



Biochemical and Biomarker Testing

> Patient information

Surname _____

First Name _____

Date of Birth MM DD YYYY

Sex Male Female

Street _____

Zip Code / Town _____

Country _____

Your Reference Number _____

Sample Collection Date MM DD YYYY

> Physician or laboratory (Reporting Address)

Name of Physician _____

Clinic _____

Department _____

Street _____

Zip Code / Town _____

Country _____

Telephone _____ Fax _____

E-Mail _____

> Billing

CENTOGENE Quotation No. _____

Invoice to Patient Clinic/Insurance
Please attach Authorization/Referral

Name _____

Department _____

Street _____

ZIP Code/Town _____

Country _____

Phone _____ Fax _____

E-Mail _____

I confirm that I have the patient's signature on file for all of the issues mentioned above and that I am aware that the patient can request us to have his/her results eliminated at any time and that I shall convey this request to CENTOGENE.

> Biomarker Testing only

Gaucher disease Fabry disease

Faber disease Niemann-Pick disease

For patients under enzyme replacement therapy (ERT) please specify the ERT:

Name of enzyme replacements: _____

Start of ERT: MM DD YYYY

Dosage: _____

> Additional report recipient

Name of Physician _____

Clinic _____

Department _____

Street _____

Zip Code / Town _____

Country _____

Telephone _____ Fax _____

E-Mail _____

> In Case of Direct Billing to the Patient

I authorize the physician to request this analysis/these analyses and I am informed about the resulting costs (and possibly applicable German 19% VAT). I herewith undertake to be liable for the payment of any invoice related to this diagnostics and I declare that the address given above is the correct billing address.

Place, Date _____

Signature of Patient/Guardian ~~X~~ _____

Minimum Sample Requirements

- Purified DNA (2 µg with min. 50ng/µl)
- EDTA Blood (1 ml)
- Centocard® (1pc, 10 fully saturated dried blood spots)

Place, Date _____

Signature of Physician ~~X~~ _____

CENTOGENE AG

Am Strande 7
18055 Rostock, Germany

Contact Details

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www.centogene.com

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> Biochemical genetics panel

Please select panel to test

Enzyme Panel = analysis of all enzyme activities within the panel

Enzyme Panel X-TRA = analysis of all enzyme activities within the panel

+ sequencing/CNVs of implicated gene(s) in event of pathological findings in any enzyme activity

✓	PANEL	12 ENZYMES	
	CentoSphingo Enzyme Panel	Acidic sphingomyelinase Beta-glucocerebrosidase Chitotriosidase	Hexosaminidase AB Alpha-N-acetylgalactosaminidase Acid lipase
	CentoSphingo Enzyme Panel X-TRA	Alpha-galactosidase Acidic alpha-glucosidase Beta-hexosaminidase	Alpha-mannosidase Beta-mannosidase Alpha-fucosidase

✓	PANEL	2 ENZYMES	
	CentonCL Enzyme Panel	Palmitoyl-protein thioesterase	
	CentonCL Enzyme Panel X-TRA	Tripeptidyl peptidase	

✓	PANEL	8 ENZYMES	
	CentomPS Enzyme Panel	Alpha-L-iduronidase Iduronate-2-sulfatase	Beta-galactosidase Arylsulfatase B
	CentomPS Enzyme Panel X-TRA	N-acetyl-alpha-glucosaminidase N-acetylgalatosamine-6-sulfate-sulfatase	Beta-glucuronidase Alpha-mannosidase

✓	PANEL	21 ENZYMES	
	CentomSD Enzyme Panel	Acidic sphingomyelinase Beta-glucocerebrosidase Chitotriosidase Alpha-galactosidase Acidic alpha-glucosidase Beta-hexosaminidase	Alpha-fucosidase Alpha-L-iduronidase Iduronate-2-sulfatase N-acetyl-alpha-glucosaminidase N-acetylgalatosamine-6-sulfate-sulfatase
	CentomSD Enzyme Panel X-TRA	Hexosaminidase AB Alpha-N-acetylgalactosaminidase Acid lipase Alpha-mannosidase Beta-mannosidase	Beta-galactosidase Arylsulfatase B Beta-glucuronidase Palmitoyl-protein thioesterase Tripeptidyl peptidase

> Biomarker Testing

Please select biomarker to test

✓	BIOMARKER	DISEASE
	C26 Ceramide (cis-C26 Cer)	Farber disease
	Glucosylsphingosine (lyso-Gb1)	Gaucher disease
	Lyso-Ceramide trihexoside (lyso-Gb3)	Fabry disease
	Lyso-SM509	Niemann-Pick disease type A/B/C



> Biochemical Testing

Please select the enzymatic activity(ies) to test according to left column

✓	ENZYME	DISEASE	GENE
	Acid lipase	Wolman disease, Cholesteryl ester storage diseases	LIPA
	Acidic alpha-glucosidase	Pompe disease	GAA
	Acidic sphingomyelinase	Niemann-Pick Type A/B disease	SMPD1
	Alpha-fucosidase	Alpha-fucosidase deficiency	FUCA1
	Alpha-galactosidase	Fabry disease	GLA
	Alpha-L-iduronidase	Hurler disease, MPS I	IDUA
	Alpha-mannosidase	Alpha-mannosidase deficiency	MAN2B1
	Alpha-N-acetylgalactosaminidase	Schindler/Kanzaki disease	NAGA
	Arylsulfatase A	Metachromatic leukodystrophy (MLD)	ARSA
	Arylsulfatase B	Maroteaux-Lamy syndrome, MPS VI	ARSB
	Beta-galactosidase	Morquio disease, MPS IVB	GLB1
	Beta-glucocerebrosidase	Gaucher disease	GBA
	Beta-glucuronidase	Sly syndrome, MPS VII	GUSB
	Beta-hexosaminidase	Tay-Sachs disease	HEXA
	Beta-mannosidase	Beta-mannosidase deficiency	MANBA
	Chitotriosidase	Gaucher disease (unspecific)	-
	Galactocerebrosidase	Krabbe disease	GALC
	Hexosaminidase AB	Sandhoff disease	HEXB
	Iduronate-2-sulfatase	Hunter disease, MPS II	IDS
	N-acetylgalactosamine-6-sulfate-sulfatase	Morquio disease, MPS IVA	GALNS
	N-acetyl-alpha-glucosaminidase	Sanfilippo syndrome, MPS IIIB	NAGLU
	Palmitoyl-protein thioesterase	Batten disease type 1, Neuronal ceroid lipofuscinosis type1, NCL1, Infantile NCL, Santavuori-Haltia disease	PPT1
	Tripeptidyl peptidase	Batten disease type 2, Neuronal ceroid lipofuscinosis type2, NCL2, Late infantile NCL, Jansky-Bielschowsky disease	TPP1
	Combined N-acetylgalactosamine-6-sulfate-sulfatase and beta-galactosidase	Morquio disease IVA, MPS IVA and Morquio disease IVB, MPS IVB	GALNS, GLB1

> Patient Clinical Information/Familial Mutations Information (Pedigree)

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Please use a separate sheet of paper if necessary

Minimum sample requirements: General: ≥ 2 ml EDTA blood or ≥ 1 Filtercard (CentoCard®)
For arylsulfatase A and beta-glucosidase from leucocytes:
≥ 5ml EDTA blood