

## Analyseauftrag für Molekulare Diagnostik - Hämatologie Teil 1 / 2

<b>Patientendaten:</b> stationär      KV                      Privat              ASV Krankenkasse bzw. Kostenträger: Name: Geburtsdatum: Diagnose:	<b>Materialnummer:</b> <b>Probentyp:</b> FFPE (Block, Schnitte) EDTA-Blut (Peripher, Knochenmark) Anderes (spezifizieren)
<b>Myeloische Erkrankungen (MPN, MDS, AML etc.)</b> <b>NGS-Panel Myeloid</b> (ABL1, ASXL1,CALR, CBL, CEBPA, CSF3R, DNMT3A, EZH2, FLT3, IDH1/2, JAK2/3, KIT, KRAS, MPL, NPM1, PTPN11, RUNX1, SETBP1, SF3B1, SRSF2, TET2, TP53, U2AF1) <b>NGS-Panel AML-Groß</b> (ASXL1, CEBPA, DNMT3A, FLT3, KIT, IDH1/2, NOTCH1, NPM1, RUNX1, SETBP1, SF3B1, SRSF2, TET2, TP53) <b>NGS-Panel AML-Klein</b> (CEBPA, FLT3, IDH1/2, NPM1) <b>MDS-Genpanel - IPSS-M (Groß)</b> (ASXL1, BCOR, CBL, EZH2, ETV6, DNMT3A, FLT3, IDH1, IDH2, KRAS, MLL/KMT2A, NPM1, NRAS, RUNX1, SF3B1, SRSF2, TP53, U2AF1) <b>MDS-Genpanel - IPSS-M (Klein)</b> (CBL, DNMT3A, ETV6, KRAS, NRAS) <b>Genfusionen (AMLplex: AML1-ETO, BCR-ABL, CALM-AF10, CBFβ- MYH11, DEK-CAN, MLL-AF6/AF9/ELL/PTD, NPM1-MLF1, PML-RARA)</b> (nur am Blut) <b>Einzelgentests</b> <b>AML - Akute myeloische Leukämie</b> IDH1-Mutation (Hotspots) - Sequenz. RUNX1_RUNX1T1-Fusion [t(8;21)] - Realtime-PCR FLT3-Duplikation/Mutation - Fragmentanalyse/RFLP KMT2A/MLL-Translokation - FISH <b>MPN - Myeloproliferative Neoplasie</b> JAK2-Mutation (V617F) - Sequenz./Droplet PCR JAK2-Mutation (Exon 12) - Sequenz. BCR-ABL t(9;22) -Translokation - FISH MPL-Mutation (W515L, W515K) - Sequenz./ Droplet-PCR Calreticulin-Mutation - Fragmentanalyse FGFR1-Translokation - FISH FIP1L1-PDGFRα-Translokation - FISH <b>MDS - Myelodysplastische Neoplasie</b> 5q (EGR1)-Deletion - FISH PDGFRB-Translokation - FISH	<b>CLL - Chronische Lymphatische Leukämie</b> <b>NGS-Panel CLL</b> (ATM, BCOR, BIRC3, BRAF, BTK, EGR2, FBXW7, KRAS, NRAS, MYD88, NOTCH1, PLCG2, POT1, SAMHD1, SF3B1, TP53, XPO1) <b>Einzelgentests</b> TP53-Mutation - Sequenz. 17p (TP53) / 11q (ATM) - Deletion - FISH IGHV-Mutationsstatus - Sequenz./Lymphotrack 13q14-Deletion - FISH Trisomie 12 - FISH
<b>MRD-Diagnostik</b> BCR-ABL p210 -Translokation - Droplet PCR	<b>Lymphomdiagnostik</b> <b>NGS-Panel B-Zelle</b> (CARD11, CD79B, CREBBP, CXCR4, EZH2, GNA13, MYD88, PIM1, PRDM1, SOCS1, STAT6, TNFAIP3, TNFRSF14) <b>NGS-Genpanel TCL</b> (ABCC9, ARID1A, ATM, BCOR, CARD11, CD28, DNMT3A, FYN, IDH2, JAK1/3, KMT2D, MSC, NCOR, PLCG1, PTEN, RHOA, SETD1B, SETD2, STAT3/5B, TET2, TP53) <b>NGS-Genpanel CTCL</b> (ARID1A, BCOR, CARD11, CD28, DNMT3A, MSC, NCOR, PLCG1, RHOA, SETD2, STAT3/5b, TET2, TP53) <b>NK-Zell-Panel (NK-LGLL)</b> (ASXL1, ATM, CCL22, DNMT3A, STAT3/5b, TET2, TNFAIP3) <b>Einzelgentests</b> STAT3/5b - Sequenz. B-Zell-Klonalität - Fragmentanalyse T-Zell-Klonalität - Fragmentanalyse κ/λ-Leichtketten - CISH <b>FL - Follik. Lymph. &amp; DLBCL - Diff. großzell. B-Zell-Lymph.</b> BCL2-IGH-Translokation - FISH BCL6 -Translokation - FISH MYC-IGH-Translokation - FISH MYC-Translokation - FISH <b>MALT-Lymphom</b> BIRC3-MALT1-Translokation - FISH <b>MCL - Mantelzell-Lymphom</b> CCND1-IGH-Translokation - FISH 17p (TP53)-Deletion - FISH <b>Mb. Waldenström (Lymphoplasmozyt. Lymphom)</b> MYD88-Mutation - Droplet-PCR CXCR4-Mutation (Codon 338) - Droplet-PCR
<b>MM - Multiples Myelom (Plasmozytom)</b> BRAF (V600Mutation) - Sequenz. KRAS-/NRAS-Mutation (Exon 2, 3, 4) - Sequenz. 17p (TP53)-Deletion - FISH CCND1-IGH-Translokation - FISH CDKN2C-Deletion - FISH CKS1B-Gain - FISH FGFR3-IGH-Translokation - FISH MAF-IGH-Translokation - FISH MAFB-IGH-Translokation - FISH	<b>Mastozytose</b> <b>NGS-Panel Mastozytose</b> (ASXL1, CBL, EZH2, IDH1/2, JAK2, KIT, KRAS, NRAS, RUNX1, SF3B1, SRSF2, TET2) <b>Einzelgentests</b> KIT-Mutation (D816V) - Droplet-PCR, RT-PCR KIT-Mutation (Ex. 8-11, 13, 17, 18) - Sequenz.
<b>HZL - Haarzell-Leukämie</b> BRAF (V600-Mutation) - Sequenz.	<b>HES - Hypereosinophiles Syndrom</b> FIP1L1-PDGFRα-Translokation - FISH PDGFRB-Translokation - FISH FGFR1-Translokation - FISH CBFβ-Translokation - FISH
<b>Hämochromatose</b> HFE-Mutation (C282Y, H63D, S65C) - Sequenz.	(Empty space for signature)
Tel. für Rücksprache:	Unterschrift: