

## MATERIALS TRANSFER AGREEMENT

**Provider:**

U.S. EPA / ORD / National Center for Computational Toxicology (NCCT)

**Recipient:**

North Carolina State University

1. Provider agrees to transfer to Recipient's Investigator named below the following Research Material:

**Chemicals and Materials**

- A list identifying selected chemicals from the ToxCast library to be tested by Seth Kullman.
- A copy of the current ToxCast chemical library consisting of chemical samples prepared as solution in dimethyl sulfoxide at a concentration of 20 millimolar. Additional chemicals may be provided in the future concurrent with expansion of the ToxCast chemical library.
- Samples of nanomaterials and characterization data on said materials

**Data and Summary Information**

- In vitro assay data derived from the ToxCast Program. This data is derived from chemicals analyzed using a variety of high throughput assay techniques. Below this is referred to as the "ToxCast Data".
- Summary descriptions of the individual data sets.
- Individual subsets of this data will be delivered to Seth W. Kullman after they have been prepared for use at EPA and cleared for release to North Carolina State University.

Recipient agrees to transfer to the EPA Investigator named below:

- All data resulting from chemical screening performed on the ToxCast chemical library.
- Relevant data on these chemicals from non-public sources.
- Unique chemicals for the ToxCast chemical library and subsequent testing.

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2. This Research Material may not be used in human subjects. The Research Material will be used only for research purposes by Recipient's investigator in his/her laboratory, for the research project described below, under suitable containment conditions. This Research Material will not be used for screening, production or sale, for which a commercialization license may be required. Recipient agrees to comply with all Federal rules and regulations applicable to the Research Project and the handling of the Research Material.

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3. Does the Research Material include specimens or data derived or collected from human subjects?

☐ Yes – Go to item #3(a).

☒ No – Skip to item #4.

3(a). Does the Research Material include specimens or data derived or collected from fetuses, children, pregnant women, or nursing women?

☐ Yes

☐ No

3(b). Was the Research Material obtained under a protocol that was reviewed and approved by an Institutional Review Board (IRB) that operated in accordance with the requirements of EPA Regulation 40 CFR 26, HHS Regulation 45 CFR 46, or any other Federal Regulation for the protection of human research subjects?

☐ Yes (Please indicate the applicable Regulation here \_\_\_\_\_ and provide copies of the protocol and IRB approval documents.)

☐ No (Please provide explanation with documentary support as appropriate.)

3(c). Can the Provider of the Research Material identify the subjects directly or through identifiers (codes) linked to the subjects?

☐ Yes – The Recipient's use of the Research Material may be human subject's research subject to 40 CFR 26. Go to item #3(d).

☐ No – The Recipient's use of the Research Material is not human subjects research subject to 40 CFR 26. Skip to item #4.

3(d). Is the Provider of the Research Material prohibited by this agreement from releasing information to the Recipient that might allow the identification of any of the subjects, including but not limited to the key to any existing code?

☐ Yes – The Recipient's use of the Research Material is not human subjects research subject to 40 CFR 26. Skip to item #4.

☐ No – The Recipient's use of the Research Material may be human subjects research subject to 40 CFR 26. Go to item #3(e).

3(e). Is the Research Material publicly available?

☐ Yes – The Recipient's use of the Research Material is human subjects research that is exempt from 40 CFR 26.

☐ No – The Recipient's use of the Research Material is human subjects research that may be subject to 40 CFR 26 and must be further evaluated accordingly by the EPA Human Subjects Review Official.

4. This Research Material will be used by Recipient's investigator solely in connection with the following research project "*Targets for Developmental Neurotoxicity*" described with specificity as follows:

Our interests are to investigate the environmental and human health consequences of Nuclear Receptor (NR) agonists and antagonists using a comparative approach with both zebrafish and medaka, small aquarium fish models of human disease. Our primary focus is to identify compounds that exhibit developmental neurotoxicity through dysregulation of nuclear receptor targets including the vitamin D receptor (VDR), Nurr1, and other nuclear receptors associated with early brain and CNS development. Current, phase one ToxCast data suggests several compounds (pesticides) that target these NR pathways. As the ToxCast phase two data set is greatly expanded, we will use this information to enhance our list of chemicals that may target NR function through these important pathways. Based upon the compilation of phase one and phase two data we will establish a priority list of chemicals to initiate a developmental neurotoxicity screen comprised of both in vitro and in vivo assays deploying , mammalian cell lines, and medaka and zebrafish developmental models.

