



**Oncology Genetics Referral Form**

|  |                 |   |  |
|--|-----------------|---|--|
| NHI:   | DOB:            | Requester:  | Sample Taken by:<br><br>Date:<br><br>Time: |
| Family Name:   | Sex: F / M      | Print name:   |  |
| Given Name:  | DHB of Domicile | Copy report to:   |  |
| Address:   |                 |   |  |
| <b>Clinical details:</b>   |                 | <b>Sample details:</b>  |  |
| <p><b>Diagnosis:</b> <input type="checkbox"/> Confirmed <input type="checkbox"/> Provisional</p> <p><b>Current state:</b></p> <p><input type="checkbox"/> Diagnosis <input type="checkbox"/> Active</p> <p><input type="checkbox"/> Pre-Treatment <input type="checkbox"/> Relapse</p> <p><input type="checkbox"/> Remission (for MRD study)</p> |                 | <p><input type="checkbox"/> Bone Marrow aspirate:</p> <p style="padding-left: 40px;">Biopsy site: <input type="checkbox"/> Left <input type="checkbox"/> Right</p> <p><input type="checkbox"/> Blood Sample</p> <p><input type="checkbox"/> Lymph node</p> <p><input type="checkbox"/> Solid tumour tissue</p> <p><input type="checkbox"/> Others</p> |  |
| <b>Reason for study/comments/further information:</b>  |                 |   |  |

| Cytogenetic Tests Requested                                  |  | Hold | Proceed Immediately | Comments   |
|--|--|------|---------------------|--|
| <b>Chromosome analysis</b> (LH or sample in transport media) |  |      |                     |  |
| <b>FISH</b>  | <b>FISH analysis</b> from peripheral blood, bone marrow, lymph node or solid tumour samples (LH or sample in transport media).<br>For FFPE slides please refer to separate FFPE referral form) |      |                     | List FISH tests required (please refer to the website for a full list of probes available) <a href="http://www.wellingtongenetics.co.nz/shop/FISH+Test+List.html">http://www.wellingtongenetics.co.nz/shop/FISH+Test+List.html</a> |
|  | <b>FISH for plasma cell disorders</b> CD138+ cell sorting is required.<br><b>Samples must be received within 24 hours.</b>   |      |                     | If plasma cell sorting is required, please indicate the plasma cell percentage ____ % (or contact the laboratory once known)   |
|  | <b>CLL workflow</b> (LH and EDTA or sample in transport media)   |      |                     | This workflow includes <i>TP53</i> FISH. If FISH result is normal, <i>TP53</i> mutations will be interrogated with the Illumina Hot Spot NGS panel.<br><b>Please contact the laboratory if mutation testing is not required.</b>   |

|   |  |  |  |
|---|--|--|--|
| <b>DNA extraction and storage</b><br>(EDTA or sample in transport media). |  |  |  |
|---|--|--|--|

| Molecular Genetics Tests Requested   | Hold | Proceed Immediately | Comments   |
|--|------|---------------------|--|
| <b>MYD88 testing</b><br>(EDTA or sample in transport media)  |      |                     |  |
| <b>Chimerism testing (please circle)</b><br>a. Same sex: EDTA required<br>b. Sex-mismatch: LH required   |      |                     | EDTA is required for same sex as STR analysis is required (this includes donor, pre-transplant or post-transplant).<br>FISH for sex-mismatch   |
| <b>Illumina Hot Spot NGS Panel testing</b> (EDTA, FFPE* or sample in transport media)<br><br>*If NGS required from FFPE sample, please provide 2-3 slides (minimum) cut at 4 microns along with an H&E slide with the area of interest circled<br><br><b>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.</b> |      |                     | Percentage of neoplastic cell nuclei: ____%<br><br>Please note: we require a minimum of 30% neoplastic cells to proceed with NGS<br>Specify genes to interrogate (please circle):<br><br><i>ABL1, AKT1, ALK, APC, ATM, BRAF, CDH1, CDKN2A, CSF1R, CTNNB1, EGFR, ERBB2, ERBB4, EZH2, FBXW7, FGFR1, FGFR2, FGFR3, FLT3, GNA11, GNAQ, GNAS, HNF1A, HRAS, IDH1, IDH2, JAK2, JAK3, KDR, KIT, KRAS, MET, MLH1, MPL, NOTCH1, NPM1, NRAS, PDGFRA, PIK3CA, PTEN, PTPN11, RB1, RET, SMAD4, SMARCB1, SMO, SRC, STK11, TP53, VHL</i> |
| <b>NGS Sendaway</b> (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 storage)?:  Yes  No

Please refer to the website (<http://www.wellingtongenetics.co.nz/>) for further information on sample requirements such as transport prerequisites for testing and the specific coverage information of the Illumina Hot Spot Panel.

**Shipping Instructions – Please send specimen with this original form to:**

**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.**