MATERIAL TRANSFER AGREEMENT FOR THE TRANSFER OF ORGANISMS (MTA-TO) to Academic/ Not-for-Profit Organizations

Provider:	National Institute of Diabetes and Digestive and Kidney Diseases
Provider Scientist:	
Recipient:	
Recipient Scientist:	
This is in response to RECIPIENT's request for the MATERIAL (specifically, the name of the gene or allele mutation that makes the organism(s) unique)	
MATERIAL/Organism, Strain or Species:	
and any unmodified derivative and unmodified progeny, as well as any biological materials (including, without limitation: zygotes, embryos, cells, tissues, fluids, etc.) which contain or incorporate the MATERIAL and are derived directly from the original organism or its unmodified progeny, to be used for the purpose of (<i>provide a brief description of the research using additional pages if necessary</i>):	

The PROVIDER requires that the RECIPIENT agree to and the RECIPIENT SCIENTIST acknowledge the following terms before the RECIPIENT receives the MATERIAL:

- 1. The above MATERIAL is the property of the PROVIDER and is made available as a service to the research community.
- 2. THIS MATERIAL IS NOT FOR USE IN HUMAN SUBJECTS.
- 3. The MATERIAL will be used for teaching or not-for-profit research purposes only.
- 4. The MATERIAL will not be further distributed to others who are not under the RECIPIENT SCIENTIST's direct supervision without the PROVIDER's written consent. The RECIPIENT shall refer any request for the MATERIAL to the PROVIDER. To the extent supplies are available, the PROVIDER or the PROVIDER SCIENTIST agree to make the MATERIAL available, under a separate Material Transfer Agreement for the Transfer of Organisms to other scientists for teaching or not-for-profit research purposes only.
- 5. The RECIPIENT agrees to acknowledge the source of the MATERIAL in any publications reporting use of it.
- 6. Any MATERIAL delivered pursuant to this Agreement is understood to be experimental in nature and may have hazardous properties. THE PROVIDER MAKES NO REPRESENTATIONS AND EXTENDS NO WARRANTIES OF ANY KIND, EITHER EXPRESSED OR IMPLIED. THERE ARE NO EXPRESS OR IMPLIED WARRANTIES OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE, OR THAT THE USE OF THE MATERIAL WILL NOT INFRINGE ANY PATENT, COPYRIGHT, TRADEMARK, OR OTHER PROPRIETARY RIGHTS. Unless prohibited by law, RECIPIENT assumes all liability for claims for damages against it by third parties which may arise from RECIPIENT's use, storage or disposal of the MATERIAL except that, to the extent permitted by law, the PROVIDER shall be liable to the RECIPIENT when the damage is caused by the gross negligence or willful misconduct of the PROVIDER.
- 7. The RECIPIENT agrees to use the MATERIAL in compliance with all applicable statutes and regulations.
- 8. If the RECIPIENT anticipates that it will generate cross-bred or genetically-modified organisms incorporating the PROVIDER's modified allele(s), RECIPIENT may transfer such cross-bred or genetically-modified organism(s) to non-profit institutions under the terms of a material transfer agreement that notifies the not-for-profit institution of the existence of PROVIDER's rights to the modified allele(s) and restricts the use of the transferred organism(s) by the not-for-profit recipient

