

## MATERIALS TRANSFER AGREEMENT

**Provider:**

U.S. Environmental Protection Agency (EPA)  
Office of Research and Development (ORD)  
National Center for Computational Toxicology (NCCT)

**Recipient:**

Craig Beeson, President, MitoHealth, Inc  
645 Meeting Street, Suite 11  
Charleston, SC 29403

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

Chemicals and Materials

- X 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.

Data and Summary Information

- X 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".

Recipient agrees to transfer to the EPA Investigator named below:

- X All data resulting from chemical screening performed on the ToxCast chemical library.

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.

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 ("Research Project") described with specificity as follows (*insert description here or use an attachment page if necessary*):



We plan to use the ToxCast phase I and II chemical libraries in our mitochondrial respirometric assay. The use of ToxCast is specifically for Aim 1 of the research project entitled "High Throughput Mitochondrial Nephrotoxicant Assay". The kidney is a target of toxicity for drugs, industrial and environmental chemicals because of its high blood flow, numerous transporters, and reliance on aerobic metabolism. Not surprisingly, mitochondria are a common intracellular target of chemicals in the kidney, damage of which leads to decreased aerobic metabolism and ATP, and cell death. These critical mitochondrial functions make them a superb proxy for cell and organ vitality. Current *in vitro* models of mitochondrial function are inadequate because cultured cells are highly glycolytic with minimal aerobic metabolism. In light of the fact that many known nephrotoxicants also have documented risks to mitochondria, the development of a high-throughput assay to measure loss of mitochondrial respiratory capacity in renal tissue will greatly enhance efforts to assess new chemicals and biological agents as potential nephrotoxicants and fill in the gap present in other high-throughput screening (HTS) efforts.

*Aim 1. The integrated metabolic and imaging assay will be used to screen the ToxCast Phase I and Phase II compounds.*

*Aim 2. Cheminformatic analysis of all the molecules will be executed and toxicophores identified. Hits will be cross-validated against the ToxCast animal testing and human exposure data.*

This RPTC/XF-96 platform will identify nephrotoxicants and mitochondrial toxicants within the ToxCast phase I and phase II libraries. This information will provide public companies and regulatory agencies with mechanisms and chemical-based criteria for assessing and predicting nephrotoxicity and mitochondrial toxicity of new drugs, consumer products, and environmental agents.

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 or which is disclosed to Recipient without a confidentiality obligation. 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.

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 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](mailto:Clausen.Rebecca@epa.gov)

**Recipient's Contact Information:**

Craig Beeson, President  
  
645 Meeting Street, Suite 11, Charleston, SC 29403

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).