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Unknown Mutation detection via Restriction hybridization Method instead of using Next generation sequencing method

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In biology mutation is a change in the nucleotide sequences of the DNA of an organism. Mainly there are three types of mutation: point mutation, deletion and insertions. Once the mutation has been defined, allele-specific oligonucleotide hybridization, amplification, heteroduplex formation method referred to as a diagnostic method. Some advanced techniques like CRISPR cas9 system are used for selected mutagenesis. Using restriction method system we can detect a mutation. Let's say you have a DNA sample with fluorescently labeled from patient and you want to make sure that gene you are interested in is healthy gene. We can design different short fragment sequences to scan through DNA or find specific gene or mutation. The sequences scan the DNA; if the sequences do not find targeted gene, it does not bind to it, which means that no fluorescence color appears under UV-light. Each different short fragment sequence is labeled with different colors. If the different short fragments sequence does not bind to the DNA or specific gene or area, this means that there will be no color appear under UV light. This part or gene will be separated from the DNA by using Restriction enzyme to do a Sanger sequencing gel electrophoresis. Result of the Sanger sequencing will provide the information about sequence of unknown part or gene of the DNA. This method is easier and cost-effective method instead of Next generation sequencing method.

Result and observation:

In the given diagram, E, G, F does not show any color under UV light, which means that E, G, F part will be separated from the DNA in order to perform a Sanger gel electrophoresis. E, G, F have unknown variation or unknown SNP which cannot be available in any Database or data sections.