

Genetics Laboratory - Credit Valley Hospital

2200 Eglinton Ave West, Room 2H144 Mississauga ON, L5M 2N1 Tel: (905) 813-1100 ext. 6288

Fax: (905) 813-3854

ONCOLOGY GENETIC TESTING REQUISITION

~ Visit community collection lab for blood draw. ~

PATIENT DEMOGRAPHICS	viole dominarity donor	Short has for shoot draw.		
Last Name:		Health Card #:	Version Code:	
First Name:		Date of Birth (DD/MM/YYYY):		
Address:		Legal Sex: ☐ Male ☐ Fer	nale □ Non-Binary	
City:		☐ Unknown ☐ X		
Province:		Phone Number:		
Postal Code:		Email Address:		
REFERRING PROVIDER		COPIES TO		
Last Name:		Last Name:		
First Name:		First Name:		
CPSO #:	OHIP Billing #:	CPSO #:	OHIP Billing #:	
Address:		Address:		
City/Prov:	Postal Code:	City/Prov:	Postal Code:	
Phone:	Fax:	Phone:	Fax:	
Signature (required):	Date:			
SPECIMEN REQUIREMENTS				
SPECIMEN COLLECTION DATE	E (DD/MM/YYYY):	TIME (HH:MM)):	
Molecular Genetics and Microarray		Cytogenetics		
☐ Blood (5-10mL EDTA, room temp)		☐ Bone Marrow (3mL NaHep, room temp); attach CBC report		
☐ Bone Marrow (3mL EDTA, room temp); attach CBC report		□ Blood (3mL NaHep, room temp); attach CBC report		
□ Extracted DNA Source: (2µg total; min.70ng/uL)		☐ Cell Suspension Source:		
□ Paraffin-Embedded (FFPE) Tissue Specimen #/ Block #: Tumour Cellularity in circled H&E region: -include circled H&E -air dry 10 x 4um sections/test ordered on uncoated slidesattach corresponding pathology report □ Tissue (fresh) Source:		□ Paraffin Embedded (FFPE) Tissue Slide Specimen #/ Block #: Tumour Cellularity in circled H&E region: - include circled H&E - air dry 2 x 4µm sections/probe on positively charged slides. - attach corresponding pathology report		
CLINICAL INFORMATION				
☐ Has patient had a blood transfusion in the last 30 days ☐ Bone Marrow Transplant recipient? Sex of donor: ☐ Male ☐ Female				
TESTING STATUS				
□ Routine □ Expedited: Reason (required):		_ Surgery Date (if applicable):		
THP - LAB USE ONLY				
Date / Time Received:		FOR THP L	_ABEL ONLY	



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CYTOGENETICS				
TEST REQUESTED	REASON FOR REFERRAL / FISH PROBE REQUEST			
☐ Karyotyping	□ AML	☐ Lymphoma		
(Chromosome analysis)	□ ALL	□ MDS		
	□ CML	☐ MPN (PV, ET,MF, etc)		
	□ CLL	☐ Multiple Myeloma		
	□ Eosinophilia	☐ Other:		
☐ Hematologic FISH	☐ Multiple Myeloma FISH panel CDKN2C (1p); CKS1B(1q); TP53 (17p); D17Z1 (17cent); IGH::MYEOV; with reflex to IGH::FGFR3; IGH::MAF; IGH::MAFB as needed			
	☐ Diffuse Large B-Cell Lymphoma Panel MYC (with reflex to BCL2 and BCL6 as needed)	☐ MYC only (8q24)		
	☐ CCND1::IGH t(11;14)	☐ <i>BCL2</i> only (18q21)		
	□ PDGFRA (4q12)	☐ <i>BCL6</i> only (3q27)		
	□ PDGFRB (5q32)	☐ <i>PML::RARA</i> t(15;17)		
☐ Paraffin-Embedded (FFPE) FISH	☐ Diffuse Large B-Cell Lymphoma Panel MYC (with reflex to BCL2 and BCL6 as needed)			
	☐ MYC only (8q24)			
	☐ <i>BCL2</i> only (18q21)			
	☐ <i>BCL6</i> only (3q27)			
MOLECULAR GENETICS and	MICROARRAY			
MOLECULAR HEMATOLOG				
☐ CML Diagnostic				
☐ CML MRD Follow Up	Require CML specimen within 24 hours of collection along with CBC report			
☐ CLL microarray panel (chromosomes: 11q, 12, 13q, 17p)				
□ B Cell Clonality				
☐ T Cell Clonality				
☐ Hemochromatosis (<i>HFE</i>) Co	ommon Variants			
☐ Hairy Cell Leukemia (BRAF)				
□ JAK2 / CALR				
□ Thrombophilia Common Variant Panel (<i>Factor II</i> , <i>Factor V</i>)				
☐ Myeloid NGS Panel¹ (Pathology report required. Bone Marrow preferred. Contact laboratory if blood testing is required). ¹ Sequence and copy number analysis on the following: ABL1 ASXL1 BCOR BCORL1 BRAF CALR CBL CDKN2A CEBPA CSF3R CUX1 DDX41 DNMT3A ETV6 EZH2 FLT3 GATA2 GNAS HRAS IDH1 IDH2 JAK2 JAK3 KIT KMT2A KRAS MPL NF1 NPM1 NRAS PHF6 PIGA PPM1D PRPF8 PTEN PTPN11 RAD21 RRAS2 RUNX1 SETBP1 SF3B1 SH2B3 SMC3 SRSF2 STAG2 STAT3 STAT5B TET2 TP53 U2AF1 WT1 ZRSR2				
DUADMACOCENOMICS				
PHARMACOGENOMICS				



