

Improved Ebola Glycoprotein-Pseudotyped Retroviruses for Gene Transfer and Gene Therapy

A technology utilizing Ebola glycoprotein-pseudotyped retroviruses with improved efficiency enables gene therapy for the lung and provides a platform for quantitatively measuring virus entry and testing antiviral agents.

Transducing cells with viral vectors to deliver novel nucleic acids requires a specific ligand or envelope glycoprotein on the virus surface to bind to a receptor on a target cell. By changing the ligand on the virus (pseudotyping), it is possible to enhance the transduction efficiency towards target cell types. For this technology, Ebola glycoprotein-pseudotyped retroviruses were produced for gene transduction into the lung and other tissues and for the study of Ebola virus entry into cells.

Researchers at Purdue University have developed a technology that allows quantitative measurement of virus entry into host cells and determination of the effectiveness of reagents that inhibit entry. This technology also allows for the production of Ebola glycoprotein-pseudotyped retroviruses with markedly improved titers, making the practical use of such viruses more feasible.

Advantages:

- Allows gene therapy of the lung using pseudotyped retroviruses
- Improved safety and efficiency

Potential Applications:

- Medical/Healthcare
- Pharmaceuticals
- Drug Development
- Antivirals

Technology ID

61136

Category

Biotechnology & Life
Sciences/Cell & Gene Therapy
Platforms
Pharmaceuticals/Drug Discovery
& Development

Authors

Beverly Davidson
Scott Jeffers
Paul McCray
Anthony Sanchez
David Sanders
Patrick Sinn

Further information

Clayton Houck
CJHouck@prf.org

View online



-Gene/Cell Therapy

TRL: 6

Intellectual Property:

Provisional-Patent, 2002-06-04, United States | Provisional-Patent, 2003-03-27, United States | PCT-Patent, 2003-06-04, WO | Utility Patent, 2004-12-02, United States

Keywords: Viral vectors, gene transduction, pseudotyping, Ebola glycoprotein, retroviruses, virus entry measurement, antiviral screening, gene therapy, cell therapy, drug development