

DNA extraction and storage

(EDTA or sample in transport media).

Wellington Regional Genetics Laboratory (WRGL)

Wellington Hospital Private Bag 7902 Wellington 6242 Tel: (04) 918 5352 Fax: (04) 385 5822

Oncol	ogy Genetics Referral Fo	orm	Email: Oncol	ogyWRGL@ccdhb.org.nz	
NHI:	DOB:		Requester	:	Sample Taken
Family Name:	Sex: F/M		Print name	:	by:
Given Name:	DHB of Domicile		Copy repo	rt to:	Date:
Address:					Time:
Clinical details	s :		Sample det	tails:	
Diagnosis: Current state: Diagnosis: Pre-Tre			□ Blo	od Sample nph node id tumour tissue	□ Right
Cytoge	enetic Tests Requested	Hold	Proceed Immediately	Comme	ents
Chromosoi	me analysis (LH or sample in transport media)		•		
	FISH analysis from peripheral blood, bone marrow, lymph node or solid tumour samples (LH or sample in transport media). For FFPE slides please refer to separate FFPE referral from)			List FISH tests required website for a full list of p http://www.wellingtongenetics List.html	robes available)
FISH	FISH for plasma cell disorders CD138+ cell sorting is required. Samples must be received within 24 hours.			If plasma cell sorting is indicate the plasma cell (or contact the laborator	percentage %
	CLL workflow (LH and EDTA or sample in transport media)			This workflow includes 7 NGS.	TP53 FISH and

Molecular Genetics Tests Requested	Hold	Proceed Immediately	Comments
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MYD88 testing			
(EDTA or sample in transport media)			
Chimerism testing (please circle) a. Same sex: EDTA required b. Sex-mismatch: LH required			EDTA is required for same sex as STR analysis is required (this includes donor, pre-transplant or post-transplant). FISH for sex-mismatch
NGS testing (FFPE or sample in transport media)			
For optimum DNA quality and test success rate fresh tissue is ideal (a minimum of 5mm³ in transport media). If testing is required from an FFPE sample, please provide a minimum of 7 de-waxed, uncoated slides cut at 4-6 microns along with 1 H&E slide with the area of interest circled and the percentage of neoplastic cell nuclei indicated.			
ldeal sample criteria and thresholds include: - Use of low-concentration (4% v/v formaldehyde) neutral-buffered formalin (12-24 h fixation)* - Specimen age below 8 years** - Minimum amount of DNA required is 50ng*** - Tumour burden ≥30%		X	Percentage of neoplastic cell nuclei:% Please note: we require a minimum of 30% neoplastic cells to proceed with NGS
*Shorter or longer times may cause enzymatic degradation of the tissue, difficulties in DNA extraction and compromise the result. **Long-term storage of FFPE tissue blocks can influence the quality of nucleic acids		^	Specify genes to interrogate (please circle BRCA1/BRCA2
***It is not possible to give a minimum size required and the concentration obtained is not known until after DNA extraction. However, the larger the sample, the better chance we have of obtaining a successful result.			
Samples with <50ng will not be processed.			
Please note: NGS tests do not distinguish between somatic and germline variants. Germline variants with significant implications for both the patient and their family may be detected.			
NGS Sendaway (EDTA required) - please specify tests required and complete appropriate consent and molecular referral form.			Specify test required:
Consent has been obtained (this includes consent	for testing	and DNA storag	e)?: ☐ Yes ☐ No

Please refer to the website (http://www.wellingtongenetics.co.nz/) for further information on sample requirements such as transport prerequisites for testing.

<u>Shipping Instructions</u> – Please send specimen with this original form to:

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Wellington Regional Genetics Laboratory Level 6 Ward Support Block Wellington Hospital Riddiford Street WELLINGTON 6021

Phone: 04 9185352

For WRGL use only		
Received by		
Date / Time		
Sample		
Volume / Condition		
Tests required		

Incomplete referral forms may result in a delay in testing and reporting time.