

### General Information

<p><b>Patient</b></p> <p>Surname: _____</p> <p>First name: _____</p> <p>Date of birth: _____</p> <p>Sex: <input type="checkbox"/> male <input type="checkbox"/> female</p> <p><b>Material</b></p> <p><input type="checkbox"/> Blood ____ ml (min. 3 ml EDTA-blood)</p> <p><input type="checkbox"/> Dried blood spot cards (at least 10 spots)</p> <p><input type="checkbox"/> DNA ____ µg (min. 5 µg DNA, concentr. ≥ 50 ng/µl) DNA-No.: _____</p> <p><input type="checkbox"/> Other specimen _____</p> <p>External ID: _____</p> <p>Date of sample collection: _____</p> <p><small>Samples can be sent by mail in a cardboard box or air cushion envelope. Samples should not be exposed to direct sunlight. Dried blood spot cards can be ordered for free (info@cegat.com).</small></p>	<p><b>Sender / Clinic</b></p> <p>Surname: _____</p> <p>First name: _____</p> <p>Institution: _____</p> <p>Street: _____</p> <p>Postcode/City: _____</p> <p>Country: _____</p> <p>Phone: _____</p> <p>Email: _____</p> <p><small>If applicable, please include a VAT number or a copy of your business registration certificate.</small></p> <p>VAT: _____</p> <p><b>Invoice</b></p> <p><input type="checkbox"/> to patient <input type="checkbox"/> to sender / clinic</p>
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**Declaration of consent**

**By signing this form, I declare that I have received comprehensive information about 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 to genetic analyses.**

**I have been informed, and agree, that the data obtained in the analysis will be recorded, evaluated, or stored in an anonymized form in scientific databases, and further, in accordance with data protection and medical confidentiality, that the request, or parts thereof, may be transmitted to a specialized cooperating laboratory. I have been informed, and agree, that all data collected by CeGaT GmbH is electronically stored, processed, and used. I also consent to the data being transmitted electronically (e.g. by e-mail or fax).**

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

I consent to the storage of my genetic material for additional tests and/or quality control (for max. 10 years).  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 anonymous storage and use of surplus genetic material and/or test results for scientific research.  Yes  No

Genetic variation may sometimes be identified, which does not fit within the scope of the requested genetic analysis (so-called secondary findings report). The reporting of these variants is limited to pathogenic alterations 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; ACMG SF V2.0; Kalia et al., 2017, PMID: 27854360). There is no claim of a comprehensive analysis of the genes included within the secondary findings report. An absence of secondary findings cannot be used to indicate a reduced disease risk.

**With regard to secondary findings I would like:**

to be informed

to NOT be informed

**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.**

_____ Patient / Legal Guardian (Block letters)	_____ Doctor (Surname, First name)
X _____ Patient / Legal Guardian (Date, Signature)	X _____ Doctor (Date, Signature)


**Doctor's stamp / Barcode**

**Contact**

To discuss the diagnostic strategy please do not hesitate to contact us.  
 Phone: +49 7071 565 44-55  
 Email: diagnostic-support@cegat.de



Deutsche  
Akkreditierungsstelle  
D-MU-13206-01-00



ACCREDITED  
COLLEGE of AMERICAN PATHOLOGISTS  
CLIA CERTIFIED ID: 9902130225

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

Indication

**Indication / Suspected Diagnosis:**

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**Clinical Major Symptoms:**

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**Preliminary genetic diagnostics:**

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**Transplants (bone marrow, tissue, stem cells)**  No  Yes, (please specify) \_\_\_\_\_

**Please include a copy of all existing reports of your patient.**

**Pedigree**                      Consanguinity:  Yes  No                      Ethnic origin: \_\_\_\_\_

- index patient
- not affected
- affected
- known carrier
- deceased
- unrelated parents
- consanguine parents
- unborn child
- abortion, stillborn child
- person of unknown sex
- identical twins (monozygous)
- fraternal twins (dizygous)

**Family medical history**

Are there other family members who currently have or have had the same or a similar disease as the patient?

Yes                       No

If yes, please list the affected family members:

Name (not required)	Relationship to the patient (e.g. mother)	Age of onset	Diagnosis / Symptoms

## Inquiry

## Inquiry

 **Large Panel Diagnostic Option (124 Genes, CAN-D45)**

In addition to the full analysis of the requested gene set(s), we will expand the analysis to all pathogenic and likely pathogenic variants (ACMG class 4 and 5) in genes which are related to the indication described for the proband on the entire diagnostic panel (124 genes in total).

