Order Form
Epilepsy & Brain Development Disorders

General Information

Patient

Surname: _____

First name: _____

Date of birth: _____

Sex: ❑ male ❑ female

Material

❑ Blood ____ ml (min. 1-2 ml EDTA-blood)

❑ Dried blood spot cards (at least 5 spots)

❑ DNA ____ µg (min. 1-2 µg DNA, concentr. ≥ 50 ng/µl) DNA-No.: _____

Source material
of extracted DNA: _____ (e.g. EDTA blood, skin biopsy)

❑ Other specimen _____

External ID: _____

Date of sample collection: _____

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).

Sender / Clinic

Surname: _____

First name: _____

Institution: _____

Street: _____

Postcode/City: _____

Country: _____

Phone: _____

Email: _____

VAT: _____

If applicable, please include a VAT number or a copy of your business registration certificate.

Invoice

❑ to sender / clinic
❑ to patient / other (KVA-No.: _____)

Surname: _____

First name: _____

Street: _____

Postcode/City: _____

Country: _____

Email: _____

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.

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 pseudonymous storage and use of surplus genetic material and/or test results for scientific research and in scientific literature. ❑ Yes ❑ No

With regard to secondary findings I would like to be informed: ❑ 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 www.cegat.com/acmg-genes/). 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

_____ _____

Patient / Legal Guardian (Block letters) | **Physician** (Surname, First name)

X _____ X _____

Patient / Legal Guardian (Date, Signature) | **Physician** (Date, Signature)

DAkkS Deutsche Akkreditierungsstelle D-ML-13206-01-00

CAP ACCREDITED COLLEGE of AMERICAN PATHOLOGISTS

CLIA CERTIFIED ID: 99D2130225

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Order Form
Epilepsy & Brain Development Disorders

Indication

Analysis type: ❑ Proband **is affected** ❑ Proband is **NOT affected** (predictive testing)

Indication / Suspected diagnosis:

Major Clinical Symptoms:

Preliminary genetic diagnostics:

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:

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

Order Form
Epilepsy & Brain Development Disorders

Medical History

Clinical features

1. Epileptic seizures

❑ No ❑ Yes; onset? _____

Etiology/Seizure types: _____

Sleep-related ❑ No ❑ Yes: _____

EEG ❑ Not performed

Further information: _____

2. Psychomotor development

❑ Progression ❑ Stagnation ❑ Regression

Intellectual disability	❑ No	❑ Yes
Speech / Language impairment	❑ No	❑ Yes
Motor deficits	❑ No	❑ Yes
Abnormal muscle tone	❑ No	❑ Yes; type? _____
Acute encephalopathy	❑ No	❑ Yes
Cerebellar dysfunction	❑ No	❑ Yes; onset? _____
Extrapyramidal dysfunction	❑ No	❑ Yes; onset? _____
Dementia	❑ No	❑ Yes; onset? _____

Remarks: _____

3. Clinical findings

Dysmorphic features	❑ No	❑ Yes; details: _____
Skin abnormalities	❑ No	❑ Yes; details: _____
Impaired vision	❑ No	❑ Yes; onset? _____

Other anomalies: _____

4. Head circumference

❑ Normal ❑ Microcephalic ❑ Macrocephalic Percentile: _____

5. MRI

❑ Not performed

Remarks: _____

6. Pregnancy history

Abnormal	❑ No	❑ Yes (please answer following questions)
Bleedings	❑ No	❑ Yes
Infection	❑ No	❑ Yes; details: _____
Medication	❑ No	❑ Yes; details: _____
Preterm birth	❑ No	❑ Yes; gestation week? _____
Hypoxia	❑ No	❑ Yes; pH umbilical cord? _____

Other noticeable occurrences: _____

7. Birth data

Size: _____ Weight: _____ Head circumference: _____

Noticeable problems: _____

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Medical History & Inquiry

8. Genetic analyses ❑ Not performed ❑ Yes **(please attach copy of results if patient agrees)**

Array CGH: ❑ No ❑ Yes

Sequencing: ❑ No ❑ Yes

Other: _____

9. Metabolic tests ❑ Not performed ❑ Yes **(please attach copy of results if patient agrees)**

Abnormalities: _____

10. Further information _____

Inquiry Array-CGH

❑ Please perform Array-CGH before Panel Diagnostics

❑ Array-CGH analysis has already been performed

❑ Array-CGH analysis not required

Inquiry – Epilepsy

All genes listed on this order form, and the entire mitochondrial genome, are sequenced in parallel. This allows you to select multiple gene sets, or choose individual genes in addition to your selected gene set. As there is no additional laboratory expenditure, there is only a moderate price increase due to the subsequent analysis and interpretation. We are happy to answer your questions or send you an individual quote. Please contact us at info@cegat.com.

