

PATIENT DETAILS		REQUESTING CLINICIAN DETAILS	
SURNAME:		NAME:	
FIRST NAME:		HOSPITAL / LAB:	
DOB:	<input type="checkbox"/> MALE <input type="checkbox"/> FEMALE	PROVIDER NO:	
ADDRESS:		EMAIL:	
MEDICARE NO:		SIGNATURE:	DATE:

CLINICAL & SAMPLE DETAILS	
<p>CLINICAL NOTES / REASON FOR TEST REQUEST (REQUIRED):</p> <p>PLEASE SEND ALL RELEVANT PATHOLOGY RESULTS (E.G. BONE MARROW REPORT, FBE REPORT, HISTOPATHOLOGY REPORT, ETC.) WITH SAMPLE</p>	<p>SAMPLE TYPE</p> <p><input type="checkbox"/> BONE MARROW <input type="checkbox"/> BLOOD <input type="checkbox"/> TISSUE <input type="checkbox"/> cfDNA</p> <p><input type="checkbox"/> OTHER (PLEASE STATE):</p> <p>COLLECTION DATE (IF ADD-ON REQUEST):</p> <p>TO BE COMPLETED BY COLLECTOR (IF PRIMARY FORM)</p> <p>COLLECTED AND LABELLED BY:</p> <p>SURNAME _____</p> <p>COLLECTION DATE _____ TIME _____</p> <p>SIGNATURE _____</p> <p>I CERTIFY THAT THE PATHOLOGY SPECIMEN AND REQUEST FORM COMPLY WITH MINIMUM LABELLING REQUIREMENTS AND THAT THE SPECIMEN WAS TAKEN FROM THE PATIENT STATED ABOVE AS ESTABLISHED BY DIRECT ENQUIRY AND/OR INSPECTION OF THE IDENTIFICATION BAND AND WAS LABELLED IMMEDIATELY.</p>

AVAILABLE ASSAYS
GENOMIC TESTING SUPPORTED BY SNOWDOME FOUNDATION THROUGH THE WILSON CENTRE FOR LYMPHOMA GENOMICS

NGS GENE PANELS	MUTATION SPECIFIC ASSAYS
<p>GENE LISTS AND FURTHER INFORMATION ON REVERSE</p> <p><input type="checkbox"/> MYELOPROLIFERATIVE NEOPLASM (MPN) DIAGNOSTIC GENE PANEL (8 GENES)</p> <p><input type="checkbox"/> HAEMATOLOGICAL MALIGNANCY (ALLHAEM) GENE PANEL (57 GENES)</p> <p><input type="checkbox"/> SELECT THIS BOX TO EXCLUDE DDX41 ANALYSIS</p> <p>PANEL AVAILABLE ON BLOOD, BM, TISSUE AND cfDNA</p> <p><input type="checkbox"/> SINGLE GENE TP53 ANALYSIS</p> <p><input type="checkbox"/> TESTING TO DETERMINE ORIGIN (SOMATIC VS GERMLINE) OF PREVIOUSLY DETECTED VARIANT</p> <p>PLEASE REFER TO SEPARATE FORM FOR INHERITED BONE MARROW DISORDER NGS PANEL REQUESTS (AVAILABLE ONLINE)</p> <p>PLEASE NOTE: THESE ASSAYS MAY DETECT GERMLINE VARIANTS WITH SIGNIFICANT IMPLICATIONS FOR BOTH THE PATIENT AND THEIR FAMILY. PLEASE ENSURE THAT YOU AND YOUR PATIENT UNDERSTAND THIS POSSIBILITY.</p>	<p><input type="checkbox"/> FLT3-ITD & TKD</p> <p><input type="checkbox"/> NPM1 (NON-QUANTITATIVE)</p> <p><input type="checkbox"/> JAK2 Val617Phe</p> <p><input type="checkbox"/> MYD88 Leu265Pro</p> <p><input type="checkbox"/> BRAF Val600Glu</p>
	QUANTITATIVE PCR (qPCR) ASSAYS
	<p><input type="checkbox"/> t(9;22) BCR-ABL1</p> <p><input type="checkbox"/> t(15;17) PML-RARA</p> <p><input type="checkbox"/> t(8;21) RUNX1-RUNX1T1</p> <p><input type="checkbox"/> inv(16) / t(16;16) CBFβ-MYH11</p> <p><input type="checkbox"/> NPM1 MRD (TYPE A – qPCR; non-TYPE A – NGS)</p>
	OTHER ASSAYS
	<p><input type="checkbox"/> CHIMERISM</p> <p><input type="checkbox"/> IGHV SOMATIC HYPERMUTATION (SHM) ANALYSIS</p> <p><input type="checkbox"/> HAVCR2 GENE VARIANT ANALYSIS</p> <p><input type="checkbox"/> UBA1 GENE MUTATION ANALYSIS</p>

