

Hayward Genetics Center Requisition Form Deliver Specimens To: 1430 Tulane Ave. Room 5301 New Orleans, LA 70112 PHONE 504-988-5101 FAX 504-988-1763 Biochemical Genetics beeper 504-501-6011		Reports To: Fax To #: Phone #: Billing Address:																																																																																																																																																									
PATIENT INFORMATION		SAMPLE INFORMATION		CLINICAL ORDER INFORMATION																																																																																																																																																							
Last Name		Date Collected		Hosp/Clinic:																																																																																																																																																							
First Name		Specimen Type ☐ Peripheral Blood ☐ Products of Conception ☐ Amniotic Fluid ☐ Bone marrow (WBC_____) ☐ Leukemic blood (WBC_____) ☐ Lymph node ☐ Tissue (fresh) ☐ Paraffin-embedded tissue ☐ Other _____		Ordering Physician:																																																																																																																																																							
Date of Birth				Diagnosis/Clinical Info./Indication:																																																																																																																																																							
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MRN																																																																																																																																																											
Molecular Genetics																																																																																																																																																											
Disease-Specific Tests ☐ Chromosomal Microarray (aCGH+SNP) ☐ Fragile X Syndrome ☐ Prader-Willi/Angelman Syndrome (Methylation PCR)																																																																																																																																																											
HemeOnc Tests ☐ JAK2 V617F mutation ☐ Myeloid NGS Panel<table><tr><td>ABL1</td><td>ANKRD26</td><td>ARID1A</td><td>ASXL1</td><td>ASXL2</td><td>ATM</td><td>ATRX</td><td>BCOR</td><td>BCORL1</td><td>BRAF</td></tr><tr><td>CALR</td><td>CBL</td><td>CBLB</td><td>CBLCL</td><td>CCND2</td><td>CDKN2A</td><td>CDKN2B</td><td>CEBPA</td><td>CREBBP</td><td>CRLF2</td></tr><tr><td>CSF3R</td><td>CUX1</td><td>DDX41</td><td>DNMT3A</td><td>EP300</td><td>ETNK1</td><td>ETV6</td><td>EZH2</td><td>FLT3</td><td>GATA1</td></tr><tr><td>GATA2</td><td>GNAS</td><td>HRA5</td><td>IDH1</td><td>IDH2</td><td>IKZF1</td><td>JAK1</td><td>JAK2</td><td>JAK3</td><td>KIT</td></tr><tr><td>KMT2A</td><td>KRAS</td><td>MPL</td><td>NF1</td><td>NPM1</td><td>NRAS</td><td>PDGFRA</td><td>PHF6</td><td>PPM1D</td><td>PTEN</td></tr><tr><td>PTPN11</td><td>RAD21</td><td>RB1</td><td>RUNX1</td><td>SETBP1</td><td>SF3B1</td><td>SH2B3</td><td>SMC1A</td><td>SMC3</td><td>SRP72</td></tr><tr><td>SRSF2</td><td>STAG2</td><td>STAT3</td><td>TET2</td><td>TP53</td><td>U2AF1</td><td>WT1</td><td>ZRSR2</td><td></td><td></td></tr></table> ☐ Lymphoid NGS Panel<table><tr><td>ABL1</td><td>ANKRD26</td><td>ARID1A</td><td>ATM</td><td>ATRX</td><td>B2M</td><td>BCL2</td><td>BCL6</td><td>BCOR</td><td>BCORL1</td></tr><tr><td>BIRC3</td><td>BRAF</td><td>BTX</td><td>CARD11</td><td>CBLB</td><td>CBLCL</td><td>CCND1</td><td>CCND2</td><td>CCND3</td><td>CD58</td></tr><tr><td>CD79A</td><td>CD79B</td><td>CDKN2A</td><td>CDKN2B</td><td>CREBBP</td><td>CRLF2</td><td>CUX1</td><td>CXCR4</td><td>DDX41</td><td>DNMT3A</td></tr><tr><td>EP300</td><td>ETV6</td><td>EZH2</td><td>FBXW7</td><td>FLT3</td><td>FOXO1</td><td>GNA13</td><td>GNAS</td><td>HRA5</td><td>ID3</td></tr><tr><td>IKZF1</td><td>IL7R</td><td>JAK1</td><td>JAK3</td><td>KDM6A</td><td>KLF2</td><td>KMT2A</td><td>KMT2D</td><td>KRAS</td><td>MAP2K1</td></tr><tr><td>MEF2B</td><td>MYC</td><td>MYD88</td><td>NFKBIE</td><td>NOTCH1</td><td>NOTCH2</td><td>NRAS</td><td>PAX5</td><td>PHF6</td><td>PIM1</td></tr><tr><td>PLCG2</td><td>POT1</td><td>PRDM1</td><td>PTEN</td><td>RB1</td><td>RHOA</td><td>RUNX1</td><td>SETD2</td><td>SF3B1</td><td>SH2B3</td></tr><tr><td>SOC51</td><td>STAT3</td><td>STAT5B</td><td>STAT6</td><td>TCF3</td><td>TET2</td><td>TNFAIP3</td><td>TNFRSF14</td><td>TP53</td><td>XPO1</td></tr></table>						ABL1	ANKRD26	ARID1A	ASXL1	ASXL2	ATM	ATRX	BCOR	BCORL1	BRAF	CALR	CBL	CBLB	CBLCL	CCND2	CDKN2A	CDKN2B	CEBPA	CREBBP	CRLF2	CSF3R	CUX1	DDX41	DNMT3A	EP300	ETNK1	ETV6	EZH2	FLT3	GATA1	GATA2	GNAS	HRA5	IDH1	IDH2	IKZF1	JAK1	JAK2	JAK3	KIT	KMT2A	KRAS	MPL	NF1	NPM1	NRAS	PDGFRA	PHF6	PPM1D	PTEN	PTPN11	RAD21	RB1	RUNX1	SETBP1	SF3B1	SH2B3	SMC1A	SMC3	SRP72	SRSF2	STAG2	STAT3	TET2	TP53	U2AF1	WT1	ZRSR2			ABL1	ANKRD26	ARID1A	ATM	ATRX	B2M	BCL2	BCL6	BCOR	BCORL1	BIRC3	BRAF	BTX	CARD11	CBLB	CBLCL	CCND1	CCND2	CCND3	CD58	CD79A	CD79B	CDKN2A	CDKN2B	CREBBP	CRLF2	CUX1	CXCR4	DDX41	DNMT3A	EP300	ETV6	EZH2	FBXW7	FLT3	FOXO1	GNA13	GNAS	HRA5	ID3	IKZF1	IL7R	JAK1	JAK3	KDM6A	KLF2	KMT2A	KMT2D	KRAS	MAP2K1	MEF2B	MYC	MYD88	NFKBIE	NOTCH1	NOTCH2	NRAS	PAX5	PHF6	PIM1	PLCG2	POT1	PRDM1	PTEN	RB1	RHOA	RUNX1	SETD2	SF3B1	SH2B3	SOC51	STAT3	STAT5B	STAT6	TCF3	TET2	TNFAIP3	TNFRSF14	TP53	XPO1
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Prenatal Studies *Gestational Age_____wks by ☐ LMP ☐ Ultrasound ☐ FISH for Aneuploidy (chr 13, 18, 21, X, Y) ☐ Chromosome analysis ☐ Other:		Peripheral Blood Studies ☐ Routine chromosome analysis ☐ High resolution chromosome analysis		Fluorescence In Situ Hybridization (Fish) Syndrome del(5)(p15) del(22)(q11.2) del(15)(q11-q13) del(17)(p11.2) del(7)(q11.23) Other FISH study: _____																																																																																																																																																							
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