Order Form Array-CGH



General Information

Patient	
Surname:	
First name:	
Date of birth:	
Sex:	male female
Material	
Blood ml (min. 1-2 ml EDTA-blood)
Dried blood spo	t cards (at least 5 spots)
DNA µg (n	nin. 1-2 µg DNA, concentr. ≥ 50 ng/µl) DNA-No.:
Source material of extracted DNA:	(e.g. EDTA blood, skin biopsy)
Other specimer	·
External ID:	
Date of sample col	lection:
	mail in a cardboard box or air cushion envelope. Samples should not inthe pried blood snot cards can be ordered for free (info@cegat.com)

Declaration of consent

By signing this form, I declare that I have received comprehensive information regarding the genetic background related to the disease in question, as well as the possibilities and limitations of molecular genetic testing. I understand that I have the right to withdraw my consent for genetic analyses.

I have been informed, and agree, that my personal data and the data obtained in the analysis will be recorded, evaluated or stored in an pseudonymized form in scientific databases, and that further, in accordance with data protection and medical confidentiality, the request, or parts thereof, may be transmitted to a specialized cooperating laboratory.

I consent to the re-evaluation of my test results within the data storage period. If significant alterations become apparent, my Physician will be informed by e-mail.

I consent that in addition to the full genetic test as requested, the analysis can be expanded to all pathogenic and likely pathogenic variants (ACMG class 4 and 5) in genes which are related to the indication described for the proband (if applicable, screen for differential diagnosis).

I have been informed, and agree to the electronic storage, processing, use, and transmission of all data collected by CeGaT GmbH.

For more detailed information on data privacy as well as your rights please refer to www.cegat.com/privacy-policy/.

Please Note

Our panels are regularly updated to reflect current scientific research. It should therefore be recognized that there is the possibility that the list of genes on the order form may have changed slightly (genes added or removed) by the time the sample is analyzed in the laboratory. By signing this form, the patient accepts that the list of genes actually analyzed may be slightly different from what is currently listed. When NGS is utilized more than the requested genes are sequenced for each sample.

This declaration of consent can be completely or partially withdrawn at any time. I have had sufficient time to consider giving my consent.

I, the referring physician, confirm that I am authorized to request genetic testing for the above-mentioned patient. For predictive testing, I confirm that I am authorized, and that I have fulfilled the requirements, to request this testing. For minors, I declare that I have the consent of all legal guardians.

If the patient did not sign this order form: I, the referring physician, confirm that the patient received genetic counseling and agrees with the genetic testing. The patient's consent has been obtained in writing.

Sender / Clinic		
Surname:		_
First name:		_
Institution:		_
Street:		_
Postcode/City:		_
Country:		_
Phone:		_
Email:		_
VAT: If applicable, please includ	e a VAT number or a copy of your business registration certificate.	-
•••	 a VAT number or a copy of your business registration certificate. to sender / clinic to patient / other (KVA-No.:)
If applicable, please includ	La to sender / clinic	
If applicable, please includ	La to sender / clinic	_)
If applicable, please includ Invoice Surname:	La to sender / clinic	
If applicable, please includ Invoice Surname: First name:	La to sender / clinic)
If applicable, please includ Invoice Surname: First name: Street:	La to sender / clinic)

If you do not check these boxes, your answer will be recorded as "No".

With regard to secondary findings I would like to be informed:	Yes	🗆 No
I consent to the pseudonymous storage and use of surplus genetic material and/or test results for scientific research and in scientific literature.	Yes	🛛 No
I consent to the storage of my test results beyond the timespan of 10 years (as required by German law).	Yes	🛛 No
I consent to the storage of my genetic material for additional tests and/or quality control (for max. 10 years).	Yes	🛛 No

Genetic variation may sometimes be identified, which does not fit within the scope of the requested genetic analysis (so-called secondary findings). The reporting of these variants is limited to pathogenic alterations (ACMG classes 4 and 5) within selected genes, for which a treatment or course of action exists for you or your family (according to the current guidelines of the American College of Medical Genetics and Genomics; details on genes and associated diseases can be found at <u>www.cegat.com/acmg-genes/</u>). There is no claim of a comprehensive analysis of this gene set. An absence of secondary findings cannot be used to indicate a reduced disease risk.

Targeted analysis of the ACMG genes according to current recommendations can be requested as "additional analyses".

According to German Genetic Diagnostic Act (GenDG) we will issue the medical report to the counselling physician. Please indicate here the contact email of the counselling physician:

Email:

Physician's stamp / Barcode



CLIA CERTIFIED ID: 9902130225 CeGaT is accredited by DAKS according to DIN EN ISO 15189:2014, the College of American Pathologists (CAP) and CLIA.

Patient / Legal Guardian

(Block letters)

Patient / Legal Guardian (Date, Signature)



Physician

(Date, Signature)

(Surname, First name)

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Indication



Analysis type:	Proband is affected	□ Proband is NOT affected (predictive testing)	
Indication / Suspected diagnosis:			
Major Clinical Symptoms:			
Preliminary genetic diagnostics:			
Transplants (bone marrow, tissu	e, stem cells) 🛛 No	□ Yes, (please specify)	
Please include a copy of all exist	ing reports of your patie	nt.	
Pedigree	Consanguinity: 🛛 Yes	No Ethnic origin:	
			<i>index patient</i><i>index patient</i>
			○ □ not affected
			 not affected affected known carrier deceased
			 not affected affected known carrier deceased unrelated parents
			 not affected affected known carrier deceased
			 not affected affected known carrier known carrier consanguine parents
			 not affected affected known carrier deceased deceased ourrelated parents consanguine parents unborn child abortion, stillborn child person of unknown sex
			 not affected affected known carrier known carrier deceased unrelated parents consanguine parents unborn child abortion, stillborn child
			 not affected affected known carrier known carrier deceased ourrelated parents consanguine parents unborn child abortion, stillborn child person of unknown sex identical twins

For further information and advice please do not hesitate to contact our Diagnostic Support team. www.cegat.com/diagnostic-support · diagnostic-support@cegat.de · Phone +49707156544-55