

Clinical exome pre-approval form

Medicare funded testing

1. CLINICIAN INFORMATION

NAME OF PAEDIATRICIAN:

PRACTICE LOCATION:

PHONE:

NOTE: Training is available to support your ability to safely order genomic testing. These resources can be found at vcgs.org.au/tests/clinical-exomes

2. PATIENT INFORMATION

FIRST NAME		WEIGHT (<i>centile</i>)		DATE OF BIRTH	
SURNAME		HEIGHT (<i>centile</i>)		HEAD CIRCUMFERENCE (<i>centile</i>)	
IS THERE ANY RELEVANT FAMILY HISTORY / SIMILARLY AFFECTED RELATIVES? <input type="checkbox"/> No <input type="checkbox"/> Yes (<i>provide details</i>)					

Parent information for trio testing

	FULL NAME	DATE OF BIRTH	ETHNICITY	AFFECTED
BIOLOGICAL MOTHER				<input type="checkbox"/> No <input type="checkbox"/> Yes
BIOLOGICAL FATHER				<input type="checkbox"/> No <input type="checkbox"/> Yes
ARE THE PARENTS CONSANGUINEOUS? <input type="checkbox"/> No <input type="checkbox"/> Yes				

3. MEDICARE ELIGIBILITY

Strict criteria apply for access to Medicare rebate for exome sequencing

Patient History (REQUIRED)

Has the patient had a non-informative chromosome microarray (CMA, molecular karyotype)?	<input type="checkbox"/> No* <input type="checkbox"/> Yes <input type="checkbox"/> Unsure (<i>include a copy of result</i>) <i>*CMA is required before an exome can be ordered</i>
Is the patient admitted (inpatient or ED)?	<input type="checkbox"/> No <input type="checkbox"/> Yes* <i>*patient ineligible while admitted</i>
Has the patient had genomic testing? <i>(defined as testing for a variant/s in multiple genes in one test and includes whole genome and whole exome sequencing or testing of a panel of genes)</i>	<input type="checkbox"/> No → continue to next question. <input type="checkbox"/> Yes → it was Medicare funded (<i>patient is ineligible</i>). Contact a geneticist to discuss. <input type="checkbox"/> Yes → Include a copy of the result.

Does the patient have:	If yes:	Features
Dysmorphic facial features <input type="checkbox"/> No <input type="checkbox"/> Yes	email photos with form AND list features →	

Does the patient have:	If yes:	Features
Congenital anomalies <input type="checkbox"/> No <input type="checkbox"/> Yes	list features →	
		Description of severity, including developmental milestones
Intellectual disability of at least moderate severity <input type="checkbox"/> No <input type="checkbox"/> Yes	provide IQ score OR detailed description →	

Other relevant features (OPTIONAL)

Differential diagnoses	
Other phenotypic features	
Results from other relevant investigations (e.g. brain MRI; metabolic investigations)	

Gene panels (REQUIRED)

What are the gene panels you wish to apply in this test?
 (Please use the PanelApp Australia website - <https://panelapp.gha.umccr.org/>)

3. OUTCOME OF DISCUSSION WITH CLINICAL GENETICIST

Medicare eligible No Yes

Test type singleton WES trio WES (In the opinion of the clinical geneticist, singleton testing is inappropriate)

Name of clinical geneticist	
Approval number	

If approved:

- Formal written consent from the patient is required (and parents if trio testing)
- A test request form must also be completed
- Forms are available on [our website](#)
- This form + the test request + patient consent must be sent to VCGSgenomics@vcgs.org.au
 - please save forms in the following format = SURNAME_firstname_dob [DDMMYY]

TESTING WILL NOT PROCEED WITHOUT ALL FORMS BEING SENT TO THE LABORATORY