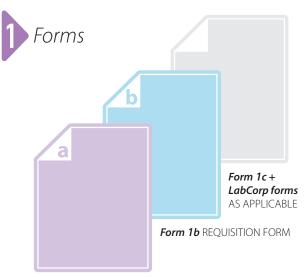
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NeuroSURESM: Epilepsy Gene Panel Genetic Test Submission Guide







Form 1a INFORMED CONSENT

3 Ship



Impact Genetics 115 Midair Court Brampton, ON L6T 5M3 1-877-624-9769

Results





Form 1a: Informed Consent to Perform Genetic Testing

The purpose of my DNA test/or my child's DNA test is to look for mutation(s) or genetic alterations known to be associated with the following genetic condition or disease:

By signing below, I acknowledge that:

- 1. My participation or, as applicable, my child's participation in this DNA testing is voluntary. The decision to consent to, or to refuse the above testing is entirely mine.
- 2. This testing is done on small biological samples.
- 3. It is possible that the quantity or quality of sample submitted may be inadequate for testing or that a mutation cannot be identified.
- 4. When DNA testing shows a mutation or alteration, then the person is a carrier or is affected with that condition or disease, or, in the case of cancer genetic testing, the person is a carrier of a mutation or alteration that may be associated with an increased risk for certain cancer(s) compared to the general population. Consulting a doctor or genetic counsellor is recommended to learn the full meaning of the results and to learn if additional testing might be necessary.
- 5. When the DNA testing does not show a known mutation or alteration, the chance that the person is a carrier or is affected is reduced, or, in the case of cancer genetic testing, the person's risk for certain cancer(s) compared to the general population will depend on additional personal factors. There is still a chance to be a carrier or to be affected because the current testing cannot find all the possible changes within a gene.
- 6. Impact Genetics will disclose the test results ONLY to the specialist designated on the Requisition Form (*Form 1b*), or to his/her agent, unless otherwise authorized by the patient or required by law, except as described in point 11 below, no information will be printed or released that discloses the patient's identity, or other confidential information.
- 7. Impact Genetics is not a DNA banking facility and patient DNA samples may not be available for future testing.
- 8. An error in diagnosis may occur if the true biological relationships of the family members are not as stated in the pedigree submitted with the Requisition Form (*Form 1b*). It is possible that the test may disclose non-paternity (someone who is not the biological father), or some other previously unknown information about family relationships, such as adoption, and I consent that this finding be reported to the referring specialist designated below.
- 9. There is a chance that the test may reveal unexpected abnormalities that may be of medical value in the patient's care. Impact Genetics will inform the referring specialist designated below.
- 10. Until the results of this test are reported, the patient and members of the patient's family should still undergo examinations as prescribed by the referring specialist.
- 11. If necessary to obtain reimbursement of test fees, Impact Genetics, its agents and legal representatives may disclose information that identifies the patient or other confidential information (including test results).
- 12. I have read or have had read to me, the above information and I understand it. I have also read or had explained to me the specific disease or condition tested for, and the specific test(s) I am having, including the test descriptions, principles and limitations. I have had the opportunity to discuss the purposes and possible risks of this testing with my doctor or someone my doctor has designated.

Consent for Future Research: After all analysis required to reach a genetic diagnosis is complete, Impact Genetics has my consent to use DNA or RNA in an anonymous fashion for research. No tests(s) will be performed and reported on my samp than the one(s) authorized by my doctor. YES NO	
Signature of Patient:	Date:
Signature of Witness:	Date:

LAB USE ONLY DO NOT FILL OUT			
Date received: Y	М	D	
Specimen type:			
Condition:			
MRN:		Tech:	

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Form 1b: NeuroSURESM: Epilepsy Gene Panel Test Requisition

Test Panel	Referring Specialist
See menu on page 2.l	Name:
☐ Epilepsy: Comprehensive (69 genes)	Specialty:
☐ Epilepsy: Single Gene(gene name)	Contact:
Patient	Telephone:Fax:
	Email:
Last name:	Cianatura
Date of birth: Y M D	Signature:
Gender: □ Male □ Female	Institution:
Pregnant: ☐ Yes ☐ No	Address:
: riegilant. 🗆 les 🗆 no	
Patient History	City:Prov/State:
☐ Confirmed clinical diagnosis	Postal code:Country:
Diagnosis Date: Y M D	Additional copies to:
☐ Suspected clinical diagnosis ☐ Unaffected	Email:Fax:
Clinical Diagnostics & Family History	Billing
☐ Isolated case ☐ Positive family history	Impact Genetics tests ordered through LabCorp test menu.
Family previously tested: Yes No	USA only.
Mutation identified:	☐ Client Bill
If mutation identified at lab other then Impact Genetics	Provide details:
please provide report.	
Proband name (first person in a family to be studied):	
Gene:	
: Mutation:	☐ Patient Pay
: Polationship to Proband (Indox Case)	Complete LabCorp Financial Responsibility Form .
Relationship to Proband (Index Case)	☐ Third party insurance (USA only)
☐ Proband ☐ Parent of proband ☐ Child of proband	Complete Form 1c: U.S. Insurance Information and to expedite testing, complete LabCorp Financial Responsibility Form.
Other:	Pre-authorization services provided.
	· · · · · · · · · · · · · · · · · · ·
Specimen Information	115 Midair Court, Brampton, ON L6T 5M3
Sample:	t: 647-478-4902 or 1-877-624-9769 f: 905-697-9786 e: info@impactgenetics.com <u>Please ensure to use secure email</u>
☐ EDTA (lavender top) Blood sample for DNA	C. Thousinpucty choices. Com <u>Freuse ensure to use secure enfull</u>
□ DNA from blood (min. 500 ng)	
☐ Buccal swab (only for preapproved familial mutation confirmation, contact lab directly before submitting)	
Date of collection: Y M D	
Time of collection: HH:MM (24hr)	



