

UNFUNDED AGREEMENT REQUEST QUESTIONNAIRE

Providing the following information will assist the Sponsored Programs Administration (SPA) to review and negotiate requested unfunded agreement with less interference and more efficiency and effectiveness. **NOTE:** Not providing a reply to all questions, even if “No” or “N/A”, may cause additional inquiry from SPA.

UMB Principal Investigator

Name:	
Study Team Contact Name & Email Address	

Collaborator

Please enter the name and email address of the POC. This is the person responsible for reviewing & executing the Agreement and may differ from the Principal Investigator (PI).

PI Name:	
*Point of Contact - Name & Email Address	
Physical Address:	

Type of Request:

Data Use/Transfer (DUA) **Non-Disclosure (NDA)/ Confidentiality Disclosure (CDA)** **TEAMING**
Domestic Memorandum of Understanding (MOU) ***MASTER** **Business Associate (BAA)** **EQUIPMENT**
Collaboration/Cooperative (CRA) **OTHER; Explain: _____**

*If MASTER; will related agreement(s) follow? Yes No

if Yes to above, select which will be determined in each agreement: IP Informed Consent IRB/IACUC

Material Transfer – **STOP HERE**; submit MTA request to CCT via UMBiz

International MOU – **STOP HERE**; submit request to International Operations (Global Hub)

Commercial collaboration – **STOP HERE**; submit request to OGC

Incoming Visiting Scientist – **STOP HERE**; submit request to Janet Simons, SPA

Agreement Purpose

If scope or purpose for agreement is not already stated in agreement, please explain. **NOTE:** For DUAs do not enter description here; it will be asked in the DUA Project Description section; page 2)

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Agreement Template Source:

UMB’s Template **Collaborator’s Template** If Collaborator, is the Agreement uploaded in KR/KB? Yes No

Related Sponsored Funding

Complete if there is any sponsored funding associated with the requested Agreement or referenced data.

Funding Source: Sponsored UMB Internal Funds

If Sponsored, enter KR award info: _____

UMB Project ID#: _____

eUMB# (if applicable): _____

Intellectual Property

Do you anticipate new intellectual property (patents/copyrights) will be developed in performance of Agreement? Yes No

Is there a reasonable possibility of commercial utility? Yes No

Is background IP included or involved? Yes No; if Yes, what is the IP reference# or title: _____

Publishing

Will any publications result from the purpose of requested Agreement? Yes No

If Yes, will this be a joint publication? Yes No

Confidential Information

In the Agreement requested, will Confidential Information be transferred between parties?

UMB Confidential Information? Yes No

External Entity’s Confidential Information? Yes No

Regulatory Compliance

(NOTE: **SKIP** this section, if the following is accurate & up-to-date in Quali Research)

Does the purpose for this Agreement involve:

Human Subjects? **Yes No**

If yes, IRB protocol #: _____

Status of IRB review: **Approved Pending**

Primary IRB: **UMB Collaborator Both**

Participant consent required? **Yes No**; If Yes, obtained? **Yes. No**

If No to obtained, briefly explain: _____

If the primary IRB is outside UMB, provide the collaborator's protocol reference #: _____

Use of Vertebrate Animals: **Yes No**

If yes, IACUC Protocol # _____

IACUC location: _____ Status of IACUC review: **Approved Pending**

Proposed Duration of the Agreement: _____

FOR DUA REQUEST ONLY; COMPLETION OF FOLLOWING SECTIONS ARE ALSO REQUIRED

Is UMB: **Data Recipient Data Provider Both**

Who Owns the Data: **UMB Collaborator UMMS** Other; if Other: _____

DUA Project Description / Justification for Use

This section should provide sufficient information such that each party understands the project that the Recipient will perform using the Data. Examples of information include: Objectives, purpose of the Recipient's work, or a general description of the actions to be performed by the Recipient using the Data and possibly the anticipated results.

NOTE: If a Justification for Use has been submitted to the Data Provider, do not complete the section above and upload justification submitted to sponsor into request routed (e.g., KR/KB).

Source & Type of Data (check all that apply):

(For information on data types: https://thefdp.org/wp-content/uploads/human_subject_data_classification_tool.pdf)

De-identified Limited Data Set Personal Health Information (PHI) Personally Identifiable Information (PII)

Electronic Medical Records (EMR) / Electronic Health Record (EHR) Other; please explain: _____

Description of Data

This section should provide sufficient information such that each party understands the information that will be transmitted under this Agreement. Examples of information include: Whether the data is obtained from human subjects and, if so, the number of subjects or a description of the population included in the data; if the data is from animal subjects, the species of animal the data was obtained using; if not from human or animal subjects, a description of the data and/or experiments included. Name of the study that the data was obtained under if there is a particular study that needs to be acknowledged/cited as the source of the data.

If UMB is the Data Provider, are there data management / disposition requirements for the Data Recipient? **Yes No**

If Yes, please explain:

Third Party Permitted / Will any other entity access the data? **Yes No**

If Yes, provide name: _____

Will the data be combined with data from any other sources? **Yes No**

Transmission Method: **Electronically Mail Repository Data Coordinating Center**

Data transferred across international borders? **Yes No**

Does the data involve personal data of a citizen or resident living in the European Economic Area or the European Union? **Yes No**

Cost associated with data transfer: **Yes No**

Length of time for use of Data: _____