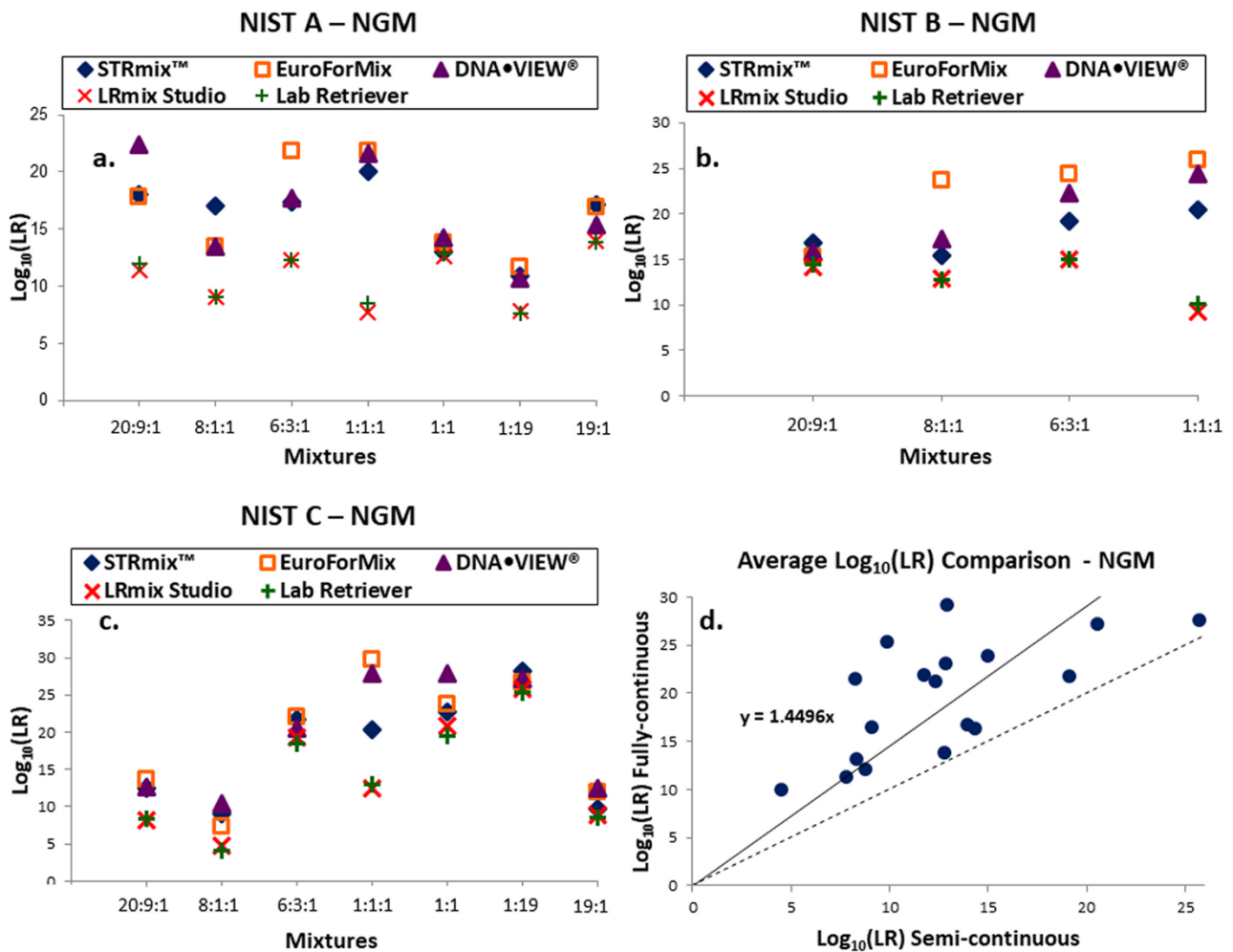
**Figure S2** Log(LR) results relative to 2-person and 3-person mixtures that were amplified with ESX DNA amplification kit. Log(LR) results of the known contributor labelled as NIST A (a), NIST B (b) and NIST C (c) are shown. In particular, log(LR) values are represented by: blue diamonds for STRmix™, orange squares for Euroformix, purple triangles for DNA•VIEW®, red addition marks for LRmix Studio and green crosses for Lab Retriever. Average log(LR) values provided by semi-continuous and fully-continuous models for ESX DNA amplification kit are remarked in (d). The dashed line represents an hypothetical situation where all the log(LR) results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms.



