

## Analyseauftrag für Molekulare Diagnostik - Hämatologie

<p><b>Patientendaten:</b></p> <p>stationär                      KV                                      privat</p> <p>Krankenkasse bzw. Kostenträger:</p> <p>Name:</p> <p>Geburtsdatum:</p> <p>Diagnose:</p>	<p><b>Materialnummer:</b></p> <p><b>Probestyp:</b></p> <p>FFPE (Gewebe, Schnitte)</p> <p>Blut (EDTA, Knochenmark)</p> <p>Anderes (spezifizieren)</p>
<p><b>Myeloische Erkrankungen (MPN, MDS, AML etc.)</b></p> <p><b>NGS-Panel Myeloid</b> (ASXL1, CALR, CBL, CEBPA, CSF3R, DNMT3A, EZH2, FLT3, IDH1/2, JAK2/3, KIT, KRAS, MPL, NPM1, NTRK1, PTPN11, RUNX1, SETBP1, SF3B1, SRSF2, TET2, TP53, U2AF1)</p> <p><b>NGS-Panel AML-Groß</b> (ASXL1, CEBPA, DNMT3A, FLT3, KIT, IDH1/2, NPM1, RUNX1, SETBP1, SF3B1, SRSF2, TET2, TP53)</p> <p><b>NGS-Panel AML-Klein</b> (CEBPA, FLT3, IDH1/2, NPM1)</p> <p><b>NGS-Panel MDS</b> (ASXL1, DNMT3A, RUNX1, SETBP1, SRSF2, TET2, TP53)</p> <p><b>Genfusionen</b> (AMLplex: AML1-ETO, BCR-ABL, CALM-AF10, CBFβ-MYH11, DEK-CAN, MLL-AF6/AF9/ELL/PTD, NPM1-MLF1, PML-RARA)</p> <p><b>Einzelgentests</b></p> <p><b>AML - Akute myeloische Leukämie</b></p> <p>RUNX1_RUNX1T1-Fusion [t(8;21)] - Realtime-PCR</p> <p>FLT3-Duplikation/Mutation – Fragmentanalyse/RFLP</p> <p><b>MPN - Myeloproliferative Neoplasie</b></p> <p>JAK2-Mutation (V617F) - Sequenz./Droplet PCR</p> <p>JAK2-Mutation (Exon 12) - Sequenz.</p> <p>BCR-ABL t(9;22) -Translokation - FISH</p> <p>MPL-Mutation (W515L, W515K) - Sequenz./ Droplet-PCR</p> <p>Calreticulin-Mutation - Fragmentanalyse</p> <p>FGFR1-Translokation - FISH</p> <p>FIP1L1-PDGFRα-Translokation - FISH</p> <p><b>MDS - Myelodysplastisches Syndrom</b></p> <p>5q (EGR1)-Deletion - FISH</p> <p>PDGFRβ-Translokation - FISH</p>	<p><b>CLL - Chronische Lymphatische Leukämie</b></p> <p><b>NGS-Panel CLL</b> (ATM, BCOR, BIRC3, BRAF, BTK, EGR2, FBXW7, KRAS, NRAS, MYD88, NOTCH1, PLCG2, POT1, SAMHD1, SF3B1, TP53, XPO1)</p> <p><b>Einzelgentests</b></p> <p>TP53-Mutation - Sequenz.</p> <p>17p (TP53) / 11q (ATM) - Deletion - FISH</p> <p>IGHV-Mutationsstatus - Sequenz./Lymphotrack</p> <p>13q14-Deletion - FISH</p> <p>Trisomie 12 - FISH</p>
<p><b>HES - Hypereosinophiles Syndrom</b></p> <p>FIP1L1-PDGFRα-Translokation - FISH</p> <p>PDGFRβ-Translokation - FISH</p> <p>FGFR1-Translokation - FISH</p> <p>CBFB-Translokation - FISH</p>	<p><b>Lymphomdiagnostik</b></p> <p><b>NGS-Panel B-Zelle</b> (CARD11, CD79B, CREBBP, EZH2, GNA13, MYD88, PIM1, PRDM1, SOCS1, STAT6, TNFAIP3, TNFRSF14)</p> <p><b>NGS-Genpanel TCL</b> (CARD11, CD28, DNMT3A, IDH2, JAK1/3, MSC, NCOR, PLCG1, RHOA, SETD2, STAT3/5b, TET2, TP53)</p> <p><b>NGS-Genpanel CTCL</b> (ARID1A, BCOR, CARD11, CD28, DNMT3A, MSC, NCOR, PLCG1, RHOA, SETD2, STAT3/5b, TET2, TP53)</p> <p><b>Einzelgentests</b></p> <p>STAT3/5b - Sequenz.</p> <p>B-Zell-Klonalität - Fragmentanalyse</p> <p>T-Zell-Klonalität - Fragmentanalyse</p> <p>κ/λ-Leichtketten - CISH</p>
<p><b>Mastozytose</b></p> <p><b>NGS-Panel Mastozytose</b> (ASXL1, CBL, EZH2, IDH1/2, JAK2, KIT, KRAS, NRAS, RUNX1, SF3B1, SRSF2, TET2)</p> <p><b>Einzelgentests</b></p> <p>KIT-Mutation (D816V) - Droplet-PCR, RT-PCR</p> <p>KIT-Mutation (Ex. 8-11, 13, 17, 18) - Sequenz.</p>	<p><b>FL - Follikuläres Lymphom &amp; DLBCL - Diffus großzelliges B-Zell-Lymphom</b></p> <p>BCL2-IGH-Translokation - FISH</p> <p>BCL6 -Translokation - FISH</p> <p>MYC-IGH-Translokation - FISH</p> <p>MYC-Translokation - FISH</p>
<p><b>MM - Multiples Myelom (Plasmozytom)</b></p> <p>BRAF (V600Mutation) - Sequenz.</p> <p>KRAS-NRAS-Mutation (Exon 2, 3, 4) - Sequenz.</p> <p>17p (TP53)-Deletion - FISH</p> <p>CCND1-IGH-Translokation - FISH</p> <p>CDKN2C-Deletion - FISH</p> <p>CKS1B-Gain - FISH</p> <p>FGFR3-IGH-Translokation - FISH</p> <p>MAF-IGH-Translokation - FISH</p> <p>MAFB-IGH-Translokation - FISH</p>	<p><b>MCL - Mantelzell-Lymphom</b></p> <p>CCND1-IGH-Translokation - FISH</p> <p>17p (TP53)-Deletion - FISH</p>
<p>Tel. für Rücksprache:</p>	<p><b>Mb. Waldenström (Lymphoplasmozyt. Lymphom)</b></p> <p>MYD88-Mutation - Droplet-PCR</p> <p>CXCR4-Mutation (Codon 338) - Droplet-PCR</p> <p><b>MALT-Lymphom</b></p> <p>BIRC3-MALT1-Translokation - FISH</p> <p><b>HZL - Haarzell-Leukämie</b></p> <p>BRAF (V600-Mutation) - Sequenz.</p> <p><b>Hämochromatose</b></p> <p>HFE-Mutation (C282Y, H63D, S65C) - Sequenz.</p> <p><b>Anderes</b></p> <p>Unterschrift:</p>