PAYMENT	
<input type="checkbox"/> BILL HOSPITAL / PATHOLOGY PROVIDER	
<input type="checkbox"/> BILL MEDICARE (MUST SIGN BELOW. NON-REBATABLE COMPONENTS WILL BE BILLED TO PATHOLOGY PROVIDED UNLESS OTHERWISE SPECIFIED)	
<p>MEDICARE ASSIGNMENT FORM (SECTION 20A OF THE HIA 1973) I OFFER TO ASSIGN MY RIGHT TO BENEFITS TO THE APPROVED PRACTITIONER WHO WILL RENDER THE REQUESTED PATHOLOGY SERVICE(S) AND ANY ELIGIBLE PATHOLOGICAL DETERMINABLE SERVICE(S) ESTABLISHED NECESSARY BY THE PRACTITIONER.</p>	
PATIENT SIGNATURE: _____	DATE: _____
<p>OR IF PATIENT UNABLE TO SIGN: I AUTHORISE PETER MACCALLUM CANCER CENTRE, WHO WILL RENDER THE REQUESTED PATHOLOGY SERVICES, AND ANY FURTHER PATHOLOGY SERVICES WHICH THE PRACTITIONER DETERMINES TO BE NECESSARY, TO SUBMIT MY UNPAID ACCOUNT TO MEDICARE, SO THAT MEDICARE CAN ASSESS MY CLAIM AND ISSUE ME A CHEQUE MADE PAYABLE TO THE PRACTITIONER, FOR THE MEDICARE BENEFIT. VERBAL CONSENT WAS PROVIDED BY PATIENT TO SUBMIT UNPAID ACCOUNT TO MEDICARE (NO SIGNATURE AVAILABLE) <input type="checkbox"/></p>	
<input type="checkbox"/> BILL PATIENT DIRECTLY (MUST SIGN HERE TO ACKNOWLEDGE COSTS HAVE BEEN DISCUSSED):	PATIENT SIGNATURE: _____
<input type="checkbox"/> BILL OTHER (PLEASE SPECIFY):	

NGS GENE PANEL LISTS

HAEMATOLOGICAL MALIGNANCY GENE PANEL (ALLHAEM) – 57 GENES

<i>ABL1</i>	<i>ARAF</i>	<i>ASXL1</i>	<i>BCL2</i>	<i>BIRC3</i>	<i>BRAF</i>	<i>BTK</i>	<i>CALR</i>
<i>CARD11</i>	<i>CBL</i>	<i>CD274</i>	<i>CD79B</i>	<i>CEBPA</i>	<i>CSF3R</i>	<i>CXCR4</i>	<i>DDX41*</i>
<i>DNMT3A</i>	<i>ETNK1</i>	<i>EZH2</i>	<i>FLT3**</i>	<i>FYN</i>	<i>GATA1</i>	<i>GATA2</i>	<i>ID3</i>
<i>IDH1</i>	<i>IDH2</i>	<i>IRF8</i>	<i>JAK2</i>	<i>JAK3</i>	<i>KIT</i>	<i>KRAS</i>	<i>MAP2K1</i>
<i>MPL</i>	<i>MYD88</i>	<i>NOTCH1</i>	<i>NPM1</i>	<i>NRAS</i>	<i>PDCD1LG2</i>	<i>PIGA</i>	<i>PFH6</i>
<i>PLCG1</i>	<i>PLCG2</i>	<i>RHOA</i>	<i>RUNX1</i>	<i>SETBP1</i>	<i>SF3B1</i>	<i>SH2B3</i>	<i>SRSF2</i>
<i>STAT3</i>	<i>STAT5B</i>	<i>STAT6</i>	<i>TCF3</i>	<i>TET2</i>	<i>TP53</i>	<i>U2AF1</i>	<i>XPO1</i>
<i>ZRSR2</i>							

**DDX41* VARIANT ANALYSIS IS EXCLUDED ON REQUEST. CLINICALLY SIGNIFICANT VARIANTS IN THIS GENE ARE LIKELY TO BE OF GERMLINE ORIGIN.