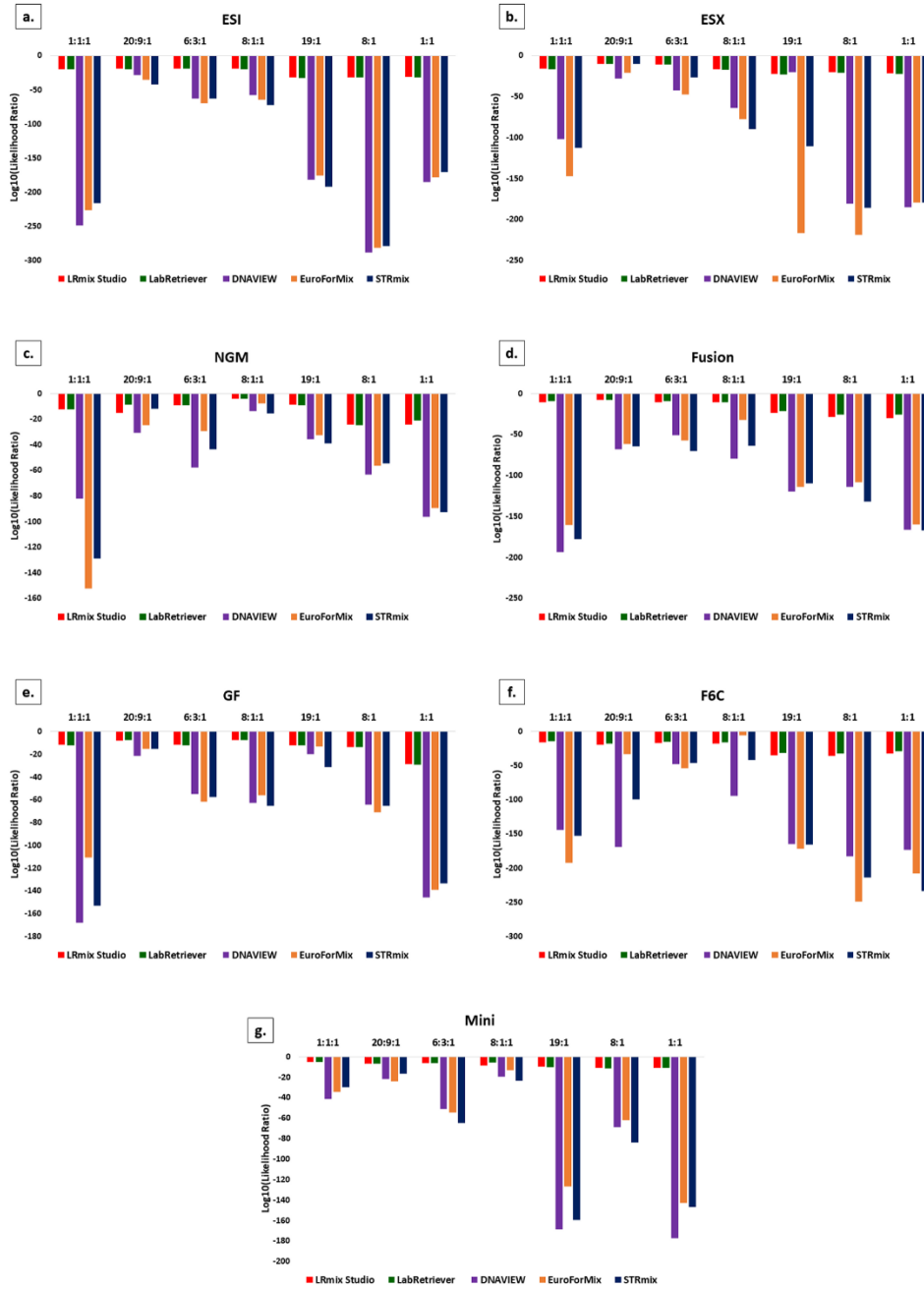
**Figure S3** Log(LR) results relative to 2-person and 3-person mixtures that were amplified with Fusion DNA amplification kit. Log(LR) results of the known contributor labelled as NIST A (a), NIST B (b) and NIST C (c) are shown. In particular, log(LR) values are represented by: blue diamonds for STRmix™, orange squares for Euroformix, purple triangles for DNA•VIEW®, red addition marks for LRMix Studio and green crosses for Lab Retriever. Average log(LR) values provided by semi-continuous and fully-continuous models for Fusion DNA amplification kit are remarked in (d). The dashed line represents an hypothetic situation where all the log(LR) results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms.



