

The Essential
Biodata Life
Science Partner in
Rare and
Neurodegenerative
Diseases

CENTOGENE (CNTG) Company Presentation August 2022



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For further information, please refer to the Risk Factors section in our Annual Report for the year ended December 31, 2020, on Form 20-F filed with the SEC on March 31, 2022, 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



- Headquarters Rostock, Germany with locations in Boston, MA, Berlin, Germany, and Rotkreuz, Switzerland
- ~500 employees
- Listed on Nasdaq in November 2019 (Ticker: CNTG)



- FY2021 revenues of €43.5 million
- Guidance* 2022:
 - Revenues of ~ €50 €52 million YoY growth 15-20%



- CENTOGENE Biodatabank, the world's largest real-world data repository for rare and neurodegenerative diseases
- State-of-the-art genomics and multiomics reference lab (ISO, CAP, & CLIA certified)



- >50 collaborations with biotech/biopharma partners, covering over 46 rare diseases
- Market access and expansion, clinical development, target and drug screening

Our ambition is to be the essential biodata life science partner in rare and neurodegenerative diseases

Fueling revenues, growing CENTOGENE Biodatabank, and building biopharma partnerships

3 STRATEGIC PILLARS

1 DIAGNOSTICS

Highly differentiated testing portfolio

Easy logistics via CentoCard & CentoCloud

WES/WGS

Multiomics

Network of ~29,000 active physicians

2 CENTOGENE BIODATABANK

> Fuel CENTOGENE Biodatabank with biomaterial, multiomics, as well as clinical data

Productize CENTOGENE Biodatabank (data monetization) 3 BIOPHARMA PARTNERSHIPS

MARKET ACCESS & EXPANSION

Real world Registry

Early Access Programs

Patient Stratification, Genetic & Biomarker Profiling, Modelling

Patient Identification & Diagnostics

CLINICAL DEVELOPMENT

Observational Studies (e.g., epidemiology, patient finding, genetic & biomarker profiling)

POC/Ph II/III:

Patient Multiomic Profiling, Stratification, Modelling, Efficacy Marker

Patient Identification & Diagnostic

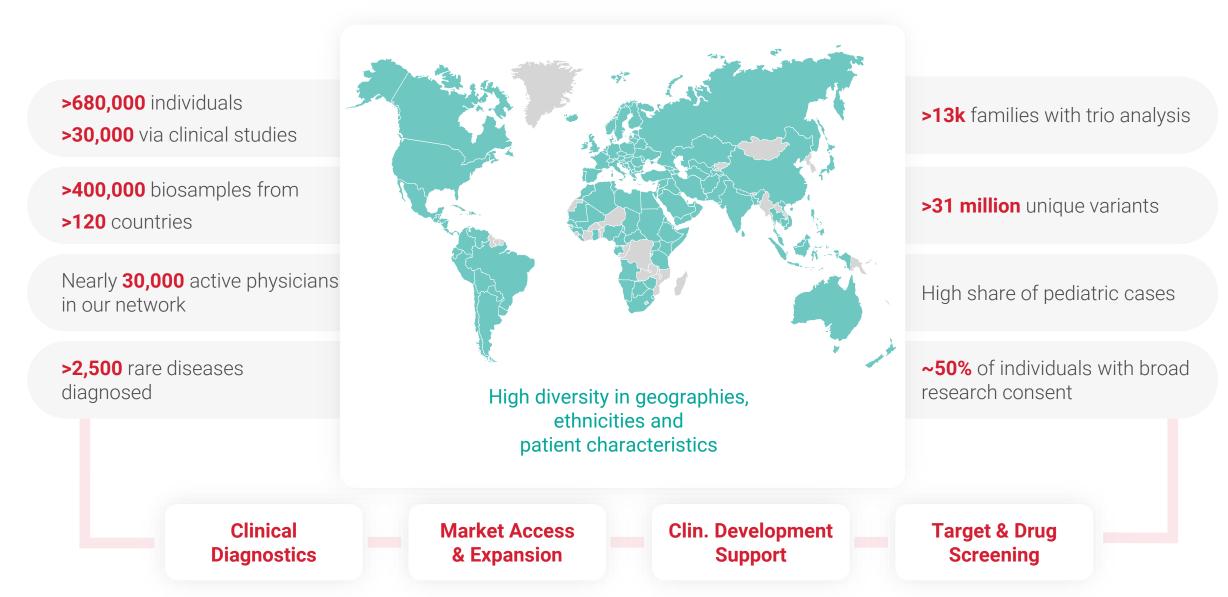
TARGET & DRUG SCREENING

Patient- derived Cell Models & Multiomics

Biomarker/ Assay Identification & Validation

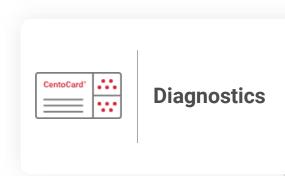
CENTOGENE BIODATA NETWORK (Insight Reports & Biodata licenses)

The Breadth and Depth of CENTOGENE's Biodatabank make it a Unique Resource

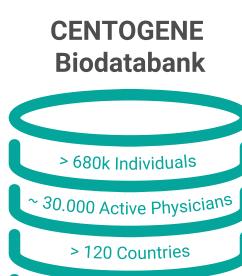


CENTOGENE Biodatabank: the world's largest real-world data repository for rare and neurodegenerative diseases

Diagnostics samples fuel the Biodatabank



Biodatabank fuels better diagnostic yields



> 2,500 Diseases

Multiomic Data

Clinical & HPO Data

Biomaterial

Sociodemographic Data

Bio-Medical Analytics

AI / ML

Insights Reports (clinical, genetic, epidemiology) & Biodata licenses



Biopharma partnerships fuel the Biodatabank

CENTOGENE Pharma Product List - Our ambition is to be the essential biodata life science partner in rare and neurodegenerative diseases

Target & Drug Screening

Patient-derived Cell Models & Multiomics

Biomarker/Assav Identification & Validation

Clinical Development

Observational Studies

POC/Ph II/III

Observational Studies (e.g., epidemiology, patient finding, genetic & biomarker profiling)

Patient Multiomic Profiling, Stratification, Modelling, Efficacy Marker

Patient Identification & Diagnostics

Market Access & Expansion

RW Registry & Early Access Programs

Patient Stratification. Genetic & Biomarker Profiling, Modelling

Patient ID & Diagnostics

CENTOGENE Biodata Network

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

Biodata Licenses

















Market access and expansion: maximize access and personalise patient, provider and biopharma value

Market Access and Expansion



- Real world registry
- Early access programs
- Patient Stratification
- Genetic and Biomarker Profiling
- Modelling
- Patient Identification
- Patient Diagnostics
- CENTOGENE Biodata Network
 - Insight Reports & Biodata Licenses





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





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



REVEAL-CP™

- 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)
- >2500 DMD & >2900 AADC samples screened

Clinical development: accelerate and expand biopharma 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 Biodata **Network**

 Insight Reports & Biodata Licenses

AstraZeneca Rare Disease

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

JENALI THERAPEUTICS

- 2018 Ongoing (ROPAD 1 & 2 Study with extensions)
- Parkinson's disease (PD)
- Enroll and genotype 12,500 patients
- 10 countries
- Success milestone: 10,000 patients enrolled and genotyped

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

→ 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)

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 Biodata Network

 Insight Reports & Biodata Licenses

CENT#GENE



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

CENT GENE



- 2020 Ongoing
- 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

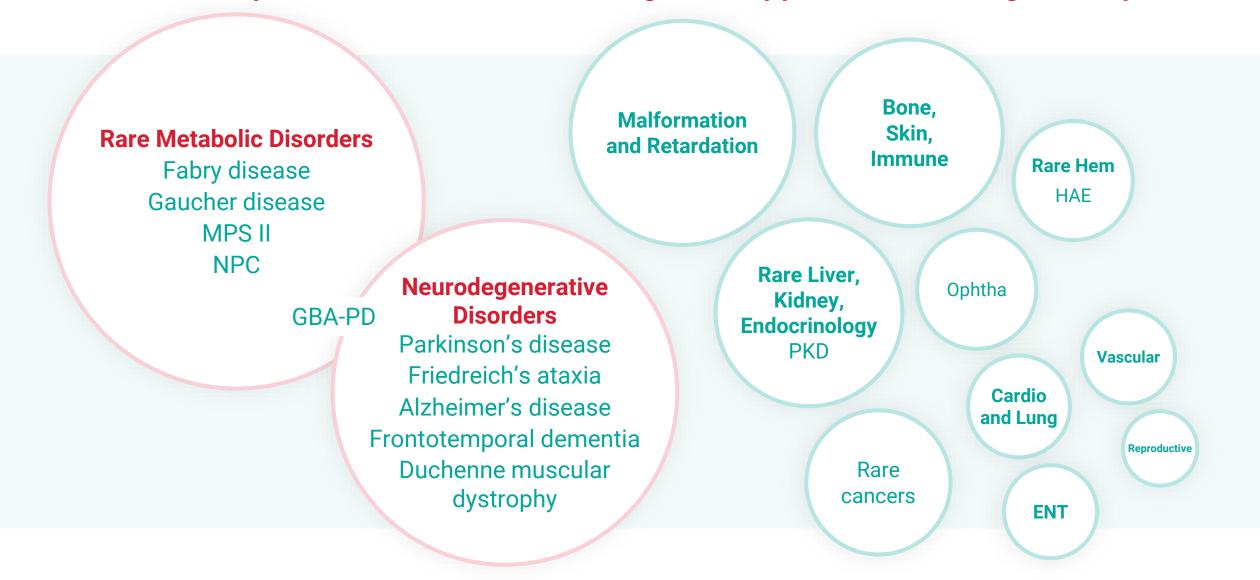
CENTGENE

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

We have knowledge on a broad footprint of over 2,500 rare and neurodegenerative diseases to deepen scientific understanding and support breakthrough therapies

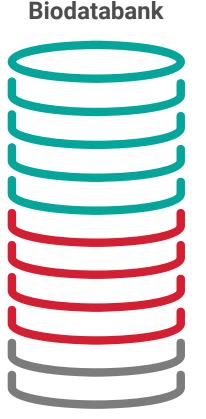


Our genetic Parkinson's disease dataset is unique in depth and breadth

Existing data and biomaterial can support work on diagnosis, disease progression, and treatment



~10,500 research consented individuals + Controls



Multiomic Data

Whole Genomes, Variants, Biomarker, Enzymatic Activity

Clinical Data

Symptoms (HPO), Medication, History, MDS-UPDRS Neuro Test, Age of Onset

Biomaterial

Dried Blood Spots Cards

Sociodemo-graphic information

Gender, Age, Geography, Ethnicity, Family History



Target & Drug Screening

Clinical Trial Support

Market Access & Expansion

First-class data capture and proprietary curation and analysis technologies established the CENTOGENE Biodatabank to predict, accelerate, and personalize treatment

Data capture

Data bioinformatics

Data utilization



Central collection and storage of biosamples

from >120 countries



Clinical data capture



Clinical data extraction & curation



Whole genome sequencing



State-of-the-art mass spectrometry



RNA sequencing



Automated pipelines for

- variant annotation
- prioritization
- · medical reporting
- > 31 million variants
- multiomic analysis and combination expertise



Biopharma Partnerships

Target & Drug Screening

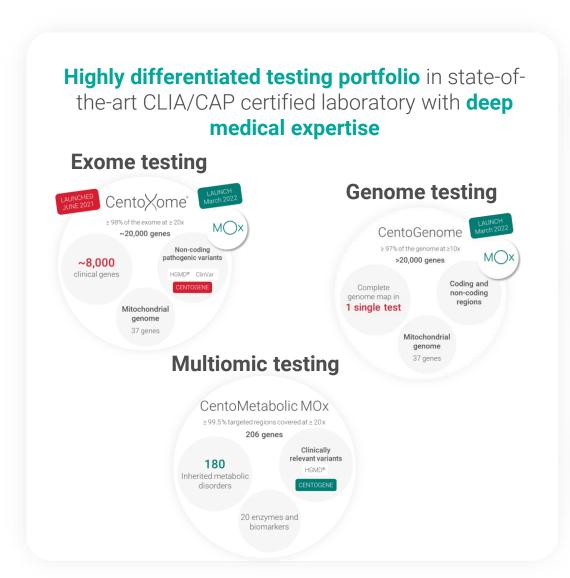
Clinical Development Support

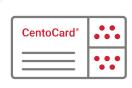
Market Access & Expansion

Value chain supported by advanced bioinformatics and AI tools



Diagnostics: A distinctive offering and services that support efficient and timely diagnosis of rare and neurodegenerative diseases, leading to better treatment and health outcomes





