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| **ETSU/Ballad Collaborative Research****Reciprocal Data Transfer and Use Agreement (“Agreement”)**Personally Identifiable Information – HIPAA |
| Party 1: INSERT ORGANIZATION NAME | Party 2:INSERT ORGANIZATION NAME |
| Party 1 ScientistName: INSERT NAME Email: INSERT EMAIL  | Party 2 ScientistName: INSERT NAME Email: INSERT EMAIL |
| Party 1 Data Type: Personally Identifiable Information – HIPAA[This template includes language and attachments for use when the data being shared is Personally Identifiable Information governed by HIPAA. Make sure you are using the appropriate template.] | Party 2 Data Type: Personally Identifiable Information – HIPAA[This template includes language and attachments for use when the data being shared is Personally Identifiable Information governed by HIPAA. Make sure you are using the appropriate template.] |
| Agreement TermStart Date: INSERT START DATEEnd Date: INSERT END DATE[In general, ETSU is prohibited from entering into agreements that exceed a 5 (five) year term.]  | Project Title: INSERT PROJECT TITLE |
| **Terms and Conditions**1. The Parties shall provide the data set(s) described on Attachment 1 (the “Data”) to each other for the research purpose set forth in Attachment 1 (the “Project”). Each Party is a Providing Party when providing Data and a Receiving Party when receiving Data. Providing Party shall retain ownership of any rights it may have in the Data, and Receiving Party does not obtain any rights in the Data other than as set forth herein.
2. Receiving Party shall not use the Data except as authorized under this Agreement. The Data will be used solely to conduct the Project and solely by Receiving Party’s Scientist and Receiving Party’s faculty, employees, fellows, students, and agents (“Receiving Party Personnel”) and Third-Party Personnel (as defined in Attachment 3) that have a need to use, or provide a service in respect of, the Data in connection with the Project and whose obligations of use are consistent with the terms of this Agreement (collectively, “Authorized Persons”).
3. Except as authorized under this Agreement or otherwise required by law, Receiving Party agrees to retain control over the Data and shall not disclose, release, sell, rent, lease, loan or otherwise grant access to the Data to any third party, except Authorized Persons, without the prior written consent of Providing Party. Receiving Party agrees to establish appropriate administrative, technical, and physical safeguards to prevent unauthorized use of or access to the Data and comply with any other special requirements relating to safeguarding of the Data as may be set forth in the applicable Attachment 2.
4. Receiving Party agrees to use the Data in compliance with all applicable laws, rules, and regulations, as well as all professional standards applicable to such research.
5. The Parties are encouraged to make publicly available the results of the Project. Before either Party submits a paper or abstract for publication or otherwise intends to publicly disclose information about the results of the Project, the other Party will have thirty (30) days from receipt to review proposed manuscripts and ten (10) days from receipt to review proposed abstracts to ensure that the Data is appropriately protected. The non-publishing Party may request in writing that the proposed publication or other discloser be delayed for up to thirty (30) additional days as necessary to protect proprietary information. The Parties will together make decisions on jointly authored publications. Authorship will be in accordance with academic and/or scholarly standards.
6. Receiving Party agrees to recognize the contribution of the Providing Party as the source of the Data in all written, visual, or oral public disclosures concerning Receiving Party’s research using the Data, as appropriate in accordance with scholarly standards and any specific format that has been indicated in Attachment 1.
7. Unless terminated earlier in accordance with this section or extended via a modification in accordance with Section 12, this Agreement shall expire as of the End Date set forth above. Either Party may terminate this Agreement with thirty (30) days written notice to the other Party’s Authorized Official as set forth below. Upon expiration or early termination of this Agreement, Receiving Party shall follow the disposition instructions provided in Attachment 1, provided; however, that Receiving Party may retain one (1) copy of the Data to the extent necessary to comply with the records retention requirements under any law, and for the purposes of research integrity and verification.
8. EXCEPT AS PROVIDED BELOW OR PROHIBITED BY LAW, ANY DATA DELIVERED PURSUANT TO THIS AGREEMENT IS UNDERSTOOD TO BE PROVIDED “AS IS.” PROVIDING PARTY 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 DATA WILL NOT INFRINGE ANY PATENT, COPYRIGHT, TRADEMARK, OR OTHER PROPRIETARY RIGHTS. Notwithstanding, Providing Party, to the best of its knowledge and belief, has the right and authority to provide the Data to Receiving Party for use in the Project.
9. Each Receiving Party shall be liable for damages, losses, claims, and demands which may arise from its use, storage, disclosure, or disposal of the Data except to the extent (a) prohibited by law and/or (b) caused by the negligence, willful misconduct, or violation of applicable privacy or security laws and regulations by the Providing Party.
10. Neither Party shall use the other Party’s name, trademarks, or other logos in any publicity, advertising, or news release without the prior written approval of an authorized representative of that Party. The Parties agree that each Party may disclose factual information regarding the existence and purpose of the relationship that is the subject of this Agreement for other purposes without written permission from the other Party provided that any such statement shall accurately and appropriately describe the relationship of the Parties and shall not in any manner imply endorsement by the other Party whose name is being used.
11. Unless otherwise specified, this Agreement and the below listed Attachments embody the entire understanding between Provider and Recipient regarding the transfer of the Data to Recipient for the Project:
12. Attachment 1: Project Specific Information
13. Attachment 2: Data-specific Terms and Conditions
14. Attachment 3: Identification of Permitted Third Parties (if any)
15. Attachment 4: Additional Terms and Conditions

In the event of any conflict between the obligations set forth in the applicable Attachment 2 and this Agreement, the obligations set forth in Attachment 2 shall prevail.1. No modification or waiver of this Agreement shall be valid unless in writing and executed by duly authorized representatives of both Parties.
2. The undersigned Authorized Officials of the Parties expressly represent and affirm that the contents of any statements made herein are truthful and accurate and that they are duly authorized to sign this Agreement on behalf of their institution.
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| By an Authorized Official of Party 1: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_ Signature DatePrinted Name: INSERT NAMETitle: INSERT TITLEContact Information for Formal Notices:Name: INSERT NAME Address: INSERT ADDRESSEmail: INSERT EMAILPhone: INSERT PHONE | By an Authorized Official of Party 2:\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_ Signature DatePrinted Name: INSERT NAMETitle: INSERT TITLEContact Information for Formal Notices:Name: INSERT NAME Address: INSERT ADDRESSEmail: INSERT EMAILPhone: INSERT PHONE |

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| **Attachment 1**Reciprocal Data Transfer and Use AgreementPersonally Identifiable Information – HIPAAProject-Specific Information |

1. Description of Project:

INSERT DESCRIPTION OF PROJECT

[This section of this attachment should provide sufficient information such that each Party understands the project that the Parties will perform using the Data. Content of this section will be very similar to the Statement of Work used in other types of Agreements. Examples of information that should be provided include:

* Objective or purpose of the Parties’ work.
* A general description of the actions to be performed by each Party using the Data and possibly the anticipated results.
* Whether or not the Parties are permitted to link the Data with other data sets (if yes, be sure to include any special disposition requirements related to the linked data sets in Section 6 and 7 of this attachment).]

2. Description of Party 1 Data:

INSERT DESCRIPTION OF PARTY 1 DATA

[This section of this attachment should provide sufficient information such that each Party understands the information that will be transmitted by Party 1 under this Agreement. If Party 1 will not be sharing any Data under this Agreement, simply indicate “None” in this section and insert “None” in the Party 1 Data Type section on the face page. Examples of information that should be provided include:

* If the Party 1 Data is obtained from human subjects, a description of the population included in the Party 1 Data.
* If the Party 1 Data is from animal subjects, the species of animal the Party 1 Data was obtained using
* If not from human or animal subjects, a description of the focus of the Party 1 Data.
* The number of subjects and/or experiments included.
* Name of the study that the Party 1 Data was obtained under - If there is a particular study that needs to be acknowledged/cited as the source of the Party 1 Data, this information should be included here.]

3. Party 1 Disposition Requirements upon the termination of expiration of the Agreement:

INSERT DISPOSITION REQUIREMENTS

[This section of this attachment should provide sufficient information such that each Party understands the Receiving Party’s obligations with regards to the Party 1 Data upon the expiration or early termination of this Agreement. If the Receiving Party is permitted to link the Data with other data sets, be sure to include any special disposition requirements related to the linked data sets in this attachment.]

4. For Party 1, send Data INSERT METHOD OF TRANSMISSION to:

 Name: INSERT NAME

Address: INSERT ADDRESS

Email: INSERT EMAIL

Phone: INSERT PHONE

5. Description of Party 2 Data:

INSERT DESCRIPTION OF PARTY 2 DATA

[This section of this attachment should provide sufficient information such that each Party understands the information that will be transmitted by Party 2 under this Agreement. If Party 2 will not be sharing any Data under this Agreement, simply indicate “None” in this section and select “None” from the Party 2 Data Type drop-down menu on the face page. Examples of information that should be provided include:

* If the Party 2 Data is obtained from human subjects, a description of the population included in the Party 2 Data
* If the Party 2 Data is from animal subjects, the species of animal the Party 2 Data was obtained using
* If not from human or animal subjects, a description of the focus of the Party 2 Data
* The number of subjects and/or experiments included
* Name of the study that the Party 2 Data was obtained under - If there is a particular study that needs to be acknowledged/cited as the source of the Party 2 Data, this information should be included here.]

6. Party 2 Disposition Requirements upon the termination or expiration of the Agreement.

INSERT DISPOSITION REQUIREMENTS

[This section of this attachment should provide sufficient information such that each party understands the Receiving Party’s obligations with regards to the Party 2 Data upon the expiration or early termination of this Agreement. If the Receiving Party is permitted to link the Data with other data sets, be sure to include any special disposition requirements related to the linked data sets in this attachment.]

