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The Synergy between Whole Genome Amplification and STR Analysis in Forensic Science

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Introduction

Forensic science has undergone a profound transformation in the last few decades, particularly in the area of genetic analysis. Among the most powerful techniques for identifying individuals and solving crimes is DNA profiling. This has revolutionized criminal investigations, exonerated the wrongfully accused, and provided undeniable evidence in courtrooms. Central to this revolution are two techniques: Short Tandem Repeat (STR) analysis and Whole Genome Amplification (WGA). STR analysis involves the examination of specific regions of the human genome that contain short, repetitive sequences of DNA. These regions are highly polymorphic, meaning they vary significantly among individuals, making them invaluable for identification. On the other hand, WGA is a technique used to amplify small amounts of DNA, which is critical when DNA samples are degraded, limited, or compromised-common situations in forensic cases where only trace amounts of biological material remain [1].

Crime scene evidence often consists of trace amounts of biological material, such as hair, skin cells, or saliva. WGA allows forensic scientists to amplify these tiny amounts of DNA, making it possible to generate an STR profile from even the most challenging samples. This is particularly important in cases where DNA is recovered from hard-to-access areas or when a suspect is not readily available for direct comparison. In cases where DNA from multiple individuals is mixed (e.g., in sexual assault cases or violent altercations), the ability to amplify the entire genome through WGA allows for better discrimination of individual profiles. This improves the chances of identifying the perpetrator and resolving complex cases where multiple DNA sources are present [2].

Description

Short Tandem Repeat (STR) analysis is a genetic technique that examines specific locations in the genome where short, repeating sequences of DNA occur. These repeats are typically 2–6 base pairs long and are found in numerous locations across the human genome. The number of repeats at a particular STR locus can vary between individuals, which allows forensic scientists to use these loci for identification purposes. The power of STRs in forensic science lies in their polymorphism, or variability. While most of the genome is conserved across individuals, STR regions differ in length from person to person. This makes them an ideal tool for distinguishing between individuals in cases such as criminal investigations, paternity testing, and missing persons cases. In forensic DNA analysis, STR markers are typically selected from a set of loci known to be highly variable. By examining multiple loci simultaneously, forensic scientists can obtain a DNA profile that is unique to an individual. This profile is used to compare samples recovered from a crime scene with those from suspects or databases [3].

A typical STR profile involves analyzing 13-15 different loci, with each

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locus containing a set of repeated sequences. The results are often presented as a series of numbers representing the number of repeats at each locus. The combination of these results creates a genetic fingerprint that is unique to an individual, except in the case of identical twins. Whole Genome Amplification (WGA) is a method used to amplify an entire genome from a small or degraded DNA sample. This technique is especially important in forensic science, where DNA samples are often available in limited quantities and can be highly fragmented due to environmental factors. The goal of WGA is to generate a sufficient quantity of DNA for downstream analysis, such as STR profiling or sequencing. WGA methods can be broadly classified into two categories: PCR-based amplification and non-PCR-based amplification. PCR-based WGA uses primers that bind to the DNA and initiate the amplification of the entire genome. Non-PCR-based WGA, on the other hand, employs chemical or enzymatic reactions to amplify the genome without the need for primers [4].

The use of WGA in forensic science is critical when working with compromised or degraded samples. For example, DNA recovered from bones, hair, or old biological samples may be fragmented or present in very low quantities. WGA allows forensic scientists to create a more complete and usable DNA sample, which increases the chances of successful STR analysis. In forensic applications, WGA helps amplify genetic material from minute or degraded sources, such as blood stains, hair follicles, semen, or tissue remnants, ensuring that even trace amounts of DNA can be analyzed effectively. While STR analysis provides a powerful method for identifying individuals based on specific regions of their genome, it is dependent on the availability of a sufficient amount of high-quality DNA. In cases where the DNA sample is degraded or present in minute quantities, the ability to amplify the genome through WGA becomes crucial. Amplifying Degraded or Low-Quantity DNA: In forensic cases, DNA samples from crime scenes can be degraded due to exposure to environmental factors such as heat, moisture, or bacteria. Traditional DNA analysis methods may fail to yield sufficient results from these compromised samples. However, WGA can amplify even tiny or degraded samples, generating enough DNA to produce a full STR profile [5].

Conclusion

The integration of Whole Genome Amplification (WGA) and Short Tandem Repeat (STR) analysis has brought about a powerful synergy in forensic science. WGA enhances the success of STR analysis by allowing forensic scientists to work with degraded or minute DNA samples, ensuring that even trace amounts of DNA can be effectively analyzed. This combination has not only improved the reliability and accuracy of forensic DNA profiling but has also expanded its applicability in cold cases, missing person investigations, and criminal justice. While challenges such as contamination risks and allelic dropout remain, the potential for using WGA and STR analysis to solve complex forensic cases continues to grow. As forensic techniques evolve and improve, the synergy between WGA and STR analysis will play a critical role in advancing forensic science and ensuring that justice is served. This union of technology has solidified its place as an invaluable tool in the forensic world, ensuring that even the most challenging cases have the potential for resolution. As forensic science continues to develop, the power of these techniques will only become more pronounced, further enhancing the ability to bring perpetrators to justice and provide closure for victims and families.

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Conflict of Interest

The author declares there is no conflict of interest associated with this manuscript.

References

- Li, Cai-xia, Jun-ping Han, Wen-yan Ren and An-quan Ji, et al. "DNA profiling of spermatozoa by laser capture microdissection and low volume-PCR." *PLoS One* 6, (2011): e22316.
- Schneider, C., U. Müller, R. Kilper and B. Siebertz. "Low copy number DNA profiling from isolated sperm using the aureka®-micromanipulation system." *Forensic Sci Int Genet* 6 (2012): 461-465.

- Lucy, D., J. M. Curran, A. A. Pirie and P. Gill. "The probability of achieving full allelic representation for LCN-STR profiling of haploid cells." *Sci Justice* 47 (2007): 168-171.
- Uchigasaki, Seisaku, Jian Tie, Erina Sobashima and Naomi Shimada. "Genotyping DNA isolated from UV irradiated human bloodstains using whole genome amplification." *Mol Biol Rep* 45 (2018): 925-929.
- Tate, Courtney M., Ada N. Nuñez, Cori A. Goldstein and Iva Gomes, et al. "Evaluation of circular DNA substrates for whole genome amplification prior to forensic analysis." *Forensic Sci Int Genet* 6 (2012): 185-190.

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