	<p align="center">Document name: REP-2-020 Attachment E- Patient Consent for Research After Testing</p>	<p align="right">Eurofins Document Reference: 1-P-QM-CF -9061961 NTD Labs SOP ID: REP-2-020 Attachment E Revision:1</p>
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REP-2-020 Attachment E- Patient Consent for Research After Testing

Eurofins Document Reference	1-P-QM-CF -9061961	Type of document	CF - Controlled Form
NTD Labs SOP ID	REP-2-020 Attachment E	Division	1-P Pharma
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Status	Effective	Periodic Review Date	Jul 31, 2020
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Written by	Margaret Palladino
Functional Document Owner	Eurofins D CDS US Quality Management Department; Eurofins D CDS US Reporting Department; Eurofins D CDS US Laboratory Management
Review and Approval	<ul style="list-style-type: none"> • Reviewers: Christina Deer; Jonathan Hayden; Lisa Schmitt • Approver (Laboratory Director Only): Terrence Hallahan
Reason for Revision	Y Chromosome - Research Consent

Revision Log

Date	Rev.	Author	Description
Aug 1, 2018	1	Eurofins D CDS US Quality Management Department; Eurofins D CDS US Reporting Department; Eurofins D CDS US Laboratory Management	Y Chromosome - Research Consent

Electronic Signatures

<p>Christina Deer;Review;Jul 2, 2018 4:04 PM EDT Jonathan Hayden;Review;Jul 3, 2018 8:47 AM EDT Lisa Schmitt;Review;Jul 11, 2018 10:41 AM EDT Terrence Hallahan;Approval;Jul 18, 2018 9:50 AM EDT</p>
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Patient Consent for Research After Testing

Eurofins NTD, LLC performs research to help develop and provide safe and effective screening tests and to contribute to advancing biomedical knowledge. My sample for **Y Chromosome Assessment** will be discarded within sixty days after taken, unless I specifically consent below. Permission to allow the use of my de-identified sample in research and development studies is entirely voluntary.

- I give permission for Eurofins NTD, LLC to retain any remaining de-identified sample for future research and development.
- I do not give permission to use the remaining sample in any research studies.

Patient/Guardian

Signature

Printed

Name

Date ____ / ____ / ____

This “Patient Consent for Research After Testing” form is to be submitted with the sample and requisition form.

<p>Revision: 1 © This document is copyright of Eurofins</p>	<p>Effective date: Aug 1, 2018 During the actual use of this form, cover page need not be printed.</p>	<p align="right">Page 1 of 1</p>
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