Form 1b: NeuroSURESM: Epilepsy Gene Panel Test Requisition

Patient name:	Data of birth. V	N.4	D
Patient name.	Date of birth: Y	IVI	D

Epilepsy Test Panels

Epilepsy: Comprehensive (69 genes)

Consider when the prognosis based on clinical and EEG findings is poor or the likelihood of lethal outcome is high, when clinical or EEG findings are not specific for a specific epilepsy syndrome.

ALDH7A1, AMT, ARX, ASAH1, ATP1A2, ATP1A3, CDKL5, CERS1, CHD2, CHRNA7, CNTNAP2, CSTB, DNM1, DOCK7, EPM2A, FOLR1, FOXG1, GAMT, GATM, GLDC, GOSR2, GRIN2A, GRIN2B, HCN1, KCNC1, KCNJ10, KCNJ11, KCNQ2, KCNQ3, KCNT1, KCTD7, LMNB2, MBD5, MECP2, MEF2C MOCS1, NECAP1, NEU1, NHLRC1, NRXN1, PCDH19, PHGDH, PLCB1, PNKP, PNPO, POLG, PRICKLE2, PRRT2, PSAT1, PSPH, SCARB2, SCN1A, SCN1B, SCN2A, SCN8A, SCN9A, SLC2A1, SLC6A8, SLC9A6, SPTAN1, STXBP1, SUOX, SYNGAP1, TBC1D24, TCF4, TSC1, TSC2, UBE3A, ZEB2.



Form 1c: U.S. Insurance Information

Impact Genetics is committed to providing the highest quality genetic testing to all patients. In many situations, genetic testing improves outcomes and quality of life and decreases total costs to the patient and healthcare system.

Processing medical insurance claims is usually challenging and time consuming. Many insurance companies require pre-authorization prior to testing. Impact Genetics supports insurance billing, completing coverage checks and pre-authorization.

It is important for patients to understand that insurance rarely covers 100% of the cost of genetic testing and that they will be financially responsible for some or all of the cost of testing. The patient is responsible for any portion of the test fee not covered by insurance for any reason, including but not limited to, co-payments, unmet deductibles, co-insurance and non-covered services. Prior determinations do not guarantee payment and the amount paid by insurance when the claim is submitted may be different from the coverage indicated during the pre-verification or pre-authorization process.

Pre-authorizations can take time to obtain depending on each individual insurance plan's policy and documentation requirements. Turnaround time for test results begins after the pre-authorization has been processed and approved.

For tests ordered through LabCorp, LabCorp administers billing. The LabCorp prior-authorization team will file a pre-verification/prior-authorization on behalf of the patient with any commercial insurance company. State managed Medicare plans cannot be billed.

Insurance process

- 1. Send Form 1c: U.S. Insurance Information as soon as possible.
- 2. Insurance coverage will be investigated and patient/specialist will be contacted to provide coverage estimate if the patient's out of pocket costs are over \$300.00.
- 3. Insurance claim will be submitted upon completion of testing.
- 4. After insurance payment is received patient will be billed for non-covered services.

Note: Timely and complete submissions will enable faster insurance checks.



Send this form to LabCorp BY FAX TO 1-888-598-7568

Form 1c: U.S. Insurance Information - NeuroSURESM: Epilepsy Gene Panel

Disease/Genetic Test	Patient Information		
☐ Test # 481518	Last name:		
NeuroSURE SM : Epilepsy Gene Panel - Comprehensive	First name:		
ICD-10 Code "provide code here"	Date of birth: Y M D		
CPT Codes: 81403, 81404, 81405, 81406, 81407, 81479	Address:		
Insurance Information			
☐ Primary insurance ☐ Secondary insurance	City:		
If Patient has secondary insurance, include the information	State:Zip code:		
on an additional copy of this form with the secondary	Country:		
insurance box checked.	Telephone:		
Name of insured (if not Patient):	. receptione.		
	Contact Information		
Insurance company:	Details of insurance coverage will be communicated.		
Claims address:	Please provide preferred telephone number(s):		
	☐ Patient ☐ Referring specialist		
City:	Telephone:		
State:Zip code:	Alternate telephone:		
Country:	Email:		
Group #:	☐ In the event Patient cannot be reached a voice message		
Subscriber/member #:	related to epilepsy genetic testing may be left at the above phone number(s)		
Physician Information	Please Attach <u>All</u> of the Following		
Physician's name:	 Copies of both the front and back of insurance membership card(s) 		
NPI:	•		
Practice name:	 Letter of Medical Necessity, signed by Referring Specialist (contact Impact Genetics for template 		
Practice Address:	if needed)		
	• Clinic notes demonstrating the Patient's need for		
Telephone:	testing and confirmation of diagnosis		
Fax:	• Insurance approval details <i>if</i> prior pre-approval completed		
LabCorp account #:	Performing Lab - Impact Genetics 115 Midair Court Brampton, ON 16T 5M3, CANADA		

