



A CLIA Accredited Laboratory  
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PLACE 1 BARCODE ON  
FORM AND 1 ON SAMPLE  
(REQUIRED: NAME/DOB)

CHECKLIST:		
<input type="checkbox"/> Demographics/Medication List	<input type="checkbox"/> ICD-10 Codes	<input type="checkbox"/> ABN (Medicare)
<input type="checkbox"/> Physician & Patient Signatures	<input type="checkbox"/> Copy of Patient Insurance Card	

## Malignant Hyperthermia Test Requisition Form

First Name	Last Name	Middle Initial	Clinic Name		
Social Security #	Date of Birth	Sex <input type="checkbox"/> F <input type="checkbox"/> M	Ethnicity <input type="checkbox"/> African American <input type="checkbox"/> Asian <input type="checkbox"/> Ashkenazi Jewish <input type="checkbox"/> Caucasian <input type="checkbox"/> Hispanic <input type="checkbox"/> Other		
Address		City	State	Zip	Phone
<b>INSURANCE:</b> Please provide a legible copy of the front and back of the patient's insurance card. IF NO INSURANCE: <input type="checkbox"/> Self Pay <input type="checkbox"/> WC/Auto (Date of Injury) <input type="checkbox"/> Other					
Name of Insured	Relationship to Patient	Insurance Company/Provider	Member/ID Number	Group Number	
Collector Name (Print)	Date Collected	Time Collected	Fasting <input type="checkbox"/> Yes <input type="checkbox"/> No		
Specimen Type <input type="checkbox"/> OCD-100 (Buccal)	Specimen Storage <input type="checkbox"/> Room Temperature <input type="checkbox"/> Refrigerated		Specimen Shipping <input type="checkbox"/> Room Temperature <input type="checkbox"/> Cooling/Ice Pack		

### MOLECULAR DIAGNOSTICS TESTING OPTIONS

**Malignant Hyperthermia Susceptibility Genomics Test** Please select the Panel to be tested. Please attach patient Medication List.

Malignant Hyperthermia Susceptibility NGS Panel

CACNA1S, BCHE, BCHE -A, BCHE-K, BCHE-F1, BCHE-F2, BCHE-S1, RYR1

### Personal/Family History Questionnaire Please complete Questionnaire

#### PATIENT'S PERSONAL HISTORY (Hx)

Clinical Details	Personal Hx	Age at Dx
Mosaicism	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Consanguinity	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Bone Marrow Transplant	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Organ Transplant	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Known Chromosomal Gain/Loss	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Known Gene Gain/Loss	<input type="checkbox"/> Yes <input type="checkbox"/> No	

#### FAMILY HISTORY

Relationship	Maternal	Paternal	Cancer Site(s)	Age at Dx
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		
	<input type="checkbox"/>	<input type="checkbox"/>		

**Clinical Presentation** Please indicate any clinical presentations and/or findings that may be relevant to genetic testing:

- Behavior  Conditions  Pedigree/Family History  
 Phenotypes  Physical  Symptoms

**Clinical Testing** Please indicate any clinical testing results and/or findings that may be relevant to genetic testing:

- Karyotype  Vision  Growth Measurements  Imaging  
 Previous Genetic Testing  Hearing  Biochemical Testing  Pathology Results

