V1.00 - 11/2024	Winc Code: 1NY44194

SW1259	
V1.0	
SW1	

. 为备款性	(Affix patient ident	tification label here)
Queensland Government	URN:	
	Family Name:	
CLINICAL OFNETIO (OFNOMIC	Given Names:	
CLINICAL GENETIC / GENOMIC TESTING CONSENT	Address:	
TESTING CONSENT	Date of Birth:	Sex: M MF MI
Test details	Date of Bitti.	Sex. W F I
	Genomic test:	
Genetic test:		
☐ Single gene ☐ Single variant ☐ Multiple variants		ome Genome
Other (specify):	Other (specify):	
Test purpose:	Test purpose:	rrier screening
☐ Diagnostic test ☐ Predictive test ☐ Carrier test ☐ Confirmation test ☐ Segregation test ☐ Propostal test	☐ Diagnostic test ☐ Ca☐ Prenatal test	rrier screening
☐ Confirmation test ☐ Segregation test ☐ Prenatal test		
Other (specify):	Other (specify):	
Clinical indications or condition tested for:		
 The test may find a genetic change of uncertain significated. More tests or analysis may be needed to understand the standard the stand	e results. This may include testing rusing new knowledge or testing rusurance. Iaboratories to assist with genonith genomic and medical database consent, or as required or permit sample has been collected, and cain in my medical records.	methods. nic testing. es that are used for patient ted by law. choose not to be told the
Name:		
Information sharing	h health professionals to hale wit	h the
 I consent to share the results and related information wi genetic testing of blood relatives. I understand that ident the relative wherever possible and not without my prior version. 	ifying information will not be discl	
 I consent to share the sample, genetic/genomic test data ethically approved research into the same or related cor information will be removed and will usually be replaced 	a, and related health information t	for Yes No

my test report is uploaded to My Health Record (MyHR).

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CLINICAL GENETIC / GENOMIC TESTING CONSENT

/ACC (* (*) (*) (*)
(Affix patient identification label here)
URN:
Family Name:
Given Names:
Address:
Date of Birth: Sex: M M F MI

Consent

Patient / proxy:

- I consent to genetic/genomic testing.
- I understand the reason for testing and the potential benefits, consequences and limitations.
- I have been given a copy of the Patient Fact Sheet Genetic and Genomic Testing (V1.0), which provides information about testing.
- I have been able to discuss the information with a health professional, ask questions and have any concerns addressed.

I am satisfied with the explanations and answers to my questions.	
Name of patient:	
Patient signature:	
Email address:	
Genetic file number (if applicable):	
Or, where consent is given on behalf of another:	
Name of proxy:	
Proxy signature:	
Relationship to patient:	
Email:	Phone number:
Health professional:	
l,	
(Name of health professional)	(Signature)
(Dasimetian)	
(Designation) have provided information on the reason for and nature of the test, p	(Date)
The patient / proxy has been able to ask questions and consider the	•
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Other declarations	
Other declarations	
Interpreter / Liaison Officer: Not applicable	
Interpreter / Liaison Officer: Not applicable	
Interpreter / Liaison Officer: Not applicable	(Signature) (Date) pplied by the health professional to the patient.
Interpreter / Liaison Officer: Not applicable I, (Name of interpreter / Liaison officer) have interpreted the content of this form and all the information su	oplied by the health professional to the patient.
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Interpreter / Liaison Officer: Not applicable I, (Name of interpreter / Liaison officer) have interpreted the content of this form and all the information su Consent for parents undergoing duo/trio genomic analy I/we consent to genomic testing for the purpose of assisting in the patient named above). I/we understand the reason for testing and Specifically, I/we understand that the details of genomic testing our related information. I/we have been able to discuss the information.	pplied by the health professional to the patient. *rsis: Not applicable interpretation of the genomic results of my/our child (the the potential benefits, consequences and limitations. tlined above apply to my/our sample(s), results and with a health professional, ask questions and have any answers to my/our questions. In information for ethically approved research into my/our eremoved and may be replaced with a unique code so
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