7. For Party 2, send Data INSERT METHOD OF TRANSMISSION to:

 Name: INSERT NAME

Address: INSERT ADDRESS

Email: INSERT EMAIL

Phone: INSERT PHONE

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| **Attachment 2**Data Transfer and Use AgreementPersonally Identifiable Information – HIPAAData-specific Terms and Conditions |

**Additional Terms and Conditions:**

1. The Data is Protected Health Information (“PHI”) as that term is defined in the Health Insurance Portability and Accountability Act of 1996 (“HIPAA”), at 45 C.F.R. §160.103 (and not a Limited Data Set).

[ ]  If checked, the Data is covered under a Certificate of Confidentiality, which must be asserted against compulsory legal demands, such as court orders and subpoenas for identifying information or characteristics of a research participant. See <https://grants.nih.gov/grants/guide/notice-files/NOT-OD17-109.html> for further information.

1. Nothing herein shall authorize the Recipient to use or further disclose the Data in a manner that would violate the requirements applicable to Provider under 45 CFR §164.514.
2. Notwithstanding any statement herein to the contrary, Provider represents that it has full authority to share the Data with the Recipient and has confirmed that the Project is consistent with such consents or authorizations, if any, as Provider has obtained from individuals who are the subjects of the Data.
3. Unless otherwise required by law or legal process, Recipient shall not use or further disclose the Data other than as permitted by this Agreement. If Recipient believes it is required by law or legal process to use or disclose the Data, it will promptly notify Provider, to the extent allowed by law, prior to such use or disclosure and will disclose the least possible amount of Data necessary to fulfill its legal obligations.
4. In the event Recipient becomes aware of any use or disclosure of the Data not provided for by this Agreement, Recipient shall take any appropriate steps to minimize the impact of such unauthorized use or disclosure as soon as practicable and shall notify Provider of such use or disclosure as soon as possible, but no later than 5 business days after discovery of the unauthorized use or disclosure. Recipient shall cooperate with Provider to investigate, correct, and/or mitigate such unauthorized use or disclosure. Recipient acknowledges that Provider may have an obligation to make further notifications as set forth in Subpart D of 45 CFR §164 or under applicable state law and shall cooperate with the Provider to the extent necessary to enable Provider to meet all such obligations.
5. Recipient will not use the Data, either alone or in concert with any other information, to make any effort to contact individuals who are the subjects of the Data without appropriate Institutional Review Board (IRB) approval, specific written approval from Provider, and informed consent and authorization from the individual or a waiver, if required.
6. Recipient agrees to implement reasonable safeguards, sufficient to meet the standards of 45 CFR §164.530(c), to limit incidental, and avoid prohibited, uses and disclosures of the Data, and to ensure that only Authorized Persons have access to the Data.
7. Recipient agrees to remove and securely destroy or return, as directed by the Provider in Attachment 1, the part or parts of the Data that identifies the individual who is the subject of the Data at the earliest time at which removal and destruction or return can be accomplished, consistent with the purpose of the Project.
8. The parties agree to take such action as is necessary to amend this Agreement, from time to time, in order for the Provider to remain in compliance with the requirements of the HIPAA Privacy Regulations.
9. By signing this Agreement, Recipient provides assurance that its relevant institutional policies and applicable federal, state, or local laws and regulations (if any) have been followed, including the completion of any IRB review or approval that may be required prior to Recipient’s use of the Data. Upon Provider’s written request to the Recipient’s Contact for Formal Notices identified in the signature block, Recipient shall provide documentation of its IRB-Approved Protocol.

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| **Attachment 3**One-Way Data Transfer and Use AgreementPersonally Identifiable Information – HIPAAIdentification of Permitted Third Parties (if any) |

For all purposes of this Agreement, the definition of “Third Party Personnel” checked below will pertain:

[ ]  “Third Party Personnel” means: None. No collaborators are permitted on the Project.

 -OR-

[ ]  “Third Party Personnel” means as set forth below and agreed upon between the Parties:

[Sample definition language for the drafter; delete if the first option is checked or after a final definition has been agreed between the Parties:

“Third Party Personnel” means: faculty, employees, fellows, or students of INSERT NAME OF THIRD PARTY INSTITUTION, an academic institution, which institution (i) has agreed to collaborate in the Project, (ii) has faculty, employees, fellows, or students who have a need to use or provide a service in respect of the Data in connection with its collaboration in the Project, and (iii) has been made aware of the terms of this Agreement and agreed to comply, and to cause it personnel to comply, with such terms.

An alternative option for (iii): “has executed an agreement that is substantially similar to this Agreement.”]

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| **Attachment 4**Data Transfer and Use AgreementPersonally Identifiable Information – HIPAAAdditional Terms and Conditions |

**Additional Terms and Conditions:**

[ ]  None. No additional terms and conditions are required.

 -OR-

[ ]  The additional terms and conditions are as set forth below and agreed upon between the Parties.

[This section should be completed if the research being conducted includes a grant or other contract. For example: Material Transfer Agreement, Sponsor Agreement, Confidentiality Agreement (e.g. NDA), MOU, Business Associate Agreement, etc. If no additional terms or conditions exist, None should be checked above.]