

The Essential
Life Science Partner
for Data-Driven Answers
in Rare & Neurodegenerative
Diseases

CENTOGENE (CNTG) Company Presentation June 2023



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Certain information contained in this presentation relates to or is based on studies, publications, surveys and other data obtained from third-party sources and the Company's own internal estimates and research. While the Company believes these third-party sources to be reliable as of the date of this presentation, it has not independently verified, and makes no representation as to the adequacy, fairness, accuracy or completeness of, any information obtained from third-party sources. In addition, all of the market data included in this presentation involves a number of assumptions and limitations, and there can be no guarantee as to the accuracy or reliability of such assumptions. Finally, while the Company believes its own internal research is reliable, such research has not been verified by any independent source.

For further information, please refer to the Risk Factors section in our Annual Report for the year ended December 31, 2022, on Form 20-F filed with the SEC on May 16, 2023, and other current reports and documents filed with the U.S. Securities and Exchange Commission (SEC). You may get these documents by visiting EDGAR on the SEC website at www.sec.gov.

CENTOGENE @ a Glance



The essential life science partner for data-driven answers in rare and neurodegenerative diseases

We are on a mission to provide data-driven, **life-changing answers** to patients, physicians, and pharma companies



2006

CENTOGENE was founded



Global

Offices in Germany, U.S., Switzerland, and Serbia



IPO

In 2019 (CNTG)



47.5m

in FY2022 revenues - Guidance FY2023: YoY growth 10-15%²



>750k

Individuals in CNTG Biodatabank, the world's largest real-world integrated multiomic data repository in rare and neurodegenerative diseases



50+

Collaborations with biotech/biopharma partners, covering over 46 rare diseases¹





>90% of the >350 Million people affected by a rare disease are have not yet been diagnosed

>90% of the >7,000 rare diseases do not have an FDA approved treatment

This market is accelerating in terms of sales and R&D spend









with CAGR of ~12%

Near-Term Opportunities: Addressing key stakeholder challenges



Patients, Patient Groups, Disease Foundations





Meet CENTOGENE

We are the essential life science partner for data-driven answers in rare and neurodegenerative diseases



CENTOGENE's 3 Strategic Pillars: Fueling revenues, growing CENTOGENE Biodatabank, and building pharma partnerships





DIAGNOSTICS

Highly differentiated testing portfolio offering state-of-the-art WES/WGS/Multiomics

Sample Collection



Proprietary DBS-solution

Multiomic Wet Lab



WES WGS

Multiomics

Multiomic Dry Lab & Bioinformatics



Bioinformatics & Medical Reporting Data Curation

Medical expertise & counselling



Physician support Medical oversight

2 CENTOGENE BIODATABANK

World's largest real-world integrated multiomic data repository in rare and neurodegenerative diseases



3 BIOPHARMA PARTNERSHIPS

Acceleration & de-risking of clinical trials from discovery to market access

Discovery & Early Clinical Development Clinical Research





Market Access & CNTG Biodatabank & Expansion HCP Network





CENTOGENE's 3 Strategic Pillars: Fueling revenues, growing CENTOGENE Biodatabank, and building pharma partnerships





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Market Access & CNTG Biodatabank & Expansion HCP Network





CENTOGENE's Diagnostic Offering: End-to-end Dx solution based on state-of-the-art sequencing & bioinformatics technology



Gold standard for DNA and multiomic sampling from >120 countries, easy logistics for centralized testing



Wet-lab: Highly differentiated testing portfolio in state-of-theart CLIA/CAP certified laboratory with deep medical expertise **Exome Testing Genome Testing** CentoXome® NEW CentoGenome ≥ 97% of the genome at ≥10x \geq 98% of the exome at \geq 20x >20,000 genes ~20,000 genes **Multiomic Testing** Complete Coding and ~8.000 genome map in Non-coding non-coding 1 single clinical pathogenic regions variants genes test CentoMetabolic MOx ≥ 99.5% targeted regions Mitochondrial Mitochondrial covered at ≥ 20 x genome genome 206 genes 37 genes 37 genes 180 Clinically Inherited

> 20 enzymes and biomarkers

Dry-lab (SaaS) solution enabling laboratories around the world to deliver leading diagnostic insights to local patients



Physician network

- Approx. 30,000 active physicians worldwide
- Composed of a range of specialties: neurology, nephrology, pediatrics, hematology

Accreditations

- CAP accredited
- CLIA accredited
- ISO 15189 certified

metabolic

disorders

CAP accredited Biobank, compliant with ISO20387

relevant

variants

Reports & Genetic Counselling

- Proactive, free-of-charge reclassification program
- Access to a network of genetic counsellors and liaisons enables better diagnostics, interpretation and physician counselling

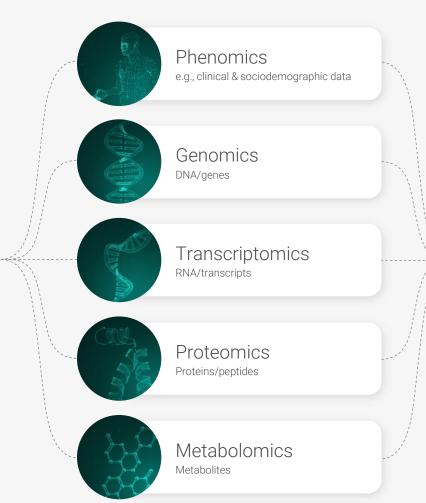
CENTOGENE's Multiomic Datasets: Feeding our bioinformatics pipeline and annotated **CENT**GENE from the CENTOGENE Biodatabank to reach a medically validated diagnosis

Deep Phenotyping

Phenotypes, structural

& functional traits, clinical data

Multiomics Datasets



Dry Lab: Data Integration & Interpretation



1. Diagnostic Bioinformatics

Highly automated bioinformatics pipeline to accelerate diagnostic timelines



2. CENTOGENE Biodatabank

Filtered variant list with annotation from proprietary CNTG Biodatabank



3. Medical Reporting

Comprehensive curation of clinical evidence & medical interpretation by medical experts

CentoCloud® (SaaS): Access CENTOGENE's dry lab capabilities with a CE IVDR-marked virtual dry lab solution to complement local wet labs¹



Your Wet Lab Analysis

Decentralized NGS Analysis

- Local laboratories sequence DNA on trusted Illumina sequencing platforms
- CENTOGENE partnership with Twist Biosciences enables local access to high quality reagents and libraries





- Genome analysis supported for standard Illumina Genome libraries
- Easy file transfer via CentoPortal®

CentoCloud® Modular Dry Lab Solution



Fully automated Bioinformatics Solution & CENTOGENE Biodatabank Annotation

- Fully automated, curated, state-of-theart, CE-marked¹ bioinformatic pipeline to accelerate diagnostic timelines
- Annotation from CENTOGENE's proprietary Biodatabank for variant classification and prioritization





Access to Variant FilterTool

CentoCloud® provides access to an intuitive and automated Variant
 FilterTool allowing for interpretation of genetic results and enabling patient diagnosis¹



High-quality Medical Reporting

 High-quality and accurate diagnostics and reporting by our board-certified human genetic experts



Full Medical Report

 Full medical report with curated medical information, clear diagnostic statement and recommendations for follow-up analyses

CENTOGENE's Accreditations: Quality driven to achieve scientific excellence







- Received accreditation from the College of American Pathologists (CAP) in February 2020
- First biorepository outside U.S. to receive such accreditation

CLIA **ACCREDITATION**

Adhering to the strictest quality criteria to ensure accurate, reliable, and timely patient test results



Meeting international standards for quality and competence particular to medical laboratories

CentoCloud is CE-marked under the In Vitro Diagnostics Directive (98/79/EC) since May 2022

CENTOGENE's 3 Strategic Pillars: Fueling revenues, growing CENTOGENE Biodatabank, and building pharma partnerships





DIAGNOSTICS

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Sample Collection



Proprietary DBS-solution

Multiomic Wet Lab



WES

WGS

Multiomic

Multiomic Dry Lab & Bioinformatics



Bioinformatics & Medical Reporting Data Curation

Medical expertise & counselling



Physician support Medical oversight

CENTOGENE BIODATABANK

World's largest real-world integrated multiomic data repository in rare and neurodegenerative diseases

