REQUISITION FORM FOR GENETIC DISEASES AND PHARMACOGENETICS



REFFERING 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 \square No \square	
If yes, please write your institutional e-mail address:	
PATIENT IDENTIFICATION (MANDATORY):	
Name:	Gender: F ☐ M ☐
Identification number:	Date of birth:/
GenoMed use only:	
Post label(s)	
Conferred by:	
CLINICAL DATA AND DIAGNOSIS*	
Affected ☐ Not affected (Asymptomatic) ☐	Ideal expected date:/
*Attach, whenever possible and can be justified, any relevant clinical information, family data including consa	nguinity other cases in the family family tree, etc.
FAMILIAL INFORMATION	
Index case (affected) ☐ Familial variant* ☐ Spouse ☐	
Known familial variant? NO YES	
If yes*: Gene/RefSeq: Variant:	
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Was the index case studied at GenoMed? NO ☐ YES ☐ *For carrier studies and predictive testing (familial variant study), please attach a copy of the index case, whe	never possible and applicable.
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 $\textbf{CONTACT PERSONS:} \ Dr.^a \ Diana \ Antunes, \ MD \ (dianaantunes@medicina.ulisboa.pt) \ / \ Dr. \ Yuri \ Chiodo, \ PhD \ (ychiodo@medicina.ulisboa.pt) \ Ext. \ 47301/48308 \ Antunes \ Antunes$

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REQUISITION FORM FOR GENETIC DISEASES AND PHARMACOGENETIC



GENETIC TEST REQUIRED*		
☐ Disease/Gene(s):		
☐ Study for familial variant (specify at "familial information")		
□ NGS genetic panel, with CNVs analysis (panel name):		
Additional genes (if you want to personalize the above panel please specify which genes do you want to add or remove):		
☐ Additional NGS genes analysis, with CNVs 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" – atta	achment III)	
\square Clinical Exome (single), with CNVs analysis (<u>mandatory</u> to fill in the "additional informed consent" – "information leaflet/Information leaf	med consent" – atta	chment III)
☐ Clinical exome (trio), with CNVs analysis (mandatory to fill in the "additional informed consent" – "information leaflet/Informed	d consent" — attachr	nent III)
Chilical exome (110), with City's analysis (manualory to limit the additional miorited consent – mioritation leaneth months	TCONSENT - attacnin	nent III)
□ 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 that I have read the Privacy Policy of GenoMed® - Diagnósticos de Medicina Molecular, S.A., available a	at https://genome	d.pt/en/privacy-
and-cookies-policy/ and I give my consent to the processing of the personal data.	☐Agree	☐ Not agree
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 ar	nd limitations were	e explained by
the aforementioned physician. Herewith I declare that I have been informed about the consequences resulting from the	ne 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	d. 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.		
(According to the Direction of General Health Standard no 015/2013 updated.)	☐ Agree	☐ Not agree
Patient's signature:	Date:/	

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