



Physician information:

Patient:

Request form for oncological laboratory diagnostics

Diagnosis:

- CML B-CLL B-ALL
 MPN B-NHL T-ALL
 MDS T-NHL
 AML MM/Plasmocytoma

Level of therapy:

- Initial diagnosis
 Control
 Recurrence
 after BMT, donor ♀ ♂

Type of sample:

- Peripheral blood
 Bone marrow

Other clinical comments:

Sampling date / time:

Signature:

Please prepare the following documents and samples for shipment:

- Request form
- Sample labeled with patient name and date of birth:
 - Karyogram and FISH: at least 5ml heparin blood resp. 3ml heparin bone marrow
 - Molecular genetics: 3ml EDTA blood resp. 3ml EDTA bone marrow
- Declaration of consent signed by the patient according to the German Genetic Diagnostics Act (GenDG) (only required for analyses under the heading "Other tests")

Patient name:

Date of birth:

KARYOGRAM

- Microscopic chromosome analysis

B-NHL/CLL

- FISH: NHL panel
- FISH: CLL panel
- SNP: Specific mutation analysis – Lymphoma panel (incl. BRAF(600), MYD88(265), NRAS(12/13/59/61/117/146), SF3B1(666), MYC(74/164))
- MG: TP53
- MG: IGHV (clonality (Lymphoma), Hypermutation status (CLL))
- MG: TP53, BIRC3, NOTCH1, SF3B1, (e.g. for therapy indication)

Multiple Myeloma / Plasmocytoma

- FISH: MM panel
- SNP: MM panel (incl. BRAF(600), NRAS(12/13/59/61/117/146), KRAS(12/13/59/61/117/146))

Waldenstrom's disease

- MG: CXCR4, MYD88

Burkitt's lymphoma

- FISH: BL panel

DLBCL

- FISH: DLBCL panel

Follicular lymphoma

- FISH: FL panel (incl. t(14;18))
- FISH / PCR: t(14;18) IGH/BCL2

Mantle cell lymphoma

- FISH: MCL panel (incl. t(11;14))
- FISH / PCR: t(11;14) CCND1/IGH

Marginal zone lymphoma

- FISH: MZL panel

Hairy cell leukemia

- FISH: HCL panel

T-cell lymphoma

- FISH: TCL panel

ALL

- PCR / FISH: t(9;22) bcr-abl1 (qualitative, diagnostic)
- PCR: t(1;19) PBX1/E2A
- PCR: t(4;11) AF4/MLL

CML

- FISH / PCR: t(9;22) bcr-abl1 (qualitative, diagnostic)
- PCR: t(9;22) bcr-abl1 (quantitative, therapy monitoring)
- MG: abl1 sequencing (TKI resistance resp. therapy failure)

MPN

- FISH: MPN panel
- FISH: Eosinophilia panel (MPN/MDS)
- SNP: Specific mutation analysis - MPN panel (incl. JAK2(V617F), CALR(364fs), MPL(505/510/515), IDH1(132), IDH2(140/172), EZH2(646/690), SRSF2(95), U2AF1(34))
- MG: JAK2 stage diagnostics (V617F and Exon12) if PV is suspected
- MG: MPN panel (ASXL1, RUNX1, TP53)

MDS

- FISH: MDS panel
- MG: MDS panel (TET2, ASXL1, TP53, RUNX1)
- SNP: Specific mutation analysis – MDS panel (incl. CSF3R(615/618), IDH1(132), IDH2(140/172), NPM1(288fs), SF3B1(666/700), SRSF2(95), DNMT3A(525/714/735/736/749/860/882/904))
- MG: Myeloid panel (ASXL1, CBL, CSF3R, DDX41, DNMT3A, EZH2, IDH1, IDH2, JAK2, NRAS, RUNX1, SF3B1, SRSF2, STAG2, TET2, TP53, U2AF1, ZRSR2)

AML

- FISH: AML panel
- SNP: Specific mutation analysis – AML panel (incl. NPM1(288fs), FLT3(690/835), IDH1(132), IDH2(140/172), cKIT(541/816))
- MG: ASXL1, RUNX1, CEBPA, TP53
- MG: FLT3 (TKD and ITD)
- FISH / PCR: t(8;21) RUNX1/RUNX1T1
- FISH / PCR: t(15;17) PML/RARA
- FISH / PCR: inv(16) MYH11/CBFB

Other tests (declaration of consent is required)

- 5-FU toxicity screening (according to DGHO position paper)
- Hemochromatosis
- β-thalassemia
- Sickle cell anemia
- Meulengracht's disease
- Hereditary pancreatitis

Request for individual or special analyses: