

<b>Providence Health Plan</b> Statement of Medical Necessity Form	<b>Hereditary Arrhythmia Genetic Testing for LongQT Syndrome (LQTS) and Catechoamlinergic Polymorphic Ventricular Tachycardia (CPVT)</b>
Patient Information	Physician Information
Name:	Physician Name:
Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Tax ID Number:
Providence Health Plan ID Number: <input type="checkbox"/> HMO <input type="checkbox"/> PPO/ASO	Specialty: <input type="checkbox"/> PCP <input type="checkbox"/> Specialist <input type="checkbox"/> Genetic Counselor (Please Identify):
Birth Date (mm/dd/yy):	Rendering Provider (Laboratory): Invitae Corporation
Date of Service (mm/dd/yy):	Rendering Provider ID Number: NPI: 1316206220
Has the patient undergone prior BRCA testing? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Coding Information	
<i>NK2, CACNA1C, CALM1, CALM2, CALM3, CAV3, KCNE1, KCNE2, KCNH2, KCNJ2, KCNQ1, SCN5A, TRDN</i> Gene Analysis CPT Codes (check all that apply): <input checked="" type="checkbox"/> <b>81413</b> (Long QT syndrome gene analyses; full sequence analysis)	
ICD-10 Codes (list all applicable codes):	
Medical Necessity Criteria	
<i>NK2, CACNA1C, CALM1, CALM2, CALM3, CASQ2, CAV3, KCNE1, KCNE2, KCNH2, KCNJ2, KCNQ1, RYR2, SCN5A, TRDN</i> <b>Mutation Testing:</b> Genetic testing for mutations in these genes may be considered <b>medically necessary</b> when <b>any</b> of the following criteria are met (please check all applicable boxes):	
<input type="checkbox"/> Individual (male or female) from a family with a known deleterious LQTS or CPVT gene mutation; identify gene and mutation: _____	
<input type="checkbox"/> Personal history of idiopathic long QT on ECG (resting or stress) or polymorphic ventricular tachycardia on stress ECG;	
<input type="checkbox"/> Personal history of ECG abnormalities associated with LQTS including Torsades de Pointes, T-wave alternans, or notched T-waves in >3 leads	
<input type="checkbox"/> Personal history of unexplained syncope	
<input type="checkbox"/> Personal history of aborted sudden death	
<input type="checkbox"/> Family history of LQTS or CPVT in a first- or second-degree relative, as documented by ECG (resting or stress) <i>Family Member:</i> _____ <i>Age:</i> _____ (years)	
<input type="checkbox"/> Family history of LQTS or CPVT in two or more first-, second-, or third-degree relatives, one of whom is a first-degree relative, as documented by ECG (resting or stress) <i>Family Member:</i> _____ <i>Age:</i> _____ (years) <i>Family Member:</i> _____ <i>Age:</i> _____ (years)	
<input type="checkbox"/> Parent of a child with <u>Sudden Unexplained Death in Infants (SUDI)</u> syndrome	
I confirm that this test is medically necessary in accordance with ( <i>Providence Health Plan</i> ) medical policy and that the information provided is accurate and factual based on the patient's medical records and history. I confirm that this test is medically necessary for the risk and assessment, diagnosis or detection of a disease, illness, impairment, symptom, syndrome or disorder. The results will determine my patient's medical management and treatment decisions. I confirm that I have been trained to provide genetic counseling, and that I have conducted a full personal and family history which includes a first, second and third degree pedigree. I have provided genetic testing information and pre-test counseling to the patient and they have consented to genetic testing. I have scheduled post-test counseling to review the test results and determine future medical management and treatment plans.	
Ordering Provider Signature: _____ Date (mm/dd/yy): _____	
Please print name: _____ NPI number: _____	
Date of genetic counseling (mm/dd/yy): _____	