Easy logistics for centralized testing enabling **broad access to multiomics**

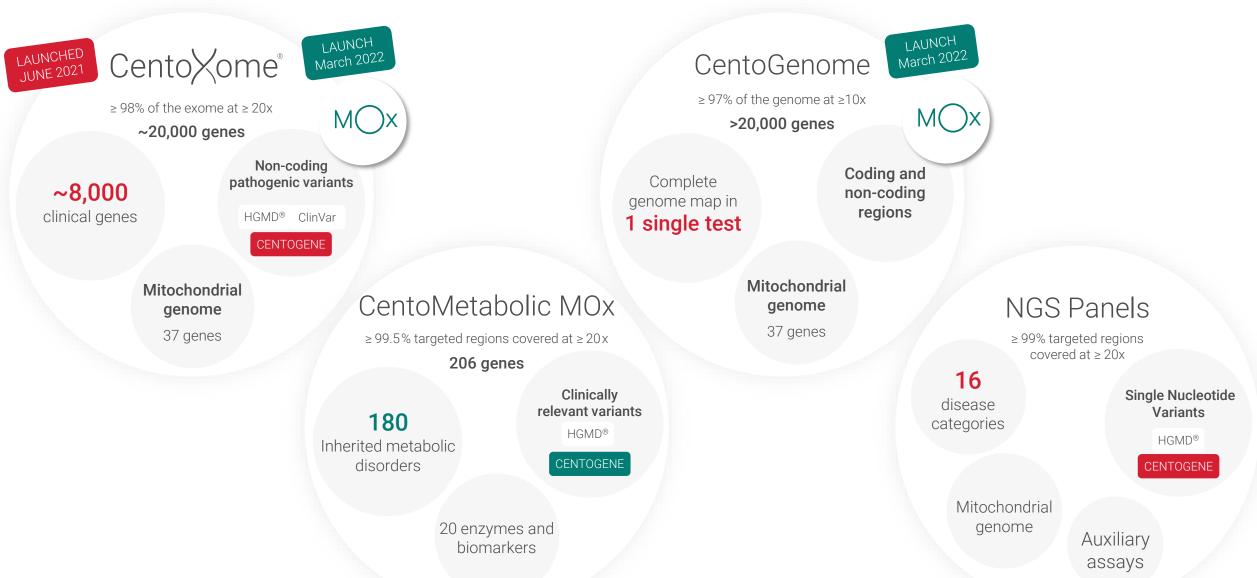


Dry-lab (SaaS) solution enabling laboratories around the world to deliver leading diagnostic **insights to local patients**



Unique global footprint with **network of** ~29,000 active physicians and focus on countries with a high prevalence of rare diseases

We have leading, differentiated core products that drive definitive diagnosis for the benefit of patients & fuel our biodatabank enabling greater understanding of diseases



Quality, data integrity & privacy are core to CENTOGENES business processes, driven by first-class data capture, multi-level QMS, & proprietary curation

CENTOGENE follows the strictest quality criteria to meet our customer's requirements in clinical diagnostics, clinical trial services and research & development. CENTOGENE's QMS integrates into every aspect of our operations, from data integrity and privacy, patient safety, and responsibility to scientific research and innovations.

