

Vás zve na seminář:

Influenza: Propagation, Quantification, and Storage

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Abstrakt



This unit covers several techniques for propagating, quantifying, and storing human influenza A viruses from existing stocks (see Basic Protocols 1

and 2) or from primary clinical specimens (see Alternate Protocols 2 and 4).

Virus isolation is a highly sensitive

and useful technique for the identification of viral infections. An important advantage

of virus isolation is the amplification of the virus from the original specimen, making it

available for further antigenic and genetic characterization. Influenza viruses are quantified

either by a "unit" of hemagglutination, which is not a measure of an absolute amount

of virus but is an operational unit dependent on the method used for the hemagglutination

assay titration (see Basic Protocol 3), or by determining infectious units using

the 50% tissue culture infectious dose assay (see Basic Protocol 4), 50% egg infectious

dose assay (see Basic Protocol 5), or plaque assay (see Basic Protocol 6). After isolating

and quantifying human influenza, the product must be properly stored to maintain virus viability.

Contributed by Kristy J. Szretter, Amanda L. Balish, and Jacqueline M. Katz *Current Protocols in Microbiology* (2006) 15G.1.1-15G.1.22 Copyright c_ 2006 by John Wiley & Sons, Inc.

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