

Area for stamps

**Referring provider** (please use **stamp**, alternatively blockletters)

Name/Clinic: .....

Department: .....

City/Country: .....

Phone: .....

**Invoice address:**

☐ Referring provider ☐ Patient/Other (declaration of cost coverage)

**Referring clinician**

Name: .....

Phone: .....

E-Mail: .....

**Required patient information** (please use **label**, alternatively blockletters)

Last name: .....

First name: .....

Gender: ☐ F ☐ M Insurance number/Date of birth: ..... / dd mm yyyy

Study: ..... Patient Study ID: .....

EUPID: .....

**Reason for referral / Indication**

- ☐ Suspected diagnosis ☐ Neuroblastoma ☐ Nephroblastoma ☐ Germ cell tumor  
☐ Initial diagnosis ☐ Ewing sarcoma ☐ Rhabdomyosarcoma ☐ Osteosarcoma  
☐ During treatment ☐ Relapse

**Remarks:**

**Sample**

Date sample taken: dd mm yyyy

Time sample taken: hh : mm

**Remarks:**

☐ **Tissue**

☐ Primary tumour ☐ Metastasis

- ☐ Native  
☐ Frozen  
☐ Touch imprints  
☐ Paraffin material (FFPE)  
Histology No. ....

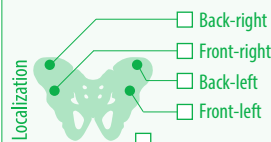
Localization / Remarks (e.g. when several tumour pieces are available):  
.....  
.....  
.....

Tumour cell content

%

☐ **Bone marrow**

- ☐ EDTA  
☐ Cytospin slides  
☐ PAXgene (to be requested)



☐ **Peripheral blood**

- ☐ EDTA  
☐ PAXgene (to be requested)  
☐ .....

☐ **Apheresis product\***

- ☐ EDTA  
☐ .....

\*Please also provide peripheral blood before and after apheresis

☐ **Body fluid**

- ☐ Ascites  
☐ CSF  
☐ Pleural effusion

☐ **Other samples**

☐ .....  
The primary sample collection manual can be found on <https://www.labdia.at>

**Moleculargenetic and immunocytologic testing**

☐ **Neuroblastoma**

- ☐ MYCN amplification (FISH)  
☐ ALK amplification / rearrangement (Break apart FISH)  
☐ ALK mutations (ddPCR, sequencing)  
☐ Segmental und numerical chromosomal aberrations (SCAs & NCAs) (SNP-Array)  
☐ .....

☐ Bone marrow infiltration / Minimal residual disease (MRD) (GD2/CD56 staining)

☐ **Rhabdomyosarcoma**

- ☐ FOXO1 (FKHR) rearrangement / fusion with:  
☐ PAX3 (FISH, qPCR)  
☐ PAX7 (FISH, qPCR)  
☐ Rare FOXO1 oder PAX3 rearrangements (Break apart FISH)  
☐ Genome-wide profile (CN/LOH) (SNP-Array)  
☐ .....

☐ Bone marrow infiltration / Minimal residual disease (MRD) (GD2/CD56 staining)

☐ **Ewing Sarcoma**

- ☐ EWSR1 rearrangement / fusion with:  
☐ FLI1 (FISH, qPCR)  
☐ ERG (FISH, qPCR)  
☐ Rare EWSR1 rearrangements (Break apart FISH)  
☐ .....

☐ Bone marrow infiltration / Minimal residual disease (MRD) (qPCR)  
information on the fusion incl. exons:  
.....

☐ **Nephroblastoma (Wilms-Tumor)**

- ☐ 1q gain (FISH)  
☐ Genome-wide profile (CN/LOH) (SNP-Array)  
☐ .....

☐ **Other tests**

- ☐ .....  
☐ .....  
☐ .....

**Asservation**

- ☐ asservation for now, feedback on desired testing follows  
☐ only for asservation in biobank / for a study:

Study: .....