❑ **Familial and Idiopathic Epilepsy (31 Genes, EPI01)**
CACNA1A, CHRNA2, CHRNA4, CHRNB2, DEPDC5, GABRA1, GABRB3, GABRG2, GRIN2A, HCN1, KCNA1, KCNMA1, KCNQ2, KCNQ3, KCNT1, LGI1, MTOR, NPRL2, NPRL3, PCDH19, PRRT2, RELN, RORB, SCN1A, SCN1B, SCN2A, SCN3A, SCN8A, SLC2A1, STX1B, TBC1D24

❑ **Epileptic Encephalopathy (151 Genes, EPI02)**
AARS1, ABAT, ACTL6B, ADAM22, ALDH7A1, ALG13, AMT, AP2M1, AP3B2, ARHGEF9, ARV1, ARX, ATP6V1A, BRAT1, CACNA1A, CACNA1B, CACNA1E, CAD, CAMK2A, CAMK2B, CDK19, CDKL5, CHD2, CLCN4, CNPY3, CPLX1, CUL3, CUX2, CYFIP2, DALRD3, DDX3X, DENND5A, DMXL2, DNM1, DOCK7, EEF1A2, EIF3F, FGF12, FGF13, FOXG1, FRRS1L, GABBR2, GABRA1, GABRA2, GABRA5, GABRB1, GABRB2, GABRB3, GABRG2, GAD1, GAMT, GLDC, GLS, GNAO1, GNB1, GOT2, GRIA4, GRIN1, GRIN2A, GRIN2B, GRIN2D, GRM7, GUF1, HCN1, HNRNPU, IQSEC2, ITPA, KCNA2, KCNB1, KCNQ2, KCNQ5, KCNT1, KCNT2, LNPK, MBD5, MBOAT7, MDH1, MDH2, MECP2, MEF2C, MOCS1, MOCS2, NARS1, NBEA, NCDN, NECAP1, NEUROD2, NEXMIF, NTRK2, NUS1, PACS2, PARS2, PCDH19, PHACTR1, PIGA, PIGB, PIGP, PIGQ, PIGS, PLCB1, PLPBP, PNKP, PNPO, POLG, PPP2CA, PPP2R1A, PPP2R5D, PPP3CA, PTPN23, PURA, RHOBTB2, RNF13, ROGDI, RORA, SCN1A, SCN1B, SCN2A, SCN3A, SCN8A, SIK1, SLC12A5, SLC13A5, SLC1A2, SLC25A12, SLC25A22, SLC2A1, SLC35A2, SLC6A1, SLC6A8, SLC9A6, SMC1A, SPTAN1, ST3GAL3, STXBP1, SYNGAP1, SYNJ1, SZT2, TANC2, TBC1D24, TBL1XR1, TCF4, TRAK1, TSC1, TSC2, UBA5, UBE3A, UGDH, UGP2, WWOX, YWHAG, ZEB2

❑ **Progressive Myoclonus Epilepsy (17 Genes, EPI05)**
AFG3L2, ASAH1, CERS1, CSTB, EPM2A, GOSR2, KCNC1, KCTD7, LMNB2, NEU1, NHLRC1, PRDM8, PRICKLE1, SCARB2, SEMA6B, SERPINI1, SLC7A6OS

❑ ***CSTB* repeat analysis not required** (associated with Unverricht-Lundborg disease, a slowly progressive myoclonus epilepsy)
The most common pathogenic alteration in CSTB is a dodecamer repeat expansion in the promotor region which cannot be detected by NGS.