Testing process will be initiated when **LabCorp Financial Responsibility Form** is received or confirmation is received from insurance provider. t: 1-877-998-7837 f: 1-888-598-7568

e: preverification@labcorp.com Please ensure to use secure email

LabCorp Statement of Financial Responsibility

S	ection A: Member/Patient Inform	nation		
Me	mber/patient name:		/ Date of b	irth:
				/ ZIP:
		/ Client account #:		
Sul	oscriber #:			
S	ection B: Requested procedure o	r service information		
	sed on information given to us by your in ysician/healthcare provider (marked bel	nsurance plan, your plan is not expecte ow).	d to pay 100% for the laboratory	test(s) ordered by your
	Test/CPT Description	Reason for Patient Ou	t of Pocket	Estimated Cost*
Tot	tal Estimated Patient Responsibility*:	*This is only an estimate. Actual amour	 nt owed may be adjusted based or	n final coverage amount.
Ne	ext Steps:	·	, ,	
	, –	to be financially responsible for the estir		listed above.
	' '	to receive the items listed in Section B a thin 5 calendar days via email, fax, or ma		Ve will not proceed until we
	receive this signed consent form with an		in to the addresses listed below. V	ve wiii not proceed until we
S	ection C: Options—Check only o	ne box. We cannot choose a box t	for you.	
	•	s) marked above to be performed. I und nally and fully responsible for payment.	erstand that my insurance plan is	not expected to pay for these
	☐ I would like to set up a payment pl	an for \$a month.**		
	Option 2—I want the services listed above, but do not bill my insurance. I understand I cannot appeal the coverage of these services with my insurance plan if they are not billed. Discounts may apply to the services listed above. Payment in full is required in order to proceed with the testing services.			
	Option 3—I do not want the laboratory test(s) marked above to be performed. I understand with this choice I am not responsible for paymen. No test will be performed and my plan will not be billed.			
Pat	ient can contact the Billing department	at 888-210-9264 to discuss payment op	tions. For non-billing questions, c	all 855-488-8750.
**Y	our first invoice will include the full balance d	lue. If your payment plan is approved, you wi	Il receive another invoice that reflects	your requested payment amount.
Sig	nature:		/ Date:_	
				
r16	ase print name:			
	LabCorp	Email this form to: Mail this form to: LabCorp Prior Au	/ Fax this f	form to:

www.LabCorp.com



Step 2: NeuroSURESM: Epilepsy Gene Panel Test Sample Requirements

Sample Requirements

Proband:

• Blood or DNA from blood.

Genetically related family member for known mutation:

- Blood or DNA from blood;
- Or buccal swab (select cases only, contact lab directly to confirm if appropriate sample).

If genetic testing has been performed at a lab other than Impact Genetics, please provide an aliquot of proband DNA and/or a copy of proband's report.

Sample Preparation Instructions

Blood samples for DNA:

• 10 mls in lavender-topped EDTA tubes (for infants 2-5 mls in pediatric or small tubes) at room temperature, to be received within 3-5 days after draw. Best results are achieved if received within 3 days.

DNA from blood:

• DNA quantity—minimum 500 nanograms. Ship with cool packs.

Sample Identification

• Label each sample with at least two patient identifiers (e.g. name and date of birth).



Step 3: Shipping Requirements

Multiple separated samples may be shipped in one box. Place multiple biohazard bags containing labeled samples into one box. Multiple boxes can be shipped in one courier envelope.

Shipping Instructions

- Ship samples to Impact Genetics at address shown on this page using a courier envelope.
- Include Informed Consent for Genetic Testing and Requisition Forms (1a and 1b) with the samples. Patients in the U.S. must also include U.S. Insurance Information (Form 1c) if required and not provided previously.
- Complete appropriate Air Waybill. If you cannot use FedEx or Purolator, please contact us.
- Place Air Waybill in the document pouch.
- For samples from outside of Canada, complete and sign appropriate customs forms (provided and available on our website; phone us if help is required). Place the customs forms in the document pouch.
- Within Canada, use **Purolator Express** (next-day) or **FedEx Priority** service (next-day). Outside of Canada use **FedEx Priority** service (next-day) and **use a FedEx "Clinical Pak"**. If you cannot use Purolator or FedEx, please contact us.
- Provide us with the parcel tracking number soon after courier pick-up: 647-478-4902, info@impactgenetics.com.
- For emailed PDF FedEx waybills and customs forms, please contact us directly.

Send to Impact Genetics

mail: Impact Genetics 115 Midair Court Brampton, ON L6T 5M3 tel: 1-877-624-9769

fax: 905-697-9786