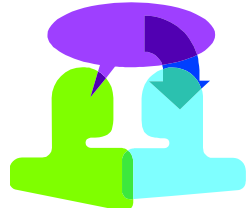




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Bayh-Dole Act

“The most inspired piece of US legislation
in the last 50 years” The Economist

- Patent and Trademark Law Amendments Act
- Bayh-Dole Act was implemented by Congress on December 12, 1980
- Permits universities and other non profits that receive federal funding for research to retain rights to their inventions in exchange for certain obligations



What is a Patent

- A patent is the right to exclude others from making, using, selling, offering for sale, or importing the patented invention
- Patents are valid only in the country that granted it
- Patents can be: Utility, Design or Plant
- Utility and Plant Patent Term: 20 years from date of filing
- Patents are limited to what is specified in the patent's claims



Patent Example



US009017935B2

(12) United States Patent
Dougherty et al.

(10) Patent No.: US 9,017,935 B2
(45) Date of Patent: Apr. 28, 2015

(54) HIV-1 LATENCY MODEL FOR HIGH THROUGHPUT SCREENING

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(73) Assignee: University of Medicine and Dentistry of New Jersey, New Brunswick, NJ (US)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 251 days.

(21) Appl. No.: 13/551,257

(22) Filed: Jul. 17, 2012

(65) Prior Publication Data

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(2013.01); *CI2N 15/86* (2013.01); *CI2N 2740/16043* (2013.01); *CI2N 2740/16122* (2013.01); *CI2Q 1/6897* (2013.01)

(58) Field of Classification Search
None
See application file for complete search history.

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Related U.S. Application Data

(62) Division of application No. 12/096,245, filed as application No. PCT/US2006/045483 on Nov. 27, 2006, now Pat. No. 8,247,167.

(60) Provisional application No. 60/742,241, filed on Dec. 5, 2005.

(51) Int. Cl.
A61K 39/12 (2006.01)
A61K 39/21 (2006.01)
CI2Q 1/70 (2006.01)
C07K 14/005 (2006.01)
CI2N 15/86 (2006.01)
CI2Q 1/68 (2006.01)

(52) U.S. Cl.
CPC *CI2Q 1/703* (2013.01); *C07K 14/005*

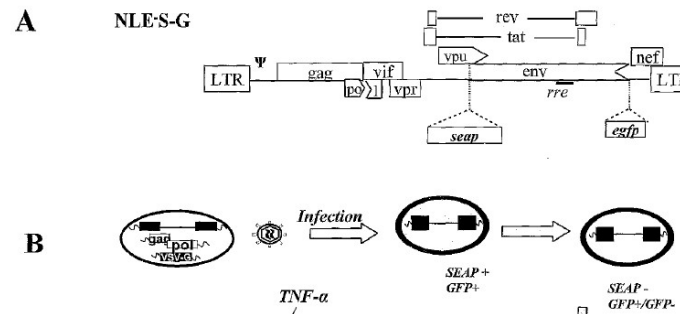
Structures at the Viral Long Terminal Repeat Drives the Progressive Entry of HIV into Latency. *J. Virol.* 2008; 82(24): 12291-12303.*
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Primary Examiner — Zachariah Lucas
Assistant Examiner — Stuart W Snyder
(74) Attorney, Agent, or Firm — Hoffmann & Baron, LLP

(57) ABSTRACT

Isolated, latently infected T cell lines are provided that can be utilized in high throughput screening to discover compounds capable of activating HIV-1. The T cell lines harbor a latent HIV-1 derived vector pro virus, which upon activation expresses a marker for late viral gene expression due to the insertion of the marker gene in the position of HIV-1 envelope.

10 Claims, 8 Drawing Sheets



Patent Example

What is claimed is:

1. An isolated, latently infected T cell line harboring a latent HIV-I derived vector pro virus, which upon activation of the pro virus expresses a secretable marker for late viral gene expression, the gene for said marker being inserted in the position of HIV-I envelope.

2. The cell line of claim 1, wherein the marker is a secretable enzyme.

3. The cell line of claim 2, wherein the secretable enzyme is alkaline phosphatase.

4. The cell line of claim 3, wherein the secreted alkaline phosphatase is capable of being detected using chemiluminescence.

5. The cell line of claim 1, wherein the pro virus further includes a gene marker for viral early gene expression at the single cell level.

6. The cell line of claim 5, wherein the expressed marker for viral early gene expression is a fluorescent protein.

20 7. The cell line of claim 6, wherein the expressed marker for viral early gene expression is enhanced green fluorescent protein (egfp).

25 8. The cell line of claim 1, wherein the provirus is capable of being activated by stimuli selected from the group consisting of tumor necrosis factor (TNF)- α , phorbol 12-myristate 13-acetate (PMA), valporic acid and combinations thereof.

9. The cell line of claim 1, wherein the provirus contains an intact HIV-I gag gene, thereby allowing the use of Gag expression as a further marker of viral gene expression.

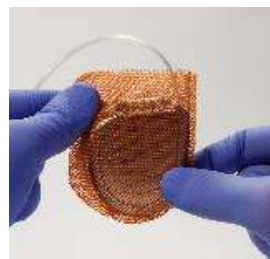
30 10. The cell line of claim 1, wherein the latent provirus in the cell line is replication-incompetent.

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