❑ **Neuronal Ceroid Lipofuscinosis (13 Genes, EPI06)**
ATP13A2, CLN3, CLN5, CLN6, CLN8, CTSD, CTSF, DNAJC5, GRN, KCTD7, MFSD8, PPT1, TPP1

❑ **GPI Anchor Deficiency with or without Hyperphatasia (24 Genes, EPI12)**
ARV1, GPAA1, PGAP1, PGAP2, PGAP3, PIGA, PIGB, PIGC, PIGF, PIGG, PIGH, PIGK, PIGL, PIGM, PIGN, PIGO, PIGP, PIGQ, PIGS, PIGT, PIGU, PIGV, PIGW, PIGY

❑ **Migraine (8 Genes, EPI14)**
ATP1A2, ATP1A3, CACNA1A, NOTCH3, PRRT2, SCN1A, SLC1A3, SLC2A1

❑ **Hyperekplexia (4 Genes, EPI15)**
ATAD1, GLRA1, GLRB, SLC6A5

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Inquiry

Inquiry – Brain Development Disorders

❑ **Primary Microcephaly and Differential Diagnoses (68 Genes, BRN01)**
ANKLE2, ASNS, ASPM, ATR, CDC45, CDC6, CDK5RAP2, CDK6, CDT1, CENPE, CENPF, CENPJ, CEP135, CEP152, CEP63, CIT, COPB2, CTNNA2, DNA2, DONSON, DYRK1A, GMNN, KIF11, KIF14, KIF2A, KIF5C, KNL1, LMNB1, LMNB2, MCM5, MCPH1, MFSD2A, NBN, NCAPD2, NCAPD3, NCAPH, NIN, NSMCE2, NUP37, ORC1, ORC4, ORC6, PCNT, PHC1, PHGDH, PLK4, PNKP, PSAT1, PSPH, RBBP8, RNU4ATAC, RRP7A, RTTN, SASS6, STIL, TRAIP, TRAPPC14, TUBA8, TUBB, TUBB2A, TUBB2B, TUBB3, TUBG1, TUBGCP4, TUBGCP6, WDFY3, WDR62, ZNF335

❑ **Neuronal Migration Disorders (76 Genes, BRN02)**
ACTB, ACTG1, ADGRG1, AKT3, APC2, ARF1, ARFGEF2, ARX, B3GALNT2, B4GAT1, CCND2, CDK5, CEP85L, COL3A1, COL4A1, COL4A2, CRADD, CRPPA, CTNNA2, DAG1, DCX, DYNC1H1, ERMARD, FAT4, FKRP, FKTN, FLNA, GMPPB, GRIN1, GRIN2B, KATNB1, KIF2A, KIF5C, KIFBP, LAMB1, LAMC3, LARGE1, MACF1, MAP1B, MAST1, NDE1, NEDD4L, OCLN, PAFAH1B1, PI4KA, PIK3CA, PIK3R2, POMGNT1, POMGNT2, POMK, POMT1, POMT2, PRUNE1, RAB18, RAB3GAP1, RAB3GAP2, RAC3, RELN, RTTN, RXYLT1, SNAP29, STAT2, TBC1D20, TMTC3, TSC1, TSC2, TUBA1A, TUBB, TUBB2A, TUBB2B, TUBB3, TUBG1, TUBGCP2, VLDLR, WDR62, WDR81

❑ **Holoprosencephaly Spectrum (15 Genes, BRN03)**
CDON, CNOT1, DHCR7, DISP1, DLL1, FGF8, FGFR1, GLI2, PTCH1, SHH, SIX3, STAG2, TDGF1, TGIF1, ZIC2

❑ **Pontocerebellar Hypoplasia (24 Genes, BRN14)**
AMPD2, CASK, CDC40, CHMP1A, CLP1, COASY, EXOSC1, EXOSC3, EXOSC8, EXOSC9, PCLO, PPIL1, RARS2, SEPSECS, SLC25A46, TBC1D23, TOE1, TSEN15, TSEN2, TSEN34, TSEN54, VPS51, VPS53, VRK1

❑ **Joubert Syndrome (43 Genes, BRN07)**
AHI1, ARL13B, ARL3, ARMC9, B9D1, B9D2, C2CD3, CC2D2A, CEP104, CEP120, CEP164, CEP290, CEP41, CPLANE1, CSPP1, FAM149B1, HYLS1, IFT172, INPP5E, KATNIP, KIAA0586, KIAA0753, KIF7, MKS1, NPHP1, OFD1, PDE6D, PIBF1, POC1B, RPGRIP1L, SUFU, TCTN1, TCTN2, TCTN3, TMEM107, TMEM138, TMEM216, TMEM231, TMEM237, TMEM67, TOGARAM1, TTC21B, ZNF423

