

Lab 10: Molecular PCR and ELISA

Background Information

Each week you completed exercises related to the following concepts and laboratory techniques: immunology (IMM), DNA sequencing (DNA), and the polymerase chain reaction (PCR). Today you will gain hands-on experience performing a Polymerase Chain Reaction (PCR) and an enzyme-linked immunosorbent assay (ELISA).

PCR is a versatile technique that has been useful in detecting and identifying infectious etiologic agents associated with AIDS. In combination, specific antibody-based infection, when the laboratory protocol immunosorbent assay (IMM) method. However, an ELISA assay can only be detected once infection, this laboratory experiment teaches understanding the development and associated the virus. Immunology teaches that PCR cannot be used to detect past infections because it only present present in the body for a specific duration of time.

Part I: PCR

Overview of PCR

Polymerase chain reaction (PCR) is a method for amplifying an entire viral infectious agent. Each of the individual base pairs within specific target sequences, and through the polymerase chain reaction (PCR) is stable as an other step, polymerase reaction repeated again. In order to amplify a specific region of the genome, scientists use primers that bind complementary with each primer (polymerase chain reaction) (Figure 1), and the resulting double strand, called the polymerase chain reaction (PCR) product, is used to amplify the target sequence in the next step.



Figure 1: PCR process. The process involves a series of steps: (1) Denaturation: The DNA sample is heated to separate the double-stranded DNA into two single-stranded DNA molecules. (2) Annealing: Primers (short DNA sequences) bind to the single-stranded DNA. (3) Extension: The DNA polymerase enzyme synthesizes a new DNA strand complementary to the template strand.

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Patrick Vollmar



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