**Figure S4**  $\text{Log}(\text{LR})$  results relative to 2-person and 3-person mixtures that were amplified with Mini DNA amplification kit.  $\text{Log}(\text{LR})$  results of the known contributor labelled as NIST A (a), NIST B (b) and NIST C (c) are shown. In particular,  $\text{log}(\text{LR})$  values are represented by: blue diamonds for STRmix™, orange squares for Euroformix, purple triangles for DNA•VIEW®, red addition marks for LRmix Studio and green crosses for Lab Retriever. Average  $\text{log}(\text{LR})$  values provided by semi-continuous and fully-continuous models for Mini DNA amplification kit are remarked in (d). The dashed line represents an hypothetic situation where all the  $\text{log}(\text{LR})$  results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms.

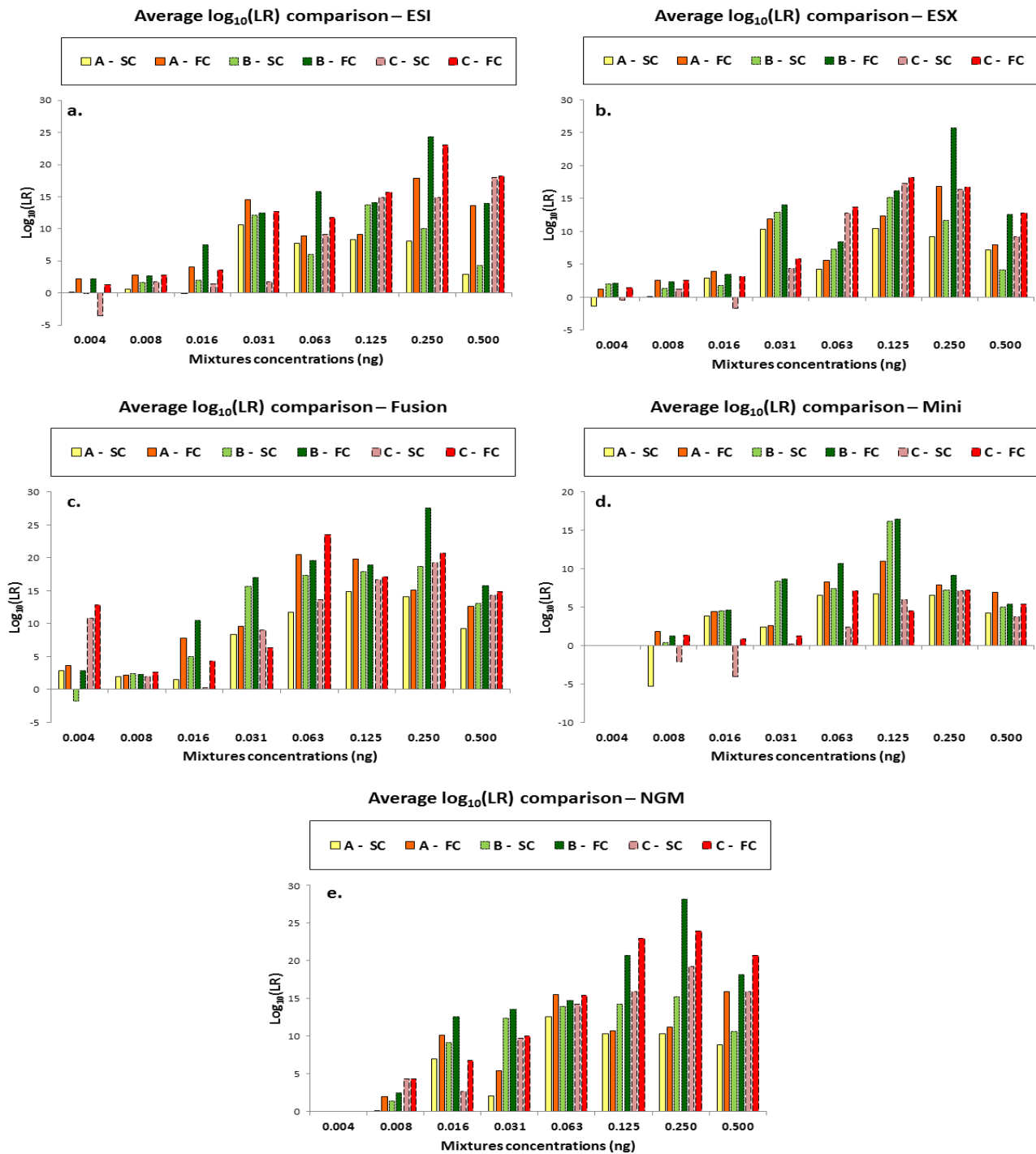


