

Genomic Medicine Service

National Genomic Test Directory Clinical Indication R89 Non-urgent Exome Sequencing Test Request

Please complete this form and email to the laboratory rduh.exterexome@nhs.net BEFORE sending any samples. Ensure that email addresses are provided for the responsible clinician and/or clinical geneticist.
CONSENT: Receipt of samples for testing assumes that **informed consent** has been obtained for all family members being tested and the possibility of incidental findings has been discussed.

Please send at least 1µg of DNA per individual to: Exeter Genomics Laboratory, RILD Level 3, Royal Devon University Healthcare NHS Foundation Trust, Barrack Road, Exeter, EX2 5DW.

Patient first name:		Life status: <input type="checkbox"/> Alive <input type="checkbox"/> Deceased		Ethnicity:		
Patient last name:		Family test: <input type="checkbox"/> Trio <input type="checkbox"/> Duo <input type="checkbox"/> Singleton		Consanguinity: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown		
Date of birth: dd/mm/yyyy	Hospital number:		Trio – affected child and both parents (gene-agnostic) Duo – affected child and one parent (gene panel) Singleton – affected child only (gene panel)			
Gender (if phenotypic sex is different please state): <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other:						
NHS number (or postcode if not known)						
<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>						
Clinical information and any relevant family history						
Reason for testing – clinical summary (this text, or an abbreviated version of it, will be included in the report): Please also include relevant information for relatives and relationship to other tested individuals, including disease status and age of onset						
Differential diagnosis: Please list						
HPO terms (https://hpo.jax.org/app/) to describe phenotypes in this individual (OPTIONAL): Please list						
Family DNA samples provided (parents for trio tests)						
Surname	Forename	Date of birth dd/mm/yyyy	NHS number	Gender <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other:	Deceased <input type="checkbox"/> No <input type="checkbox"/> Yes	Status <input type="checkbox"/> Affected <input type="checkbox"/> Unaffected
		dd/mm/yyyy		<input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other:	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Affected <input type="checkbox"/> Unaffected
		dd/mm/yyyy		<input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other:	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Affected <input type="checkbox"/> Unaffected
Clinician details						
Responsible clinician / consultant paediatrician: Name, Department, Hospital			Email address for report: (nhs.net)			
			Telephone number:			
Clinical geneticist: Name, Department, Hospital			Email address for report: (nhs.net)			
			Telephone number:			