

REQUISITION FORM FOR GENETIC DISEASES AND PHARMACOGENETICS

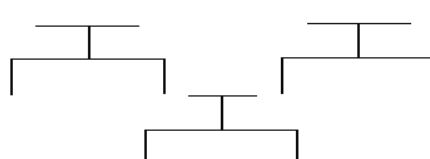
REFERRING DOCTOR (MANDATORY):	
Name (or print label):	ID number:
Hospital:	Service:
E-mail:	Telephone:
Signature:	Date: ____/____/____
Do you authorize the report to be sent by e-mail? Yes <input type="checkbox"/> No <input type="checkbox"/>	
If yes, please write your institutional e-mail address: _____	
PATIENT IDENTIFICATION (MANDATORY):	
Name:	Gender: F <input type="checkbox"/> M <input type="checkbox"/>
Identification number:	Date of birth: ____/____/____
<i>GenoMed use only:</i>	
Post label(s)	
<i>Conferred by:</i>	

CLINICAL DATA AND DIAGNOSIS*	
Affected <input type="checkbox"/>	Not affected (Asymptomatic) <input type="checkbox"/>
Ideal expected date: ____/____/____	
*Attach, whenever possible and can be justified, any relevant clinical information, family data including consanguinity, other cases in the family, family tree, etc.	

FAMILIAL INFORMATION	
Index case (affected) <input type="checkbox"/>	Familial variant* <input type="checkbox"/>
Spouse <input type="checkbox"/>	
Known familial variant? NO <input type="checkbox"/> YES <input type="checkbox"/>	
If yes*: Gene/RefSeq: _____ / _____ Variant: _____	
Was the index case studied at GenoMed? NO <input type="checkbox"/> YES <input type="checkbox"/>	
*For carrier studies and predictive testing (familial variant study), please attach a copy of the index case, whenever possible and applicable. According to the Portuguese law, (article 9º, law nº 12/2005), carrier studies and predictive testing (conducted in healthy individuals) it is ESSENTIAL that the clinical geneticist makes the request and that the patient informed consent is obtained.	

Information for Pedigree construction:

<input type="checkbox"/> Man	<input type="checkbox"/> Woman	<input checked="" type="checkbox"/> Affected	<input checked="" type="checkbox"/> Carrier	<input checked="" type="checkbox"/> Deceased	<input checked="" type="checkbox"/> Consanguinity	<input checked="" type="checkbox"/> Index case
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SAMPLE	COLLECTION
Blood (EDTA) <input type="checkbox"/> DNA <input type="checkbox"/> Tumor <input type="checkbox"/> ____% (infiltration percentage)	Date: ____/____/____
Other (specify) <input type="checkbox"/> _____	Hour: ____:____

INFORMED CONSENT (to be filled by the referring doctor)	
I hereby declare that the patient informed consent for diagnosis was obtained.	YES <input type="checkbox"/> NO <input type="checkbox"/>
I hereby declare that the patient informed consent for investigation was obtained.	YES <input type="checkbox"/> NO <input type="checkbox"/>

CONTACT PERSONS: Dr.ª Diana Antunes, MD (dianaantunes@medicina.ulisboa.pt) / Dr.ª Ana Coutinho, PhD (anacoutinho@medicina.ulisboa.pt) Ext. 47301/47308

GENETIC TEST REQUIRED*

Disease/Gene(s): _____

Study for familial variant (specify at "familial information")

NGS genetic panel (panel name): _____

Additional genes (if you want to personalize the above panel please specify which genes do you want to add or remove): _____

With CNVs analysis** (applicable to tests/panels with that option, marked at attachments I and II. For other CNVs studies, please contact us)

**Additional cost.

Additional NGS genes analysis (based on total exome) for:

NGS panel (panel name): _____

Gene(s) (specify): _____

Clinical Exome (single) (mandatory to fill in the "additional informed consent" – "information leaflet/Informed consent" – attachment III)

With CNVs analysis** (applicable to testes/panels with that option, marked at attachments I and II. For other CNVs studies, please contact us)

**Additional cost.

Clinical Exome (single) (mandatory to fill in the "additional informed consent" – "information leaflet/Informed consent" – attachment III)

With CNVs analysis**

Clinical exome (trio) (mandatory to fill in the "additional informed consent" – "information leaflet/Informed consent" – attachment III)

With CNVs analysis**

**Additional cost.

DNA extraction

Other studies*: _____

*See attachment I and/or II from our requisition or consult our website. **If the test you want isn't listed, please contact us.**

INFORMED CONSENT (mandatory – to be filled by the patient)

I hereby declare to authorize the collection of mine/my _____ [affiliation], _____ [name], born on ____/____/____, to the execution of the genetic testing described above, whose purposes and limitations were explained by the aforementioned physician. Herewith I declare that I have been informed about the consequences resulting from the teste results. I agree that the sample may be stored in order to allow repetition of the tests or further related tests at GenoMed or authorized partners around the world. The data/test results are subject to medical confidentiality and should only be disclosed to family members or other physicians with my permission. I am entitled to revoke this consent at any time.

Agree Not agree

I also declare that the data/test results may be used in scientific investigations and publications in an anonymized form when and only approved by the Ethics Committee.

Agree Not agree

(According to the Direction of General Health Standard nº 015/2013 updated.)

Patient's signature: _____ **Date:** ____ / ____ / ____