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party has any patent, invention or other intellectual property rights in the organism(s), these rights are not transferred to RECIPIENT by PROVIDER. 9. If NIH is the PROVIDER, the following checked addenda are attached and are incorporated by reference (check all that apply): [] Cre-Lox Addendum, [] OncoMouse® Addendum, [x] Animal Transfer Addendum, or [] Animal Transfer Agreement (not required for transfers within NIH), [] Other The PROVIDER, RECIPIENT and RECIPIENT SCIENTIST must sign both copies of this letter and return one signed copy to the PROVIDER. The PROVIDER will then send the MATERIAL. PROVIDER INFORMATION and AUTHORIZED SIGNATURE: Provider Scientist: National Institute of Diabetes and Digestive and Kidney Diseases Provider Organization: Address: 9000 Rockville Pike, Building 12A, Room 3011 Bethesda, MD 20892-5632 Name of Authorized Official: Rochelle S. Blaustein, J.D. Title of Authorized Official: Director NIDDK, Office of Technology Transfer and Development Signature of Authorized Official Date RECIPIENT INFORMATION and AUTHORIZED SIGNATURE: Recipient Scientist: Recipient Organization: Authorized Signature for Recipient's Institution Name of Authorized Signatory: Title of Authorized Signatory: Certification of Recipient Scientist: I have read and understood the conditions outlined in this Agreement and I agree to abide by them in the receipt and use of the MATERIAL. Date: __ Recipient Scientist Signature Name of Recipient Scientist: Title of Recipient Scientist: Recipient's address for documents: E-mail for documents: Recipient Scientist's address for materials:

to teaching or not-for-profit research purposes only. This Agreement does not transfer any of PROVIDER's patent, invention, or other intellectual property rights in the organism(s) to RECIPIENT. Additionally, to the extent that any other

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E-mail for Recipient Scientist:

DUPONT ADDENDUM FOR USE IF NIH IS THE PROVIDER

NIH has entered into certain Memoranda of Understanding (MOUs) with DuPont, in which DuPont agrees that NIH may receive, use, and transfer mice containing Cre-Lox or OncoMouse® technologies to other entities for research purposes. The MOUs state that NIH provide the following respective notices when shipping such mice.

Cre-Lox Technology:

The above MATERIALS are provided to the RECIPIENT with the knowledge of "Dupont Patent Rights" (U.S. Patent 4,959,317 and any patents granted on any divisional and continuation applications thereof) that have been assigned to DuPont, under the following conditions:

- a) The RECIPIENT may use the MATERIAL, and any progeny or derivatives containing cre DNA and/or lox DNA derived directly or indirectly therefrom, for its internal noncommercial research purposes only, provided however, that such research purposes specifically excludes (i) any activity associated with higher plants or agricultural applications and (ii) the alteration of mouse embryonic stem cells or other pluripotential mouse cells for the purpose of preparing a library of such mouse embryonic stem cells or other pluripotential mouse cells containing cre DNA and/or lox DNA. The MATERIAL, and any progeny or derivatives containing cre DNA and/or lox DNA derived directly or indirectly therefrom, will not be used for any commercial purpose or for the direct benefit of any for-profit institution (except as may be permitted under a written agreement between the non-profit institution and DuPont).
- b) The MATERIAL, and any progeny or derivatives containing cre DNA and/or lox DNA derived directly or indirectly therefrom, may not be transferred by the RECIPIENT to any third parties (except as may be permitted under a written agreement between the RECIPIENT and DuPont).
- c) With respect to further license rights under U.S. patent number 4,959,317, the RECIPIENT should contact:

Debra Desmond Bristol Myers Squibb Company Route 206 & Province Line Road EST&L, Mail Stop B24-05 Princeton, NJ 08543-4000

Tel: 302-467-9431 Fax: 609-252-7337

Email: debra.desmond@bms.com

OR Drew E. Van Dyk

Associate Director, Commercial Development E.I. du Pont de Nemours and Company Rt. 141 and Henry Clay

Wilmington, DE 19880-0268 Voice: (302) 695-3538 Fax: (302) 355-2831

Cell: (302) 354-4943

drew.e.van-dyk@usa.dupont.com

Oncomouse® Technology:

The above MATERIALS are provided to RECIPIENT with knowledge of "Dupont Patent Rights" (U.S. Patent 4,736,866, and 5,087,571, and 5,925,803 and corresponding foreign patents, and any patents granted on any divisional and continuation applications thereof (collectively known as "Harvard Patent Rights")), and that have been exclusively licensed to Dupont by Harvard University, under the following conditions:

- a) The RECIPIENT institution may use the MATERIAL for its internal noncommercial research purposes only. The MATERIAL will not be used for any commercial purpose or for the direct benefit of any for-profit institution (except as may be permitted under a written agreement between the RECIPIENT and DuPont). Accordingly and without limitation, the RECIPIENT is not permitted under this Material Transfer Agreement to use any MATERIAL to test compounds for any commercial purpose or for the direct benefit of any for-profit institution or use the MATERIAL for the production of products for any commercial purpose or for the direct benefit of any for-profit institution.
- (b) The MATERIAL may not be transferred by the RECIPIENT to any third parties (except as may be permitted under a written agreement between the RECIPIENT and DuPont).
- c) The RECIPIENT is notified by PHS of the existence of Harvard Patent Rights and the exclusive license thereof to DuPont, and that the restrictions set forth under (a) and (b) above shall exist only during the term of the Harvard Patent Rights.
- d) With respect to further license rights under the Harvard Patent Rights, the RECIPIENT should contact:

Greg Townsend

DuPont Intellectual Assets and Licensing

Chestnut Run Plaza 708/179 Wilmington, DE 19880-0708 USA

Tel: 877-881-9787 Fax: 302-999-4149

Email: <u>j-gregory.townsend@usa.dupont.com</u>

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ANIMAL TRANSFER ADDENDUM FOR USE IF NIH IS THE PROVIDER

The terms of this Addendum are directed to the use and transfer of the tangible animal(s).

For transfers of any live animals from NIH to outside laboratories, a fully signed National Institutes of Health Animal Transfer Agreement should be completed by the parties prior to the actual shipment of the requested animal. The Animal Transfer Agreement has been adopted for use by the National Institutes of Health (NIH) for use in transferring animals for research purposes pursuant to Section 301 of the Public Health Service Act.

The PROVIDER agrees to transfer the animal(s) described in this Agreement to the RECIPIENT.

RECIPIENT agrees to use the animal(s) solely in connection with biomedical or behavioral research.

Relevant documents concerning the medical history, health status, and research uses of the animal(s), including prior surgical procedures and any infectious disease (human or zoonotic) to which the animal(s) may have been exposed, will be provided in a separate document.

For domestic recipients, RECIPIENT agrees that it will comply with the Animal Welfare Act and its implementing regulations, as applicable. RECIPIENT agrees that it will adhere to all applicable national standards for humane care and use of the animal(s) and assures the PROVIDER that it has appropriate animal care and use policies in place. The "Public Health Service Policy on Humane Care and Use of Laboratory Animals" and "Guide for the Care and Use of Laboratory Animals" are examples of acceptable standards for humane care and use of research animals.

RECIPIENT agrees that it will adhere to appropriate biosafety practices and use the animals in a safe and responsible manner. The National Institutes of Health/Centers for Disease Control publication, "Biosafety in Microbiological and Biomedical Laboratories" is an example of acceptable standards for biosafety practices. RECIPIENT agrees that it will comply with applicable import/export regulations.

In accepting the animal(s), RECIPIENT accepts full ownership, custody, and control of the animal(s), except that to the extent the Government has any patent, invention or any other intellectual property rights in the animal(s), the Government retains these rights, as described in the attached Material Transfer Agreement for the Transfer of Organisms.

PROVIDER is transferring the animal(s) as a service to the research community. The animal(s) is transferred to the RECIPIENT with no warranties, express or implied, including any warranty of merchantability or fitness for a particular purpose. Unless prohibited by law from doing so, RECIPIENT agrees to hold the United States Government harmless and to indemnify the Government from all liabilities, demands, damages, expenses and losses arising out of the RECIPIENT's care, use or treatment of the animal(s).

RECIPIENT agrees not to claim, infer, or imply Governmental endorsement of the RECIPIENT, the research project, the institution or personnel conducting the research, or any resulting product(s).

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