❑ **Leukodystrophy (86 Genes, BRN05)**
AARS1, AARS2, ABCD1, ACBD5, ACOX1, ADAR, AIMP1, AIMP2, ALDH3A2, ARSA, ASPA, BCAP31, CLCN2, CLDN11, CNP, CSF1R, CTC1, CYP27A1, DARS1, DARS2, DEGS1, EARS2, EIF2AK2, EIF2B1, EIF2B2, EIF2B3, EIF2B4, EIF2B5, EPRS1, FAM126A, GALC, GAN, GBE1, GFAP, GJC2, HEPACAM, HIKESHI, HSD17B4, HSPD1, HTRA1, IFIH1, KARS1, L2HGDH, LMNB1, LSM11, MLC1, NAXD, NAXE, NKX6-2, NOTCH3, OCLN, PLAA, PLEKHG2, PLP1, POLR1C, POLR3A, POLR3B, POLR3K, PSAP, PYCR2, RAB11B, RARS1, RNASEH2A, RNASEH2B, RNASEH2C, RNASET2, RNU7-1, SAMHD1, SCP2, SLC16A2, SLC17A5, SLC25A12, SNORD118, SOX10, STAT2, STN1, SUMF1, TMEM106B, TMEM63A, TREM2, TREX1, TUBB4A, TYROBP, UFM1, VPS11, ZNHIT3

❑ **Aicardi-Goutières Syndrome (10 Gene, BRN06)**
ADAR, IFIH1, LSM11, RNASEH2A, RNASEH2B, RNASEH2C, SAMHD1, RNU7-1, STAT2, TREX1

❑ **Macrocephaly (62 Genes, BRN04)**
AKT1, AKT2, AKT3, APC2, ASPA, ASXL2, BRWD3, CCND2, CDKN1C, CHD3, CHD4, CHD8, CRADD, DIS3L2, DNMT3A, EED, EZH2, GCDH, GFAP, GLI3, GPC3, H1-4, HEPACAM, HERC1, HRAS, HUWE1, KIF7, KPTN, L1CAM, MITF, MLC1, MPDZ, MTOR, NF1, NFIA, NFIB, NFIX, NONO, NSD1, PAK1, PHF6, PIGA, PIK3CA, PIK3R2, PPP1CB, PPP2R5B, PPP2R5C, PPP2R5D, PTCH1, PTEN, RAB39B, RIN2, RNF125, SETD2, SOS1, STRADA, SUFU, SUZ12, TBC1D7, TRIO, UPF3B, ZBTB20

❑ **Kabuki Syndrome (3 Genes, BRN11)**
CHD7, KDM6A, KMT2D

❑ **Coffin-Siris Syndrome (13 Genes, BRN12)**
ARID1A, ARID1B, ARID2, BICRA, DPF2, SMARCA2, SMARCA4, SMARCB1, SMARCC2, SMARCD1, SMARCE1, SOX11, SOX4

❑ **Cornelia de Lange Syndrome (8 Genes, BRN13)**
ANKRD11, BRD4, HDAC8, NIPBL, RAD21, SMC1A, SMC3, UBE2A

Additional analyses (additional fees may apply)

❑ **HLA-Typing (HLA01)**
I would like to receive an additional report stating the HLA alleles (HLA class I (Gene A, B, C) and HLA class II (Gene DPA1, DPB1, DQA1, DQB1, DRB1, DRB3, DRB4, DRB5)).

❑ **ACMG genes diagnostics**
I would like to be informed of relevant alterations within the list of recommended genes for secondary analysis, according to the current guidelines of the American College of Medical Genetics and Genomics. The analysis is restricted to the sequence data, re-sequencing of regions with poor sequence coverage will not typically be performed. A negative “ACMG genes” report cannot be used to rule out (genetic) disease risk. Additional fees may apply. According to German legislation, predictive tests for minors may not be performed for diseases which have an onset in adulthood. Therefore, some genes will not be analyzed for minors, unless the phenotypic spectrum is within the scope of the primary medical indication of the patient. Details on genes and associated diseases can be found at https://www.cegat.com/acmg-genes/

❑ **Pharmacogenetics (PGX) (22 genes)**
ABCG2, CACNA1S, CYP2B6, CYP2C19, CYP2C9, CYP2D6, CYP3A4, CYP3A5, CYP4F2, DPYD, G6PD, HLA-A, HLA-B, IFNL3, MT-RNR1, NUDT15, POR, RYR1, SLCO1B1, TPMT, UGT1A1, VKORC1

I would like to receive an additional report analyzing known variants in 22 genes that are involved in the metabolism of pharmaceutical products.

❑ _____

For further information and advice please do not hesitate to contact our Diagnostic Support team at any time.
www.cegat.com/diagnostic-support | diagnostic-support@cegat.com | Phone +49 7071 565 44-55