**Figure S5** Log(LR) results relative to 2-person and 3-person mixtures that were amplified with NGM DNA amplification kit. Log(LR) results of the known contributor labelled as NIST A (a), NIST B (b) and NIST C (c) are shown. In particular, log(LR) values are represented by: blue diamonds for STRmix™, orange squares for Euroformix, purple triangles for DNA•VIEW®, red addition marks for LRmix Studio and green crosses for Lab Retriever. Average log(LR) values provided by semi-continuous and fully-continuous models for NGM DNA amplification kit are remarked in (d). The dashed line represents an hypothetic situation where all the log(LR) results are the same for both the investigated models, while the solid line (with intercept equal to zero) indicates the average trend that is observed among the results provided by semi- and fully-continuous algorithms.

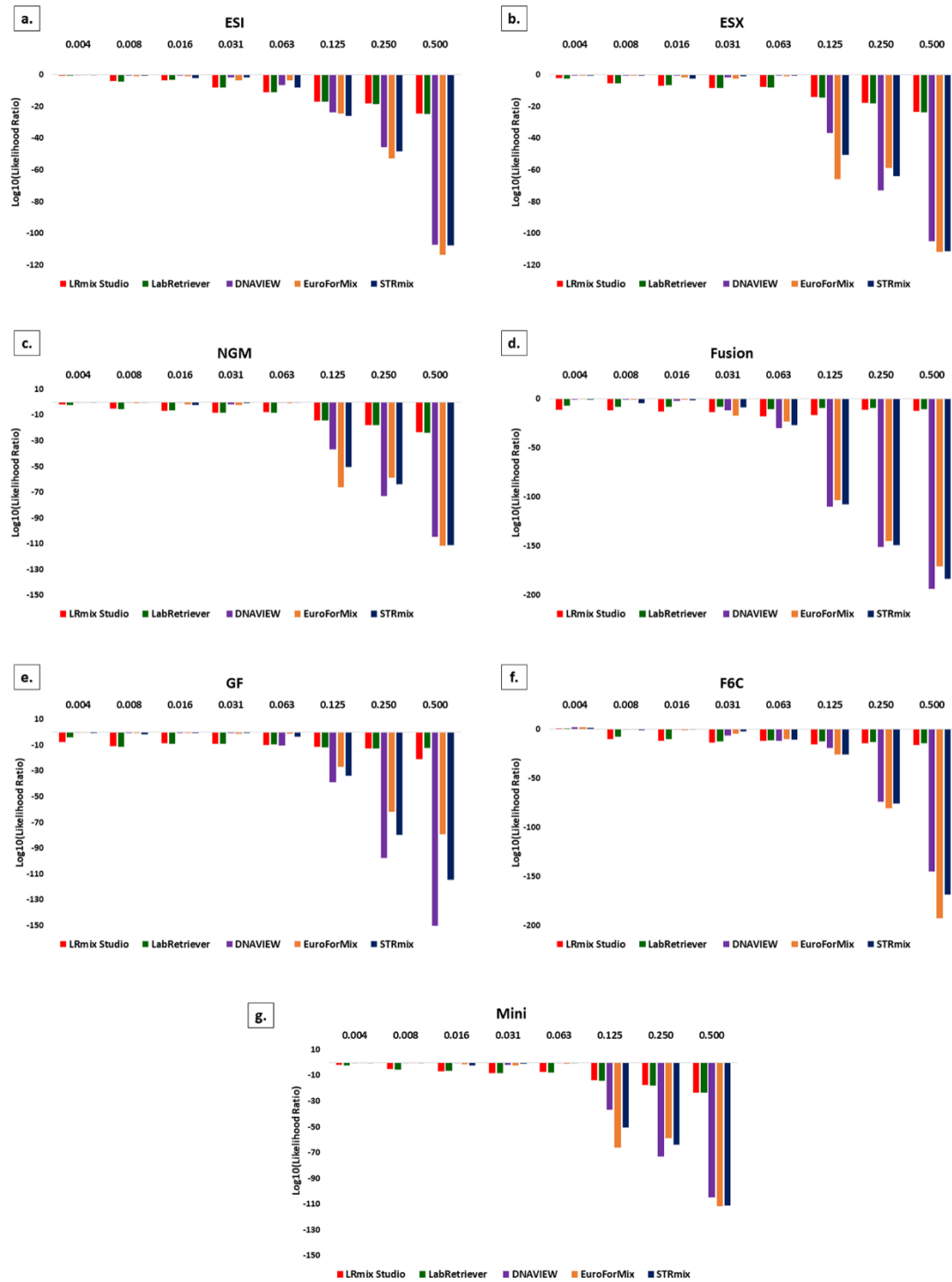