**ICD-10 DIAGNOSIS CODES:** Additional documentation supporting Medical Necessity may be attached.

- |  |   |
|--|---|
| <input type="checkbox"/> T88.3XXA Malignant hyperthermia due to anesthesia, initial encounter                      | <input type="checkbox"/> E87.2 Acidosis   |
| <input type="checkbox"/> T88.3XXD Malignant hyperthermia due to anesthesia, subs encounter                         | <input type="checkbox"/> E87.5 Hyperkalemia   |
| <input type="checkbox"/> T88.3XXS Malignant hyperthermia due to anesthesia, sequela                                | <input type="checkbox"/> I10 Hypertension   |
| <input type="checkbox"/> P81.9 Disturbance of temperature regulation of new-born, unspecified (New-born 0-28 days) | <input type="checkbox"/> I47.2 Ventricular tachycardia  |
| <input type="checkbox"/> T41.0X5A Adverse effect of inhaled anesthetics, initial encounter                         | <input type="checkbox"/> I49.01 Ventricular fibrillation  |
| <input type="checkbox"/> T41.0X5D Adverse effect of inhaled anesthetics, subsequent encounter                      | <input type="checkbox"/> M62.82 Rhabdomyolysis  |
| <input type="checkbox"/> T41.0X5S Adverse effect of inhaled anesthetics, sequela                                   | <input type="checkbox"/> M62.838 Other muscle spasm   |
| <input type="checkbox"/> T41.0X6A Underdosing of inhaled anesthetics, initial encounter                            | <input type="checkbox"/> M62.89 Other specified disorders of muscle (Muscle rigidity)             |
| <input type="checkbox"/> T41.0X6D Underdosing of inhaled anesthetics, subsequent encounter                         | <input type="checkbox"/> R00.0 Tachycardia, unspecified   |
| <input type="checkbox"/> T41.0X6S Underdosing of inhaled anesthetics, sequela                                      | <input type="checkbox"/> R00.1 Bradycardia, unspecified   |
| <input type="checkbox"/> T41.1X5A Adverse effect of intravenous anesthetics, initial encounter                     | <input type="checkbox"/> R03.0 Elevated BP reading, w/o diagnosis of hypertension                 |
| <input type="checkbox"/> T41.1X5D Adverse effect of intravenous anesthetics, subsequent encounter                  | <input type="checkbox"/> R06.82 Tachypnea, not elsewhere classified                               |
| <input type="checkbox"/> T41.1X5S Adverse effect of intravenous anesthetics, sequela                               | <input type="checkbox"/> R50.9 Fever  |
| <input type="checkbox"/> T41.1X6A Underdosing of intravenous anesthetics, initial encounter                        | <input type="checkbox"/> R61 Generalized hyperhidrosis  |
| <input type="checkbox"/> T41.1X6D Underdosing of intravenous anesthetics, subsequent encounter                     | <input type="checkbox"/> Z92.84 Personal history of unintended awareness under general anesthesia |
| <input type="checkbox"/> T41.1X6S Underdosing of intravenous anesthetics, sequela                                  | <input type="checkbox"/> Z92.89 Personal history of other medical treatment                       |
| <input type="checkbox"/> T41.295A Adverse effect of other general anesthetics, initial encounter                   | <input type="checkbox"/> Z84.89 Family history of other specified conditions                      |
| <input type="checkbox"/> T41.3X5A Adverse effect of local anesthetics, initial encounter                           | <input type="checkbox"/> Z78.9 Other specified health status                                      |
| <input type="checkbox"/> T41.45XA Adverse effect of unspecified anesthetic, initial encounter                      | <input type="checkbox"/> Other  |

**Medical Necessity**  
Required for insurance

I, the provider, attest that I am the ordering physician or am authorized under applicable laws and regulations to order genetic testing for the patient. I attest that this test is medically necessary for the diagnosis or detection of a disease or disorder, and that the results will be used in medical management and care decisions for the patient. I further attest that any information entered on this Test Requisition Form, or otherwise provided by me on behalf of the patient, is true and correct to the best of my knowledge, and that the patient has consented to receive communications about his/her genetic test from RDL.

**Patient Informed Consent**  
Patient must consent

I, the patient, voluntarily consent to the collection and testing of my specimen. I certify that the specimen is fresh and has not been adulterated in any manner. I authorize the laboratory to release the results of this testing to the ordering provider. I further authorize my insurance benefits to be paid directly to RDL for services rendered. I acknowledge that the lab may be treated as an out-of-network provider. In the event I receive payment for laboratory services from my insurer, I will remit said payment to the lab within 14 days of receipt. I will either endorse the original check, or produce a personal check for the entire payment amount, and forward it to the lab. When selecting Self Pay above, I acknowledge financial responsibility for all lab charges associated with the processing of this test requisition. All rights to the samples will belong to the laboratory conducting the testing. There will be no compensation in the event of an invention resulting from research and development using this sample. I agree to allow my provided specimen to be used for the purpose of (diagnosis/research) (development/quality control). I understand that if I agree, any information identifying me will be kept confidential so that it will not be possible to determine from whom the sample was drawn. Your signature on this form indicates that you understand to your satisfaction the information about RDL and agree to have the test done. In no way does this waive your legal rights or release anyone from their legal and professional responsibilities. If you have further questions concerning matters related to this consent, you may wish to seek professional genetic counseling prior to signing this form. Consultation with a medical geneticist, genetic counselor, or your referring healthcare provider also may be warranted after the test has been completed.

**Opt In for Research**

I give permission for my specimen and clinical information to be used in de-identified studies at Tesis Biosciences and for publication, if Tesis deems it appropriate. I understand that my name and/or other identifying information will NOT be used in or linked to the results of any studies and publications. More information is available at www.tesisbiosciences.com.

Provider Name (Print)	Provider NPI #	Clinic Address	Clinic Phone/Fax
Provider Signature	Date	Patient Signature (or Legal Guardian)	Date
X		X	