



Please paste here the patient identification or provide the following data:

NAME _____

DATE OF BIRTH / / _____

Also paste the identification tag, or write the name on the tube(s)

REQUISITION OF GENETIC AND GENOMICS STUDIES OF SOLID TUMORS

I. REQUESTING ENTITY

REQUESTING PHYSICIAN _____

TELEPHONE _____

E-MAIL _____

II. SAMPLE SENT

1. SENT PRODUCT _____ COLLECTION DATE / / _____

SOLID TUMOR/METASTASIS IN PARAFFIN LYMPH NODE BIOPSY PERIPHERAL BLOOD

III. CLINICAL INFORMATION

1. DIAGNOSIS (DISCRIMINATE AS MUCH AS POSSIBLE) _____

2. THERAPY (CURRENT AND PRECEDING) _____

IV. INTENDED ANALYSIS

GENETIC RISK ASSESSMENT OR DIAGNOSIS OF FAMILIAL CANCER

(PANELS NGS + SANGER + MLPA, and evaluation of the *Alu* insert in *BRCA2*. The genes included in each panel are those recommended by the colleges of international specialties)

- 4001** Hereditary Breast and Ovary Cancer - *BRCA1* and *BRCA2* genes
 - 4000** Hereditary Breast and Ovary Cancer -12 gene panel (*BRCA1*, *BRCA2*, *PTEN*, *TP53*, *STK11*, *CDH1*, *ATM*, *PALB2*, *CHEK2*, *RAD51C*, *RAD51D*, *BRIP1*)
 - 4002** Hereditary Colon cancer - 13 gene panel (*EPCAM*, *ENG*, *BRAF*, *PTEN*, *BMPR14*, *SMAD4*, *STK11*, *PMS2*, *MSH6*, *MSH2*, *MLH2*, *MLH1*, *APC*, and *MUTYH*)
 - 4003** Lynch Syndrome- 5 gene panel (*MLH1*, *MSH2*, *MSH6*, *PMS2* and *EPCAM*)
 - 4004** Familial Adenomatous Polyposis - 2 gene panel *APC* and *MUTYH* genes
 - 4005** Hereditary Gastric Cancer - gene *CDH1*
 - 4006** Hereditary Melanoma - 5 gene panel (*CDKN2A*, *CDK4*, *BAP1*, *MITF*, *MC1R*)
 - 4007** Hereditary Pancreatic Cancer - 5 gene panel (*BRCA1*, *BRCA2*, *CDKN2A*, *PALB2*, *ATM*)
 - 4008** Hereditary Prostate Cancer - 3 gene panel *BRCA1*, *BRCA2*, *HOXB13* genes
 - 4009** Li Fraumeni Syndrome - gene *TP53*
 - 4010** Peutz Jeghers Syndrome - gene *STK11*
- other gene (s), which _____

FAMILY HISTORY

(Family tree with family history of cancer is indispensable for analysis)

TISSUE AND FLUID BIOPSY

The **tissue biopsy** (point out) is usually used in the diagnosis, the **liquid biopsy** (indicate) is typically used in patients in progression and / or treatment, or when it is not possible to access Tissue Biopsy

<input type="radio"/> BT 4410 NGS Mutational panorama- 50 genes (extended pharmacogenomics)	<input type="radio"/> BT 4458 qPCR <i>BRAF</i> V600E (response to vemurafenib and dabrafenib)
<input checked="" type="radio"/> BL 4413 test to identify therapeutic alternatives - multiple classes of drugs	<input checked="" type="radio"/> BL 4459
<input type="radio"/> BT 4415 panorama of mergers- 4 genes (<i>ALK</i> , <i>RET</i> , <i>ROS1</i> , <i>NTRK</i>)	<input type="radio"/> BT 4450 qPCR <i>EGFR</i> 29 mutations (response to EGFR inhibitors)
<input checked="" type="radio"/> BL 4416 predicting susceptibility tyrosine kinase inhibitors	<input checked="" type="radio"/> BL 4451
<input type="radio"/> BT 4424 NGS Panel <i>BRCAness</i> - 4 genes (<i>BRCA1</i> , <i>BRCA2</i> , <i>PALB2</i> and <i>ATM</i>)	<input type="radio"/> BT 4452 qPCR <i>EGFR</i> T790M (resistance to Erlotinib and other inhibitors)
<input checked="" type="radio"/> BL 4425 susceptibility to platinum-based drugs and <i>PARP</i> inhibitors	<input checked="" type="radio"/> BL 4453 of crosstalk tyrosine of the 1st and 2nd generation <i>EGFR</i>)
<input type="radio"/> BT 4422 NGS <i>TP53</i> (identification of pathogenic mutations for	<input type="radio"/> BT 4454 qPCR <i>KRAS</i> 19 mutations
<input checked="" type="radio"/> BL 4423 susceptibility to <i>TP53</i> activators)	<input checked="" type="radio"/> BL 4455 (resistance to EGFR inhibitors)
<input type="radio"/> BT 4420 NGS NGS <i>Her2</i> (identifies activating mutations for	<input type="radio"/> BT 4456 qPCR <i>NRAS</i> 16 mutations
<input checked="" type="radio"/> BL 4421 susceptibility to <i>Her2</i> receptor inhibitors)	<input checked="" type="radio"/> BL 4457 (resistance to EGFR inhibitors)
<input type="radio"/> BT 4418 NGS <i>ESR1</i> (early identification of resistance	<input type="radio"/> BT 4460 qPCR <i>PIK3CA</i> 5 mutations (response to mTOR inhibitors,
<input checked="" type="radio"/> BL 4419 mutations to hormone therapy)	<input checked="" type="radio"/> BL 4461 resistance to EGFR inhibitors)
<input type="radio"/> NGS Other(S) which _____	<input type="radio"/> qPCR Other(s), which _____

INFORMED CONSENT

I CONSENT to having the above genetic testing performed on which I have previously been clearly and objectively informed about the application and limitations thereof. I AUTHORIZE the collection of biological samples necessary for the performance of the genetic test (s) indicated by the Germano de Sousa Laboratory Medicine Center or, if necessary, by other laboratories designated by it. I AUTHORIZE that the data contained in this form be registered and treated only by duly authorized professionals, guaranteeing the protection and confidentiality according to the law in force. I GIVE MY CONSENT for the result (s) to be sent to the medical assistant. I WAS INFORMED about my right to revoke consent at any time without justification by sending an email.

SIGNATURE OF THE PATIENT OR THE LEGAL RESPONSIBLE (MINOR OR MAJOR INCAPABLE) OBLIGATORY _____

DATE _____

SIGNATURE OF THE DOCTOR OBLIGATORY _____

CONTACT FOR SUBMISSION OF RESULTS OBLIGATORY _____

IMPORTANT: Send the sample on the collection day. Do not send on Friday, not on the eve of holiday (consult central laboratory).