5. In all oral presentations or written publications concerning the Research Project, Recipient will acknowledge Provider's contribution of this Research Material unless requested otherwise. To the extent permitted by law, Recipient agrees to treat as confidential, any of Provider's written information about this Research Material that is stamped "CONFIDENTIAL" for a period of three (3) years from the date of its disclosure to recipient. The foregoing shall not apply to information that is or becomes publicly available, which is disclosed to Recipient without a confidentiality obligation, which is independently developed, which is known to the Recipient prior to disclosure, or is disclosed pursuant to law or judicial order. Any oral disclosures from Provider to Recipient which Provider wishes to be treated as confidential shall be identified as being Confidential at the time of the disclosure and by written notice delivered to Recipient within thirty (30) days after the date of the oral disclosure. Recipient may publish or otherwise publicly disclose the results of the Research Project, but if Provider has given Confidential information to Recipient, such public disclosure may be made only after Provider has had thirty (30) days to review the proposed disclosure to determine if it includes any Confidential information, to the extent such review period is permitted by law.

6. This Research Material represents a significant investment on the part of Provider and is considered proprietary to Provider. Recipient's investigator therefore agrees to retain control over this Research Material and further agrees not to transfer the Research Material to other people not under his/her direct supervision without advance written approval of Provider. Provider reserves the right to distribute the Research Material to others and to use it for its own purposes. When the Research Project is completed, the Research Material will be returned to the Provider or disposed, if directed by Provider. Recipient shall be entitled to retain one archival copy of all Confidential Information in order to meet its obligations under this agreement and to meet any record retention laws or policies of North Carolina.

7. This Research Material is provided as a service to the research community. It is being supplied to Recipient with no warranties, express or implied, including any warranty of merchantability or fitness for a particular purpose. Provider makes no representations that the use of the Research Material will not infringe any patent or proprietary rights of third parties.

8. Recipient shall retain title to any patent or other intellectual property rights in inventions made by its employees in the course of the Research Project. However, if said inventions contain any portion of the Research Material, are derived from the Research Material, or could not have been produced but for the use of the Research Material, Recipient agrees to contact the Provider to determine what ownership interests, if any, the Provider may have, and, where applicable, to negotiate in good faith the terms of a commercial license. Inventorship for a patent application or a commercialized product based on said inventions shall be determined according to United States patent law.

9. When Provider is the EPA: Recipient agrees not to claim, infer, or imply endorsement by the Government of the United States of America (hereinafter referred to as "Government") of the Research Project, the institution or personnel conducting the Research Project or any resulting product(s). Recipient agrees, to the extent allowed under law, to hold the Government harmless and to indemnify the Government for all liabilities, demands, damages, expenses and losses arising out of Recipient's use for any purpose of the Research Material.

10. When Recipient is the EPA: Provider will not be liable to EPA for any claims or damages arising from EPA's use of the Research Material.

11. The Provider shall have the right to terminate this Agreement at any time if Recipient breaches any of the terms of this Agreement. Upon termination, Recipient shall return to the Provider all unused portions of the Research Materials.

12. Will EPA develop any products or services from information or materials provided by the Recipient?

☐ Yes – go to item A

☒ No – skip to #13 (next clause)

Item A: The EPA has a long history of applying principles of quality assurance/quality control to all technical work conducted by or for the Agency (CIO 2106: USEPA Quality Policy). Given EPA is receiving metabolomics and screening data and will use the metabolomics and screening data for Agency purposes, the Recipient is required to provide EPA with documentation such as a quality manual, describing their organization's quality system. In lieu of such documentation, Standard Operating

Protocols for compound handling and the assays performed are acceptable or documentation showing third party accreditation to a relevant standard and scope is also acceptable for documenting an organization's quality system. EPA requirements for quality management plans can be found at this URL:  
[http://www.epa.gov/quality/qa\\_docs.html](http://www.epa.gov/quality/qa_docs.html)

13. All notices pertaining to or required by this Agreement shall be in writing and shall be signed by an authorized representative and shall be delivered by hand (including private courier mail service) or sent by certified mail, return receipt requested, with postage prepaid, addressed as follows:

**Provider's Administrative Contact Information:**

Rebecca Clausen  
National Center for Computational Toxicology (NCCT)  
US EPA  
109 TW Alexander (MD-B-205-01)  
Research Triangle Park, NC 27711  
For commercial courier address use:  
4930 Old Page Rd.  
Durham, NC 27703  
919-541-3002  
Clausen.Rebecca@epa.gov

**Recipient's Contact Information:**

Rob Whitehead, Director of Licensing

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North Carolina State University  
Raleigh, NC 27695-8210

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14. Paragraphs 2, 7, 9 and 10 shall survive termination.

15. (Deleted)

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Any false or misleading statements made, presented, or submitted to the Government, including any material omissions, under this Agreement and during the course of negotiation of this Agreement are subject to all applicable civil and criminal statutes including 31 U.S.C. ' ' 3801-3812 (civil liability), 18 U.S.C. ' 1001 (criminal liability), and 31 U.S.C. ' ' 3729-33 (False Claims Act).