> 750k Individuals

 $^{\sim}$ 30,000 Active Physicians

> 120 Countries

> 2,500 Diseases

Multiomic Data

Clinical & HPO Data

Biomaterial

Sociodemographic Data

Bio-Medical Analytics

AI/ML

3 BIOPHARMA PARTNERSHIPS

Acceleration & de-risking of clinical trials from discovery to market access

Discovery & Early Clinical Development Clinical Research





Market Access & CNTG Biodatabank & Expansion HCP Network

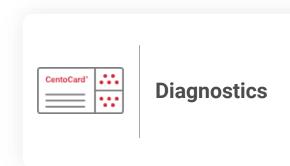




The CENTOGENE Biodatabank: Highly diverse real-world data repository for rare and neurodegenerative diseases



Diagnostics samples fuel the CNTG Biodatabank



CNTG Biodatabank fuels higher diagnostic yields



Diagnostics generate realworld evidence & real-world insights and data business



Biopharma partnerships fuel the CNTG Biodatabank

The CENTOGENE Biodatabank: The breadth and depth makes it a unique resource





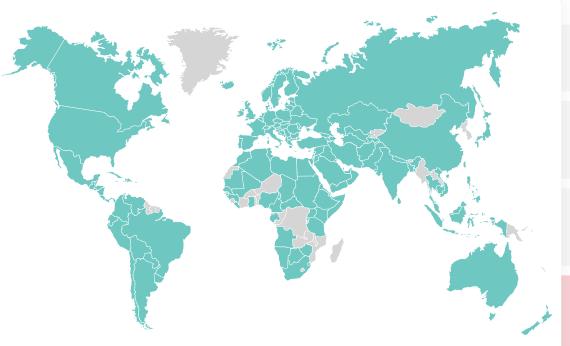
>30,000 via clinical studies

>400,000 biosamples from

>120 countries

~30,000 active physicians in our network

~50% broad research consent
(>70% in recent years)