***FLT3*-ITD ANALYSIS PERFORMED USING A SEPARATE ASSAY AND INCLUDED WHEN CLINICALLY APPROPRIATE OR REQUESTED.

THE HAEMATOLOGICAL MALIGNANCY GENE PANEL HAS BEEN DESIGNED TO PROVIDE DIAGNOSTIC, PROGNOSTIC AND THERAPEUTIC INFORMATION ACROSS THE SPECTRUM OF HAEMATOLOGICAL MALIGNANCY AND IS AVAILABLE FOR ORDER ON THE FOLLOWING SAMPLE TYPES: PERIPHERAL BLOOD, BONE MARROW ASPIRATE, FRESH OR FFPE TISSUE AND cfDNA. SEE SAMPLE REQUIREMENTS BELOW.

MYELOPROLIFERATIVE NEOPLASM (MPN) DIAGNOSTIC GENE PANEL – 8 GENES

<i>JAK2</i>	<i>CALR</i>	<i>MPL</i>	<i>SF3B1</i>	<i>KIT</i>	<i>CSF3R</i>	<i>ETNK1</i>	<i>SH2B3</i>
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SAMPLE REQUIREMENTS

GENE SPECIFIC ASSAYS, NGS ASSAYS & IGHV SHM ASSAY	<ul style="list-style-type: none"> ○ 4ml PERIPHERAL BLOOD (EDTA) ○ 1-2ml BONE MARROW ASPIRATE (EDTA) ○ TISSUE: FRESH / FROZEN / FFPE BLOCK / FFPE SLIDES – PLEASE PROVIDE HISTOPATHOLOGY REPORT ○ DNA (MINIMUM 10µL at >50ng/µL) ○ CELL FREE DNA (cfDNA) (EXCLUDING IGHV ANALYSIS): 10ml PERIPHERAL BLOOD IN STRECK TUBE SAMPLES FOR cfDNA MUST BE RECEIVED WITHIN 72 HOURS OF COLLECTION ○ OTHER (E.G. CSF) – PLEASE CALL / EMAIL TO DISCUSS PRIOR TO SENDING ○ HAIR (FOR FOLLOW UP TESTING OF PREVIOUSLY DETECTED VARIANTS ONLY) – PLEASE CALL / EMAIL TO DISCUSS PRIOR TO SENDING
QPCR ASSAYS	<ul style="list-style-type: none"> ○ 20ml PERIPHERAL BLOOD (EDTA) ○ 1-2ml BONE MARROW ASPIRATE (EDTA) <p>ALL SAMPLES MUST BE RECEIVED WITHIN 48 HOURS OF COLLECTION</p>
CHIMERISM	<ul style="list-style-type: none"> ○ 4ml (PRE-TRANSPLANT) / 18ml (POST-TRANSPLANT) PERIPHERAL BLOOD (EDTA) ○ 1-2ml BONE MARROW ASPIRATE (EDTA) <p>POST-TRANSPLANT SAMPLES MUST BE RECEIVED WITHIN 24 HOURS OF COLLECTION</p>

ADDRESS & CONTACT DETAILS

PLEASE SEND SPECIMEN AND COMPLETED FORM TO:

PATHOLOGY – SPECIMEN RECEPTION (LEVEL 4)
PETER MACCALLUM CANCER CENTRE
305 GRATTAN STREET
MELBOURNE VIC 3000

TEL: +61 3 8559 7284

EMAIL: MOLECULAR.HAEMATOLOGY@PETERMAC.ORG

FAX: +61 3 8559 5437

WEBSITE: www.petermac.org/about/signature-centres/centre-clinical-cancer-genomics/molecular-haematology