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Genome Diagnostics - SOLID TUMOUR

Test	Regions of Interest		
☐ Bladder NGS Panel ^{1,2}	0 (1)00 D		
☐ Breast NGS Panel 1,2,3,4	Gene content of NGS Panels Sequence and copy number analysis of all genes is performed,		
☐ BRAF/MLH1 Specify tissue type:	unless otherwise indicated. Partner-agnostic fusion analyses are performed for all RNA panels. The specific exons targeted for each gene is available upon request. 1 Solid Tumour DNA Panel: AKT1 ALK BRAF CCND1 CTNNB1 DDR2 EGFR EIF1AX ERBB2 FGFR1 FGFR2 FGFR3 GNAS HRAS IDH1 IDH2 KIT KRAS MAP2K1 MDM2 MET NRAS PDGFRA PIK3CA PTEN RET ROS1 STK11 TERT TP53 TSHR 2 Solid Tumour RNA Fusion Panel: ALK BRAF EGFR FGFR1 FGFR2 FGFR3 KRAS MET NRG1		
☐ Colorectal NGS panel ¹			
☐ Endometrial NGS Panel ^{1,3}			
☐ Endometrial <i>MLH1</i> Methylation			
☐ GIST NGS Panel ¹			
☐ Hepatobiliary/Cholangiocarcinoma NGS Panel ^{1,2}			
☐ Lung NGS Panel ^{1,2,3}			
☐ Lung Resistance Mutations ^{1,2,3}	NTRK1 NTRK2 NTRK3 RET ROS1		
☐ Melanoma NGS Panel ¹	³ Additional DNA Analysis:		
□ Ovarian (BRCA1/BRCA2)	ATRX DICER1 ESR1 FOXL2 GNA11 GNAQ H3F3A HIST1H3B POLE SMARCA4		
☐ Ovarian SCCOHT NGS Panel 1,3	4 Additional RNA Fusion Analysis: ESR1 PPARG RELA YAP1 ZFTA		
☐ Prostate NGS Panel (ATM, BRCA1, BRCA2, PALB2)			
☐ Sex Cord Stromal NGS Panel ^{1,3}			
☐ Solid Tumour Fusion Panel (<i>NTRK</i> , <i>etc</i>) ^{2,4}			
☐ Thyroid NGS Panel ^{1,2,4}			
☐ Uveal Melanoma NGS Panel ^{1,3}			
Central Nervous System (CNS) SOLID TUMOURS			
☐ <i>MGMT</i> Methylation			
☐ CNS Microarray Panel (1p/19q, 1q, 6, 7, 10, <i>MYCN</i> , <i>EGFR</i> , <i>CDKN2A/2B</i>)			
☐ CNS NGS Panel 1,3			
☐ CNS NGS Fusion Panel ^{2,4}			