High diversity in geographies, ethnicities, and patient characteristics

Country represented in Biodatabank Country not represented in Biodatabank

>70 million unique variants

High share of pediatric cases

>13k family data sets

>2,500 rare diseases covered with diagnosed patients

>70% of individuals of non-European descent*

*as compared to ~78% European ancestry in GWAS catalogue per 2019¹

Clinical Diagnostics

Target & Drug
Discovery

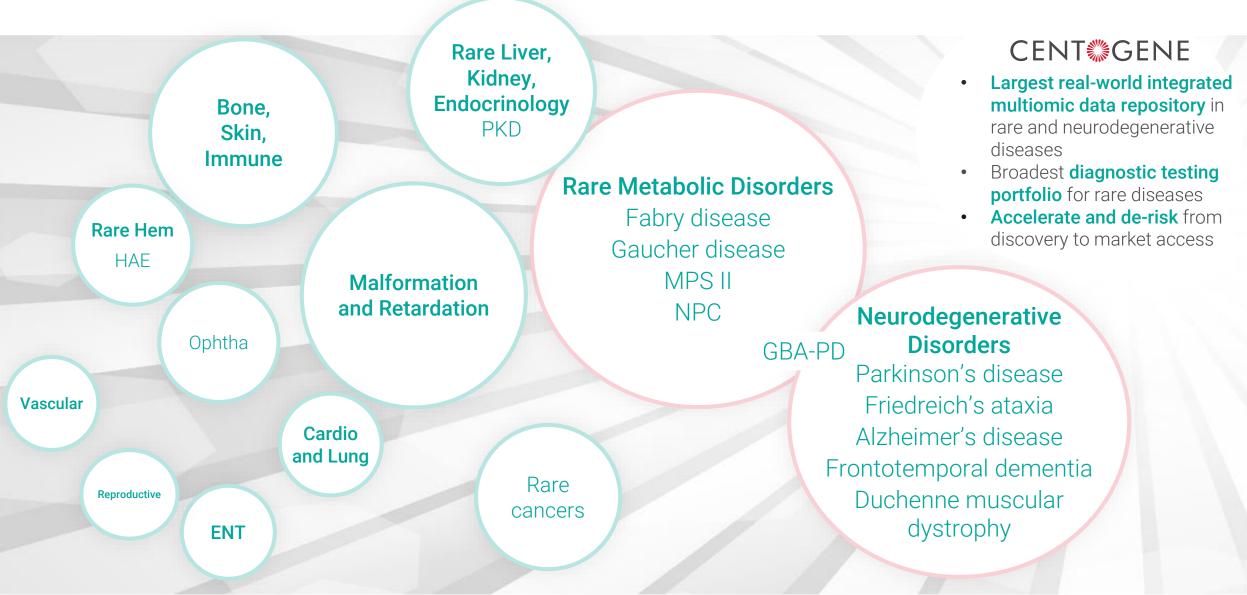
Clinical Study Support

Market Access & Expansion

The CENTOGENE Biodatabank: Insights into 2,500 rare diseases to support



life-saving therapies



CENTOGENE's 3 Strategic Pillars: Fueling revenues, growing CENTOGENE Biodatabank, and building pharma partnerships





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World's largest real-world integrated multiomic data repository in rare and neurodegenerative diseases



120 00011110

> 2,500 Diseases

Multiomic Data

Clinical & HPO Data

Biomaterial

Sociodemographic Data

Bio-Medical Analytics

AI / ML

3 BIOPHARMA PARTNERSHIPS

Acceleration and de-risking of clinical trials from discovery to market access

Discovery & Early Clinical Development Clinical Research





Market Access & CNTG Biodatabank & Expansion HCP Network





CENTOGENE's Unique Pharma Offering: Accelerating and de-risking drug discovery, development, and commercialization



Discovery & Early Clinical Development

Patient-derived Cell Models & Multiomics

Biomarker/Assav Identification & Validation

Clinical Research

POC/ Ph II/III

Patient Multiomic Profiling, Stratification. Modellina

Patient Identification &

Diagnostics for Clinical Recruitment

Observational Studies

Observational Studies & Registries (e.g., epidemiology, Natural History Studies genetic & biomarker profiling)

Market Access & Expansion

Early Patient Access Programs

Patient Stratification. Genetic & Biomarker Profiling, Modelling

Patient ID & Diagnostics for Market Expansion

CENTOGENE Biodatabank & HCP Network

Real-World Evidence and Real-World Insights

(e.g., new and existing reports, clinical feasibility, genetic, epidemiology)

Data Business

(de-identified data cuts & patient cohorts for analysis on third-party platforms)

CNTG Biodata License Fee (e.g., on TRE Platform)

HCP Network

(~30,000 physicians across 120 countries) **Outreach Services**

















Market Access and Expansion: Maximize access and personalize patient, provider, and pharma value



Market Access and Expansion



- Real world registry
- Early access programs
- Patient Stratification
- Genetic and Biomarker Profiling
- Modelling
- Patient Identification
- Patient Diagnostics
- CENTOGENE Biodatabank
- HCP Network





- 2015 Ongoing
- Rare metabolic and rare neurodegenerative diseases
- Provide diagnostic testing services to identify patients with rare metabolic and rare neurodegenerative diseases
- 42 Countries



- 2019 2022
- Duchenne muscular dystrophy (DMD)
- DMD Sponsored testing program (250 samples)
- 5 countries: UAE, KSA, Lebanon, Kuwait, Egypt





- 2015 2022
- Hereditary transthyretin amyloidosis (hATTR) disease
- Sponsored testing program with > 600 samples from 10 countries (Europe) & 125 samples (U.S.)



- 2019 Ongoing
- Identify patients in **DMD &** Aromatic L-amino Acid Decarboxylase (AADC)
- Genetic testing and biomarker analytics for AADC deficiency in 65 countries (LATAM, Europe, MENA)
- >2,500 DMD & >2,900 AADC samples screened

Clinical Development: Accelerate and expand pharma partnerships



Clinical Development



Observational studies

- Epidemiology & Patient finding
- Genetic & biomarker profiling

POC/Ph II/III

- Patient multiomic profiling
- Stratification, Modelling, and Efficacy markers
- Patient identification & diagnostics

CENTOGENE Biodatabank & HCP Network



- 2021
- Hypophosphatasia (HPP)
- Strensiq (innovative enzyme replacement therapy)
- De novo variant identification for HPP and identification of potential new genes causing HPP
- Germany