**Figure S6** Histograms displaying false donor (i.e. NIST F) log(LR) values log(LR) results relative to 2-person and 3-person mixtures that were amplified with different DNA amplification kits. DNA mixtures were amplified by ESI (a), ESX (b), NGM (c), Fusion (d), GlobalFiler (e), Fusion 6C (f) and MiniFiler (g) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. Log(LR) values are represented by red (LRmix Studio), green (Lab Retriever), purple (DनावIEW®), orange (EuroForMix) and blue (STRmix™) histograms.

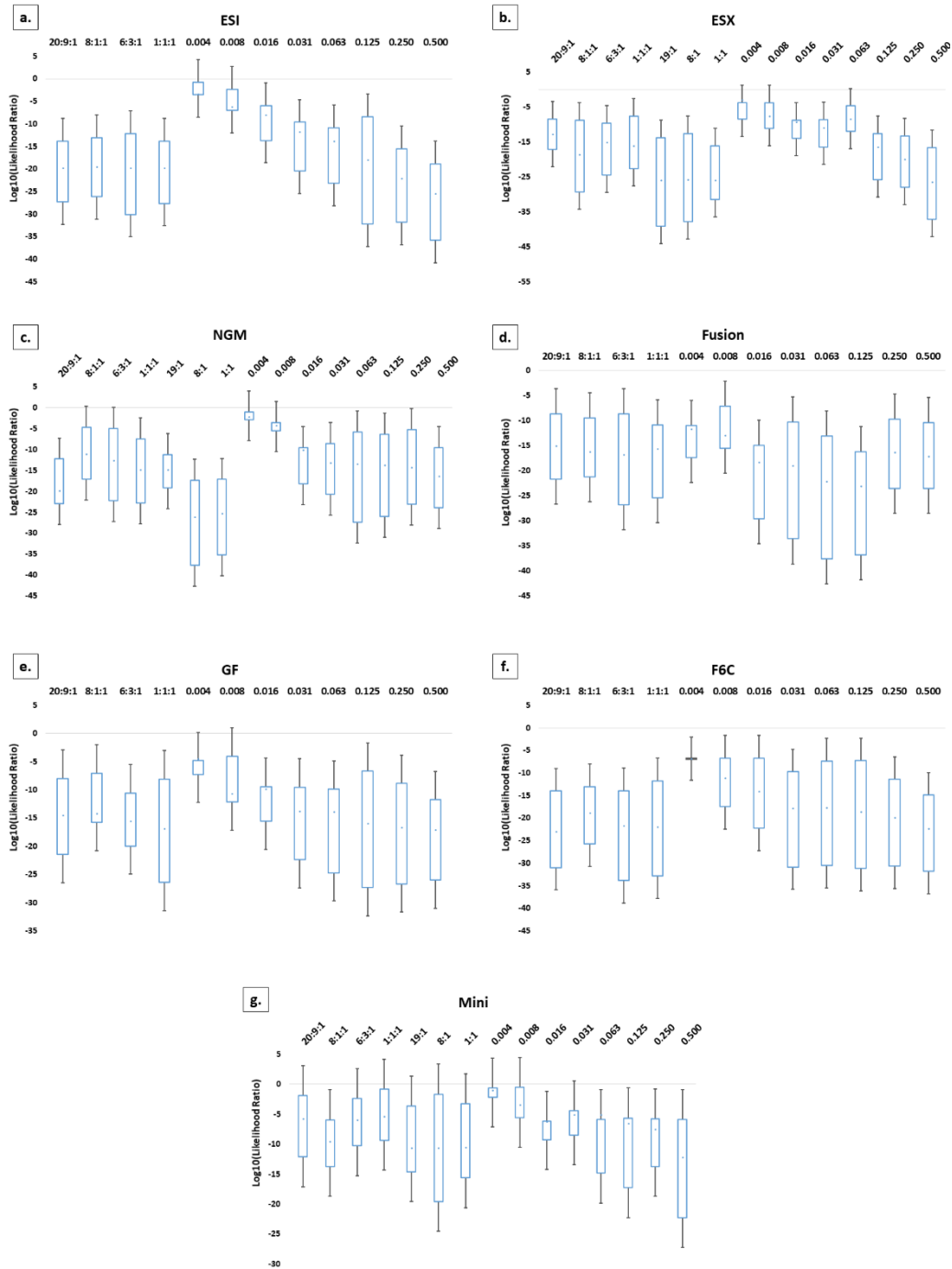




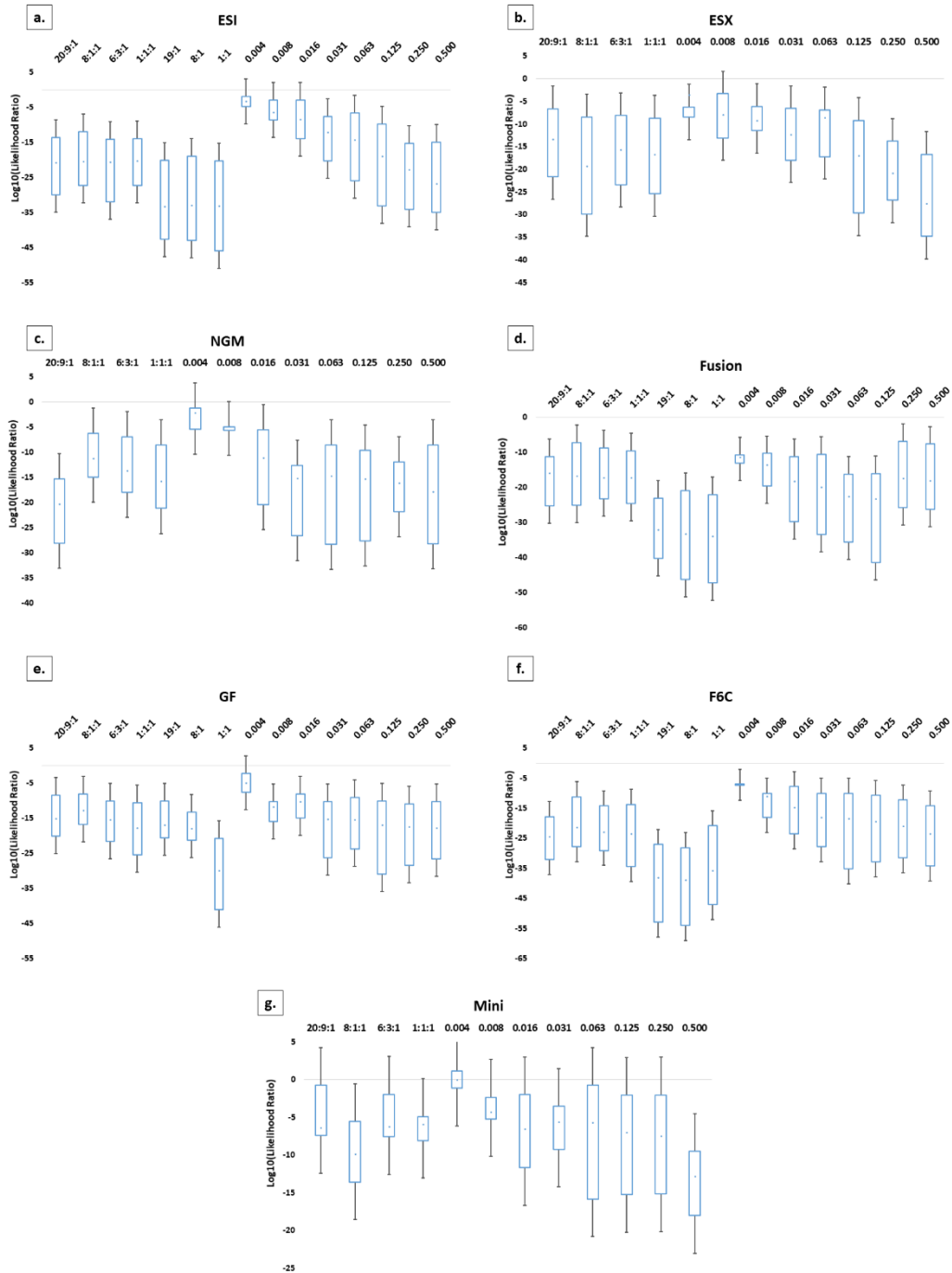
**Figure S7** Histograms displaying average  $\log(\text{LR})$  values provided by semi-continuous (SC) and fully-continuous (FC) models for 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified by ESI (a), ESX (b), Fusion (c), Mini (d) and NGM (c) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. Average  $\log(\text{LR})$  values relative to NIST materials composing the mixtures are represented by yellow (SC) and orange (FC) histograms for NIST A contributor, light green (SC) and dark green (FC) histograms for NIST B contributor, pink (SC) and red (FC) histograms for NIST C contributor. Data relevant to 0.004 ng mixtures for Mini and NGM are not available as such DNA mixtures didn't provide any accessible results when amplified by such DNA amplification kits.



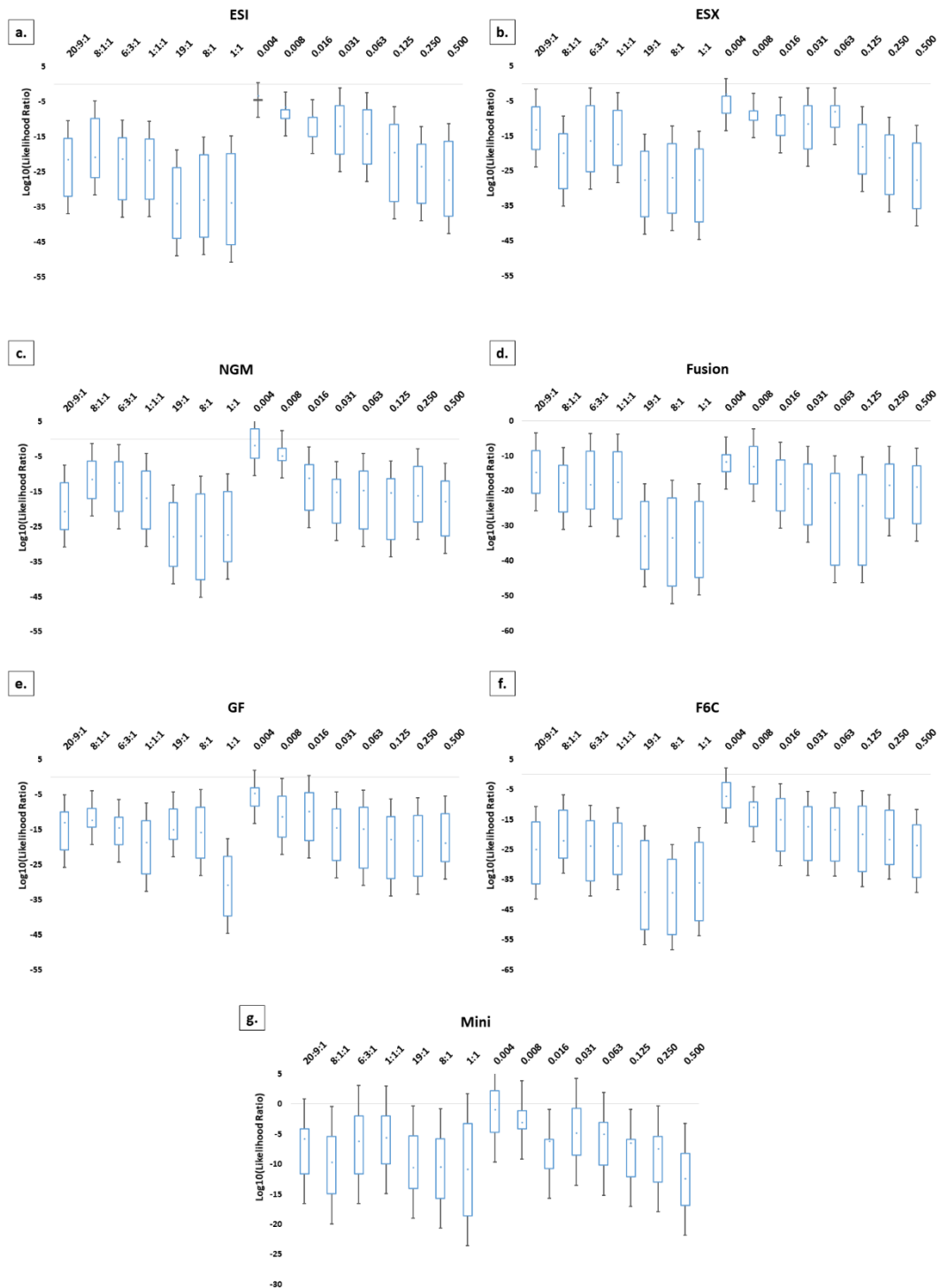
**Figure S8** Histograms displaying false donor (i.e. NIST F) log(LR) values log(LR) results relative to the 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified with different DNA amplification kits. DNA mixtures were amplified by ESI (a), ESX (b), NGM (c), Fusion (d), GlobalFiler (e), Fusion 6C (f) and MiniFiler (g) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. Log(LR) values are represented by red (LRmix Studio), green (Lab Retriever), purple (DनावIEW®), orange (EuroForMix) and blue (STRmix™) histograms.



**Figure S9a** Boxplots displaying non-contributor tests for NIST A and their relative log(LR) values relative to the 2-person and 3-person mixtures and the 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified with different DNA amplification kits. DNA mixtures were amplified by ESI (a), ESX (b), NGM (c), Fusion (d), GlobalFiler (e), Fusion 6C (f) and MiniFiler (g) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. The dot within the boxplot represents the median value.



**Figure S9b** Boxplots displaying non-contributor tests for NIST B and their relative log(LR) values relative to the 2-person and 3-person mixtures and the 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified with different DNA amplification kits. DNA mixtures were amplified by ESI (a), ESX (b), NGM (c), Fusion (d), GlobalFiler (e), Fusion 6C (f) and MiniFiler (g) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. The dot within the boxplot represents the median value.



**Figure S9c** Boxplots displaying non-contributor tests for NIST C and their relative log(LR) values relative to the 2-person and 3-person mixtures and the 1:1:1 serially scalarly-diluted 3-person DNA mixtures that were amplified with different DNA amplification kits. DNA mixtures were amplified by ESI (a), ESX (b), NGM (c), Fusion (d), GlobalFiler (e), Fusion 6C (f) and MiniFiler (g) DNA amplification kits. The codes indicating the different DNA mixtures are reported on the x axis. The dot within the boxplot represents the median value.