

Analyseauftrag für Molekulare Diagnostik - Hämatologie

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