- CENTOGENE's holistic quality management system (QMS) has been recognized by our CAP, CLIA, and ISO 15189 certifications. CENTOGENE's biobank is the first CAP accredited repository outside the U.S., compliant with ISO 20387
- As a full service company we follow applicable and market-standard good laboratory practice (GLP) and good manufacturing practice (GMP) guidelines
- We continually participate in the international external proficiency testing schemes of the European Molecular Genetics Quality Network (EMQN) and the College of American Pathologists (CAP) to ensure technical and medical competency and proficiency
- CENTOGENE's processes and IT systems are ISO/IEC 27001:2017 certified
 by the independent accreditor datenschutz cert GmbH, ensuring a high level
 of confidentiality, availability, and integrity to all processed data
- CentoPortal is certified for providing safety and privacy according to the "internet privacy standards" by German accreditor datenschutz-cert





Rostock Lab

#8574447 (biorepository)

#99D2049715 #22D2154474

Boston Lab #8417628





Certificate-ID: DSC.843.09.2020





Urkunden-ID: DSC.1096.11.2021

A unique diagnostic portfolio of products, technologies, and enablers

PRODUCTS

CentoCard®

Unique and proprietary CElabeled DBS sample collection kit

CentoCloud®

Cloud-based automated bioinformatics and medical reporting solution

CentoXome[®]

Enhanced WES with uniform coverage across the entire exome and increased Dx yield of up to 20%*

CentoGenome®

Advanced WGS solution capturing one of the most extensive ranges of disease-causing genetic variants

MOx Products

Multiomic
WES/WGS/Panel
solutions integrating
NGS w/ biochemical
analysis to provide a
complete clinical
picture (e.g.
CentoMetabolic®)

TECHNOLOGY

Logistics

SaaS**

NGS**

NGS

Multiomics/ Biomarkers

ENABLERS

Physician network

- Nearly 30,000 active physicians worldwide
- Composed of a range of specialties, with emphasis on neurology

Reports & Genetic Counselling

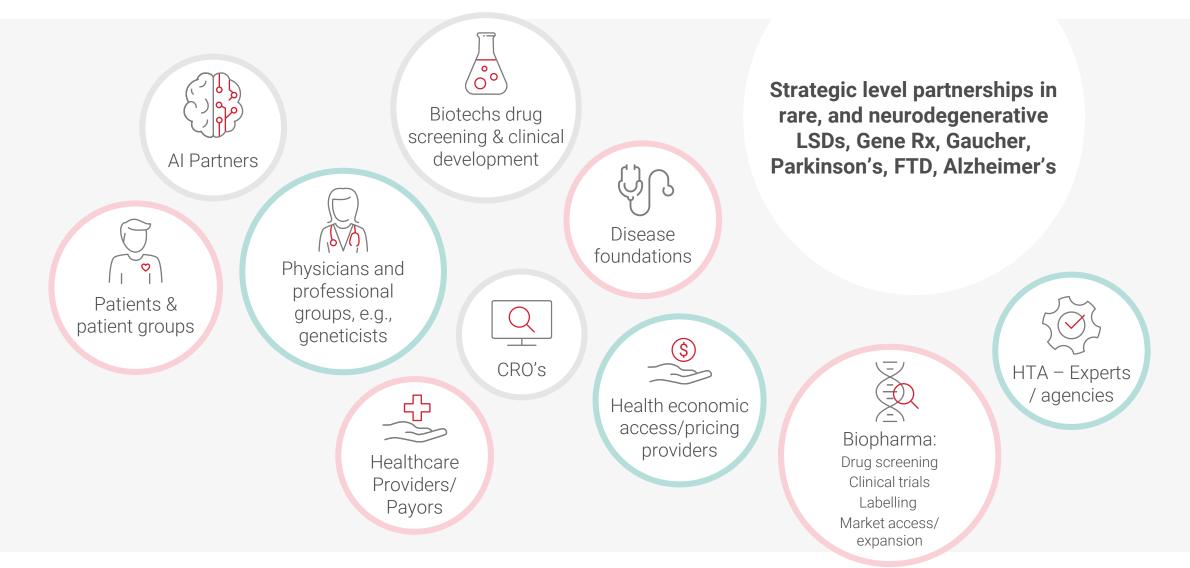
- Proactive, free-of-charge reclassification program
- Access to a network of genetic counsellors and liaisons enables better diagnostics, interpretation and physician counselling