 **Colorectal Cancer (26 Genes, CAN01)**

APC, AXIN2, BMPR1A, CDH1, CHEK2, EPCAM, FLCN, GREM1, MLH1, MSH2, MSH3, MSH6, MUTYH, NBN, NTHL1, PMS2, POLD1, POLE, PTEN, RINT1, RNF43, RPS20, SCG5, SMAD4, STK11, TP53

 **Colorectal Cancer – Polyposis syndrome (14 Genes, CAN11)**

APC, BMPR1A, CHEK2, GREM1, MSH3, MUTYH, NTHL1, POLD1, POLE, PTEN, RNF43, SCG5, SMAD4, STK11

 **Colorectal Cancer – Hereditary Nonpolyposis Colorectal Cancer (HNPCC)/Lynch Syndrome (5 Genes, CAN12)**

EPCAM, MLH1, MSH2, MSH6, PMS2

 **Gastric Cancer (10 Genes, CAN13)**

BRCA2, CDH1, EPCAM, IL1B, IL1RN, KIT, MLH1, MSH2, MSH6, PMS2

 **Cowden Syndrome (6 Genes, CAN14)**

AKT1, PIK3CA, PTEN, SDHB, SDHD, SEC23B

 **Pancreatic Cancer (17 Genes, CAN06)**

APC, ATM, BRCA1, BRCA2, CDKN2A, EPCAM, MLH1, MSH2, MSH6, PALB2, PALLD, PMS2, PRSS1, SPINK1, STK11, TP53, VHL

 **Breast and Ovarian Cancer (12 Genes, CAN02)**

ATM, BRCA1, BRCA2, CDH1, CHEK2, NBN, PALB2, PTEN, RAD51C, RAD51D, STK11, TP53

 **Breast and Ovarian Cancer – extended (40 Genes, CAN21)**

ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, EPCAM, FAM175A, FANCA, FANCC, FANCD2, FANCE, FANCF, FANCG, HOXB13, MEN1, MLH1, MRE11A, MSH2, MSH3, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, PTCH1, PTEN, RAD50, RAD51C, RAD51D, RINT1, SDHB, SDHC, SDHD, SLX4, STK11, TP53, XRCC2

 **Prostate Cancer (11 Genes, CAN03)**

BRCA1, BRCA2, CHEK2, EPCAM, HOXB13, MLH1, MSH2, MSH6, NBN, PMS2, TP53

 **Pheochromocytoma and Paraganglioma (14 Genes, CAN04)**

CDKN1B, MAX, MEN1, NF1, PRKAR1A, RET, SDHA, SDHAF2, SDHB, SDHC, SDHD, SRGAP1, TMEM127, VHL

 **Other Familial Tumor Syndromes (53 Genes, CAN05)**

AIP, AKT1, ALK, APC, ATR, BAP1, BLM, BRCA2, CDC73, CDH1, CDKN1C, CDKN2A, CYLD, DICER1, EPCAM, FH, HRAS, IL1B, IL1RN, KIT, LIG4, LZTR1, MET, MLH1, MSH2, MSH6, NBN, NF1, NF2, PIK3CA, PMS2, PTEN, RASAL1, RB1, RECQL4, RET, RHBDF2, SDHA, SDHB, SDHC, SDHD, SEC23B, SMARCA4, SMARCB1, SMARCE1, SPRED1, SUFU, TP53, TSC1, TSC2, VHL, WRN, YAP1

 **Tumors of the Central Nervous System (18 Genes, CAN51)**

AIP, APC, BRCA2, MLH1, MSH2, MSH6, NF1, NF2, PMS2, PTEN, SDHA, SDHB, SDHD, SMARCE1, SPRED1, SUFU, TP53, VHL

 **Renal cell carcinoma (28 Genes, CAN07)**

BAP1, CCND1, CDC73, CHEK2, DIS3L2, EPCAM, FH, FLCN, HNF1A, HNF1B, MET, MITF, MLH1, MSH2, MSH6, PALB2, PMS2, PTEN, SDHA, SDHAF2, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL, WT1

 **Xeroderma Pigmentosum (9 Genes, CAN08)**

DDB2, ERCC1, ERCC2, ERCC3, ERCC4, ERCC5, POLH, XPA, XPC

 **Melanoma (15 Genes, CAN09)**

BAP1, BRCA2, CDK4, CDKN2A, EPCAM, MC1R, MITF, MLH1, MSH2, MSH6, PMS2, POT1, PTEN, RB1, TP53

 **Fanconi-Anemia (21 Genes, CAN10)**

BRCA1, BRCA2, BRIP1, ERCC4, FANCA, FANCB, FANCC, FANCD2, FANCE, FANCF, FANCG, FANCI, FANCL, FANCM, MAD2L2, PALB2, RAD51, RAD51C, SLX4, UBE2T, XRCC2

## Additional analyses

For further information and advice please do not hesitate to contact our Diagnostic Support team.

[www.cegat.de/en/diagnostic-support](http://www.cegat.de/en/diagnostic-support) · [diagnostic-support@cegat.de](mailto:diagnostic-support@cegat.de) · Phone +49 7071 565 44-55