→ agios

- 2021 Ongoing
- Pyruvate kinase ("PK") deficiency
- Genetic testing and identification of causative mutations, incl. HBA1, HBA2, and HBB genes, in Ph. II/III trials
- 20 Countries (North America, Europe, MENA, APAC, & LATAM)

DENALITHERAPEUTICS

- 2018 Ongoing (ROPAD Study with extensions)
- Parkinson's disease (PD)
- Enroll ~15,000 enrolled patients to date
- 10 countries
- Patients identified with LRRK2 genetic variations may be eligible for interventional clinical studies

Takeda

- 2018 2021
- Gaucher disease
- Longitudinal natural history study (LysoProof) with >1,600 samples analyzed
- 13 countries (EU, LATAM, APAC, MENA)
- Identified and genetically tested
 >250 Gaucher patients

2AInylam

- 2020 Ongoing
- Hereditary transthyretin-related amyloidosis (hATTR)
- Longitudinal study providing a molecular genetic diagnosis of hATTR via NGS and MLPA
- Germany
- 5,000 patients enrolled

alector

- 2021 Ongoing
- Frontotemporal dementia (FTD)
- Enroll and genetically test over 3,000 FTD patients in EFRONT Study
- 7 countries (Belgium, Germany, Greece, Italy, Portugal, Spain, and Turkey)



Target & Drug Screening: Build partnerships around precision and confidence

Target and Drug Screening



Target & Drug Screening

- Patient-derived cell models & multiomics
- Biomarker/Assay identification & validation

CENTOGENE Biodatabank & HCP Network

CENTGENE



- 2021 2022
- Niemann-Pick type C
- Collaboration to generate data set to enable start of drug discovery

CENT#GENE



- 2020 2023
- Gaucher Disease joint drug discovery project
- Joint drug discovery project to identify small molecules reducing biomarker Lyso-GB1 in disease cell models
- Transcriptomic and metabolomic data set enabling patient selection with highest unmet need

CENTGENE

- 2021 Ongoing
- Friedreich's Ataxia, Duchenne Muscular Dystrophy, Hereditary transthyretin-related amyloidosis
- Novel biomarker discovery to stratify and monitor patients for disease severity/progression and to enable the discovery of disease modifiers explaining heterogeneity



- · 2018 2022
- Rare neurodegenerative diseases
- Data Access and Collaboration Agreement granting access to the CENTOGENE Biodatabank for discovery & validation of novel genetic and biochemical targets for the potential development of new therapies for rare diseases

CENTOGENE Capabilities: Fully integrated multiomic solutions for Diagnostics & Pharma



Wet Lab Capabilities:

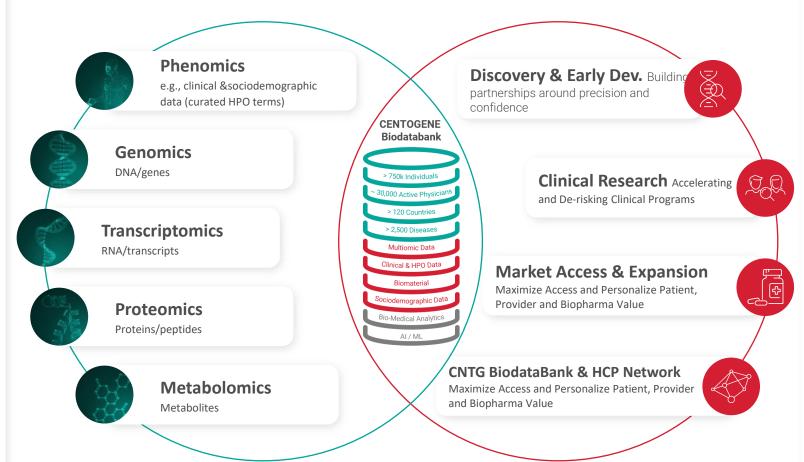
- **Genomics**: CentoGenome®, CentoXome®
- Multiomics: CentoGenome MOx®, CentoXome MOx®, CentoMetabolic MOx®
- Reproductive testing: CentoNIPT®
- NGS Panels Testing: Centogene NGS Panels
- Carrier/Family testing: CentoScreen®, CentoArray®
- Biochemistry Testing: CentoMetablic MOx®
- Single Gene Sequencing: NGS, Sanger, MLPA, FLA, RPA