Accreditations

- CAP accredited
- CLIA accredited
- ISO 15189 certified
- CAP accredited Biobank, compliant with ISO20387

^{17 *} Compared to routine WES

We are an essential partner across Biopharma for diagnosing, understanding, and treating rare and neurodegenerative diseases



Pharma partnerships - The opportunity: By 2024, 18% of Rx worldwide will be for rare diseases

Rare diseases CENTOGENE Biodatabank

Largest WW biorepository for rare diseases*

- Estimated 80% of ~7,000 rare diseases are genetic in origin: ~5,6001
- CENTOGENE Biodatabank covers >2,500 diseases²
- Public datasets are ~80% of European descent⁷
- CENTOGENE database rich in bio samples, ethnic diversity and patient consent

Rare disease market

- Rare diseases market expected to grow 11+% to 2024³
- By 2024 rare disease products account ~18%
 ^R sales⁴
- FDA have approved 23 gene/cell therapies to date⁵
- >50% of FDA approvals in 2021 were orphan drugs⁶

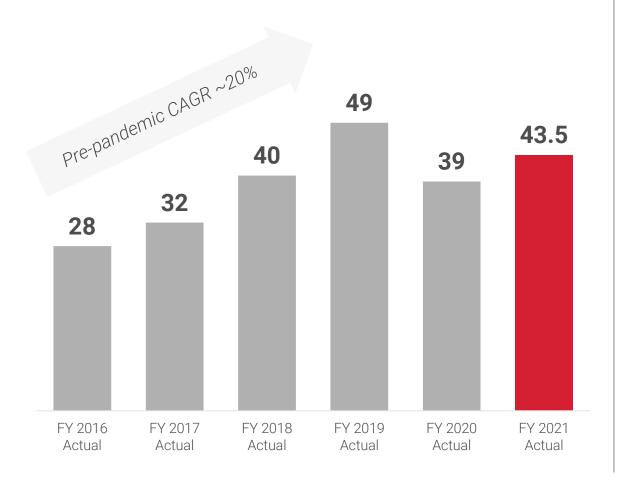
Regulatory and payor scrutiny*

- RD, NDD and gene therapies are typically premium price products with high unmet need and patient pressure
- Regulatory/payor scrutiny will raise standards for approval, access and entry
- Patient profiling/stratification can help to improve labelling/pricing optimization and probability of success

¹ Compare Centogene 20-F for FY2021 p. 56 | 2 CENTOGENE Biodatabank statistic per Dec 31, 2021. | 3 & 4 EvaluatePharma — Orphan Drug Report 2020 p. 2 | 5 Based on FDA approvals per May 2022. [Approved Cellular and Gene Therapy Products <u>LFDA</u>] | 6 EvaluatePharma — Orphan Drug Report 2022 p. 2 | 7 The Scientist (2019) "Lack of Diversity in Genetic Datasets is Risky for Treating Disease", reflecting distribution of ancestry of individuals in genome-wide association studies (GWAS) as of January 2019 | * Statements reflect internal management estimates.

CENTOGENE 2021 Financials

Core Business - Dx and Biopharma revenues¹



Record year for Diagnostics

Diagnostics

- Full year 27.9 € million +26% vs FY 2020
- 57,000 test requests +36% vs FY 2020
- WES and WGS ~ 20 € million +25% vs FY 2020

Impacted by COVID

Biopharma

- Full year 15.6 € million -8% vs FY 2020
- Q4 revenues 6.5 € million +40% YoY
- 45 active collaborations by end 2021
- Driven by partnerships in patient identification and clinical development

CENTOGENE Q1 2022 Financials

Dx and Biopharma revenues in € million¹

Double-digit growth rate

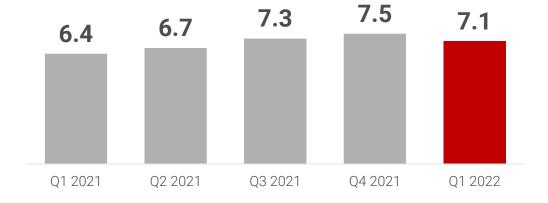
Diagnostics

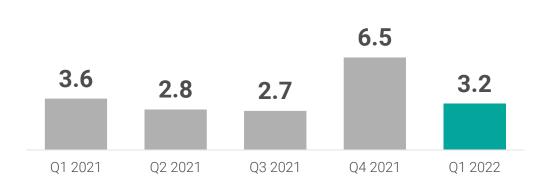
- Q1 2022 €7.1 million +11% vs Q1 2021
- Q1 2022 16,300 test requests +24% vs Q1 2021
- WES and WGS ~ 3.7 € million +18% vs Q1 2021
- Product portfolio updates on CentoCloud & MOx

