John M. Garvey, Ph.D., Associate, Greenberg Traurig, Boston, MA

Dr. Garvey works with pharmaceutical and biotechnology clients, primarily those companies engaged in drug discovery, genomics, proteomics, small molecule drug design, immunology and vaccine technologies, drug delivery, gene therapy technologies, stem cell technologies and biomedical devices. His practice includes advising clients on the creation and development of patent portfolios, and their strategic use in corporate collaborations and financial transactions, as well as patent validity, freedom to operate and infringement opinions. He also supports the firm's litigation and patent interference practices.

Dr. Garvey has authored numerous articles on patent law, technology transfer, and the development of intellectual property portfolios for biotechnology companies. Most recently, aspects of his scientific work, originating out of a collaboration with researchers at the University of Birmingham, U.K., were presented at the 9th Annual Human Antibodies and Hybridomas Conference in Berne, Switzerland.

Education

- Ph.D., Molecular and Cellular Biology, Brandeis University, 2000
 - J.D., Franklin Pierce Law Center, 1993
- B.S., Fairfield University, 1990

Admitted to Practice

Massachusetts

U.S. Patent and Trademark Office

William LeePh.D., President & CEO, Cofounder, eMembrane, Inc.

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Dr. Lee received his B.Eng., M.Eng. and Ph.D. from the University of Tokyo. Following a post-doctoral program at Harvard Medical School/Massachusetts General Hospital/Shriners Research Center, he joined JAFCO, Japan's largest venture capital firm, as an Assistant Investment Officer for the Life Science Division. Shortly after his career in venture capital, Dr. Lee returned to Boston and cofounded eMembrane, Inc. in Cambridge, MA based on his and his advisors' years of research work. eMembrane is a US company developing a broad spectrum of multi-functional polymeric materials and membranes for chemical and biological applications. The Company's "Made in Japan" technology has been funded by Asian investors to target multiple global markets with diverse applications.

Michel Morency, Ph.D., Shareholder, Greenberg Traurig, LLP, Boston, MA

Dr. Morency works with pharmaceutical and biotechnology clients, primarily those companies engaged in drug discovery, neurobiology, genomics, proteomics, small molecule drug design, immunology and vaccine technologies, drug delivery, gene therapy technologies, stem cell technologies and biomedical devices. Mike advises clients on all aspects of patent strategy, including the creation and management of patent portfolios as well as the evaluation of both client and competitor portfolios, preparation of validity and infringement opinions, and freedom-to-operate opinions. His practice also involves the acquisition and exploitation of intellectual property rights through licensing and strategic collaboration agreements. In addition, Mike represents companies and investors with regards to intellectual property assets and the formation and funding of ventures built around technology portfolios.

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- Member, American Intellectual Property Law Association
- Member, Patent and Trademark Institute of Canada and a support of the application

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Education in the second s

LL.B. (J.D.), cum laude, University of Ottawa in Ottawa, Canada, 1994

Author of several legal and scientific publications many lever interval and an authority.

• Ph.D., Medical Sciences-Neurosciences, McMaster University in Hamilton, Canada, 1992

B.S., with honors, Life Sciences, Queen's University at Kingston, Canada, 1984

Admitted to Practice

Massachusetts

United States Patent and Trademark Office

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- Weak infrastructure
- Highly dependent on imports

 40%-60% in 1990s
- Market for domestic biotechnology products should reach \$7.5 billion by 2005
- Government committed to foster Biotech as nation's strategic industry
- Aims to be world's G-7 level in Biotech by 2010

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Downside Protection (Exclusivity-1)

- Priority of this Agreement. In the event KJK seeks to grant any license under the KJK Patent Rights or any sublicense under the JAERI Patent Rights to any manufacturer or seller of XXXX Products (a "Third Party License"), and where the Third Party License may permit such third party licensee to manufacture or sell XXXX Products within the scope of the license granted in this Agreement, then KJK will: - promptly provide written notice to eMembrane that it is
- promptly provide written notice to eMembrane that it is negotiating such Third Party License, such notice detailing the proposed royalty rates and terms of the Third Party License, and
 extend to eMembrane the opportunity to negotiate with KJK, in good faith, for a similar license (a "Supplemental License"), such that the Supplemental License taken in combination with this Agreement shall afford eMembrane rights of equivalent scope and equivalent terms with respect to the Third Party License.



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Downside Protection (Exclusivity-2)

 <u>Equivalency of Terms</u>. If any royalty rates or terms in the Third Party License are more favorable to the third party licensee than the royalty rates or terms in this Agreement are to eMembrane, then irrespective of whether eMembrane obtains such Supplemental License, KJK shall extend to eMembrane under this Agreement the more favorable royalty rates or terms, effective as of the date on which they became effective with respect to the Third Party License. For the avoidance of doubt, this includes the royalties or terms applicable to any Project Plans.



Downside Protection (Exclusivity-3) KJK shall not enforce any KJK Patent Rights or JAERI Patent Rights against eMembrane, its Affiliates, or any third party manufacturer of XXXX Products if eMembrane elects to manufacture or have manufactured any XXXX Products under Section 3.1 or 3.2, and will indemnify and hold harmless eMembrane, its Affiliates or third party manufacturers from any such infringement suit brought by JAERI to enforce the JAERI Patent Rights.

Japanese-Style Provisions-1

<u>Dispute Resolution</u>. The Parties recognize that disputes as to certain matters may from time to time arise which relate to either Party's rights and/or obligations hereunder. The Parties hereby agree that they will attempt in good faith to resolve any controversy, claim or dispute (collectively, a "Dispute") arising out of or relating to this Agreement promptly by negotiations. Any such Dispute which is not settled by the Parties which rifteen (15) days after notice of such Dispute is given by one Party to the other in writing shall be referred to a senior executive of eMembrane and of KJK who are authorized to safel such Disputes in the respective companies ("Senior Executives") and who, if possible, are not involved in the Dispute. The Senior Executives will meet for negotiations within fifteen (15) days after ned of the 15-day negotiation period referred to above, at a time and place mutually acceptable to both Senior Executives. The Dispute has not been resolved within thirty (3) days after the end of the 15-day negotiation period referred to above (which period may be extended by mutual agreement), the other Party can immediately bring an action relating to the Dispute before a court of competent jurisdiction.



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