Dry Lab Capabilities:

- Software as a Service
 offer: CentoCloud®
- Fully automated genetic annotation and interpretation pipeline
- Medical Reporting

Sample Logistics:

- CentoCard®
- CentoPortal®



Discovery & Early Clinical Development:

- Patient-derived Cell Models & Multiomics
- Biomarker/Assay Identification & Validation

Clinical Research:

- Patient Multiomic Profiling, Stratification, Modelling,
- Patient Identification & Diagnostics (Clinical Recruitment)
- Observational Studies & Registries

Market Access and Expansion:

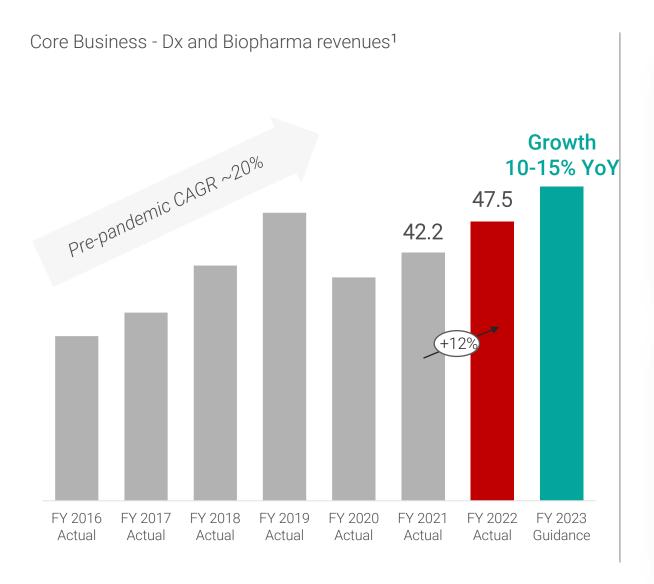
- Early Patient Access Programs
- Patient Stratification, Genetic & Biomarker Profiling, Modelling
- Patient ID & Diagnostics (Market Expansion)

CNTG BiodataBank & HCP Network:

- Real World Evidence & Insights
- Data Business / TRE Platform
- HCP Network

CENTOGENE Financials - FY 2022 - and new Guidance for 2023





Diagnostics

Outgrowing the market

- FY'22 revenues of €31.4 million +18% vs FY'21
- Sales of CentoXome®/CentoGenome® (Whole Exome/Genome Sequencing) up 34% YoY
- 25% of CentoXome® & CentoGenome® upsold to CENTOGENE MOx (multiomic solutions)
- Launched CentoCloud®, CE-marked decentralized SaaS and clinical decision support platform

Returned to growth

Pharma

- FY'22 revenues of €16.1 € million +3% vs FY'21
- Increased activity in the clinical studies of pharmaceutical partners
- GP margins more than doubled to 55%
- Launched data-driven partnering solutions for biopharma and research

Summary



Growth

- Focus on unique and transformative business model
- Expand pharma partnerships
 - Fully execute on our ongoing partnerships and target new pipeline deals
- Keep growing Dx at above-market level
 - Focus on profitable growth
 - Commercial excellence, CentoCloud®, and multiomics

Cost management

- Drive fit-for-purpose organization
- Focus on efficient operations and margin improvement

Cashflow

- Sector is not about growth at all costs
- Diligently manage cash and extend runway





Topline

Bottomline

Runway

Revenues of €47.5M up +12% yoy driven by Diagnostics +18% yoy and Pharma +3% yoy



GP% expansion from 32% to 42% and ongoing cost control resulted in >30% improvement in net results

Segment adj. EBITDA of €13.2M up 69%

^{*} Revenues reflect the Diagnostics and Pharma reporting segments, and do not include the COVID-19 Testing revenues recorded in the period 2020 – Q1 2022. COVID-19 Testing has been reported as discontinued operations since Q1 2022. Prior period financials as revised per FY2022 20F results filing on May 16, 2023. Guidance as communicated per FY2022 earnings announcement on May 16, 2023. Image source: flaticon.com

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Thank you