Longer recovery post COVID impact

Biopharma

- Q1 2022 € 3.2 million, -10% vs Q1 2021
- Primarily attributable to impact of COVID-19 on slowing pharma programs and longer sales cycle
- 42 active collaborations per March 31, 2022
- Post Q1 2022 extended Agios & Takeda contracts





CENTOGENE Q1 2022 Financials

P&L and balance sheet highlights in € million¹

	Q1 2021	Q1 2022	delta
Revenue	10.0	10.3	+0.3
Cost of sales	6.2	6.5	+0.3
Gross Profit	3.8	3.9	+0.1
Gross Profit %	38%	38%	-
Research & development expenses	4.3	4.6	+0.3
General Administrative expenses	11.6	7.9	(3.7)
Selling expenses	1.9	2.4	+0.5
Impairment of financial assets	0.1	0.2	+0.1
Other operating income	0.4	0.7	+0.3
Other operating expenses	0.0	0.0	-
Operating loss	(13.9)	(10.5)	+3.4

Balance sheet highlights

(in € million)

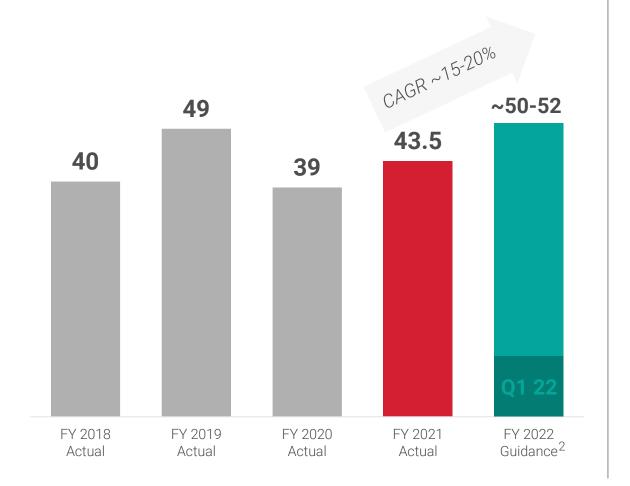
(in € million)	Q1 2022	Q4 2021	delta
Cash & cash equivalents	42.7	17.8	24.8
Debt outstanding ²	(43.0)	(22.5)	(20.5)
Net debt	(0.3)	(4.7)	4.4

^{22 1} Selected information, may include rounding differences

² Debt outstanding includes non-current loans, non-current lease liabilities, current loans and current lease liabilities.

CENTOGENE Guidance: 2022 poised for post-COVID recovery and refocus on core business and growth strategy

Core Business - Dx and Biopharma revenues¹



FY 2022 Guidance²



- \$62 million financing (~€55 million) announced Q1 2022:
 - €15 million PIPE & \$45 million debt ³
- Q1 of € 10.3 million up 3% year-over-year acceleration in second half driven by biopharma
- Q1 COVID-19 revenues of ~€19 million; exited in Q1 20224

CENTOGENE 2022 Priorities: Focus on growth, execution, fit-for-purpose organization



Meet/Exceed Revenues Guidance

- 50-52 € million total revenues
- ~15-20% growth y-o-y



Fuel Biopharma Partnerships

- Fully execute on our existing >20 ongoing partnerships
- Target ~20+ new pipeline deals



Optimize Dx Revenues Growth

- Focus on profitable growth
- Commercial excellence, CentoCloud & multiomics



Fit-for-Purpose Organization

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



Thank you

To our investors
To our partners
To our associates

MISSION

Together let's cure 100 rare diseases (and others) in the next 10 years

