

Title (en)

DIRECT LYSIS BUFFER AND THE DETECTION OF HIV-1 PLASMA VIREMIA

Title (de)

EIN PUFFER ZUR DIREKTEN LYSE UND DIE FESTSTELLUNG DER HIV-1 PLASMA VIRÄMIE

Title (fr)

TAMPON DE LYSE DIRECTE ET DETECTION DE LA PRESENCE DE VIH-1 DANS LE PLASMA

Publication

**EP 0698082 A1 19960228 (EN)**

Application

**EP 94915927 A 19940428**

Priority

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Abstract (en)

[origin: WO9426867A1] Immunocapture of plasma HIV-1, coupled with direct lysis of the virions and a simplified method of reverse transcription and amplification of the HIV-1 cDNA by the Polymerase Chain Reaction (PCR) represents a rapid and highly sensitive method to monitor HIV-1 disease progression. This method is also less time and labor intensive than quantitative culture. In addition, the development of a method to directly lyse the immunocaptured virions and a simplified single step reverse transcription (RT)/PCR procedure eliminated the need for organic solvent extraction and reduced the number of steps in the procedure. A direct lysis buffer was formulated to isolate plasma HIV-1 RNA for direct use in the RT and PCR reactions, thus eliminating the need for organic solvent extraction and ethanol precipitation. This resulted in a significant saving of time needed to complete the assay and significantly reduces the possibility of contamination associated with PCR reactions. The immunocapture-RT/PCR assay was used to show that vertical transmission of HIV-1 from a mother to her child depended largely on factors other than viral load. Conversely, the plasma viral load played a significant role in transfusion associated transmission of HIV-1 infection. Finally, the detection and quantitation of plasma associated viral load by immunocapture-RT/PCR may provide an additional marker of disease progression and may aid in determining the efficacy of various HIV therapeutics.

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**C11D 17/00; C12N 9/50**

IPC 8 full level

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