

2Q25 Presentation

10 July 2025

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Efficient diagnostics for better treatment decisions



Gentian Diagnostics at a glance





A MedTech company targeting \$1.8bn serviceable diagnostic market with 5-10% annual growth



Appealing value proposition, focused strategy and lean business model.



Industry-leading capabilities - strong focus on in-house R&D and Operations



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At commercial phase and making profit

High quality standards (IVDR and ISO13485 certified) and focus on ESG





Appealing value proposition









Value proposition

Many clinically relevant diagnostic biomarkers are available only on slow and inefficient platforms

By leveraging existing, open-channel instrumentation, Gentian converts these tests to high-throughput analysers

Faster results \rightarrow better treatment decisions

Up to 10x improved efficiency and cost savings



Lean business model

Partnerships with global IVD companies

OEM partnerships to secure broad roll-out and acceptance of product

Distributors in select markets

In selected markets we do not serve directly

Direct end-users

Large central laboratories in selected markets





Focused strategy targeting large, existing market with our world-leading knowledge on PETIA*



Highly specialised with PETIA assays on high volume diagnostic segments.

Addressing customer needs with worldclass R&D, production, clinical data generation and regulatory support.

Leveraging growing volumes, cost pressure and market consolidation trends.

Sustained growth with diverse product pipeline, technological improvements for PETIA, or via adjacent new technologies.







Key disease areas: inflammation & infection, kidney disease, heart failure





Key drivers for long-term growth and value creation

Five **established products** with solid growth potential

Prove clinical relevance of GCAL® and bring NT-proBNP to market

Bring a steady stream of new high-impact diagnostic tests to market

Secure one new contract with a global commercial partner per year

Grow gross margin from ~50% to 60%+ through economies of scale

Long-term EBITDA margins of 40%







Targeting a serviceable market of USD 2.2bn*









*Kalorama 2024, Company estimates including RBP



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Highlights

2Q highlights: Good sales growth in Q2 and H1 vs. '24 – some one-off COGS events impacted gross margin and EBITDA

2Q25 financials and key milestones

Sales MNOK 43.6 +14% vs 2Q24	Sales to US +156% vs 2Q24
EBITDA	Gross margin
MNOK 1.7	44%
MNOK 6.8 in 2Q24	57% in 2Q24

Additional highlights

- Revenue of NOK 88.1 million in 1H25 up 15% vs 1H24 (13% organic growth).
- Cystatin C +31% in 2Q25 vs.2Q24. Continued momentum in China indicates a return towards a normalised supply situation.
- US sales growth of 157% in 2Q25, and 93% in 1H25 with additional new accounts established for Cystatin C and cCRP.
- EBITDA of NOK 15.7 million in 1H25 versus NOK 11.6 million in 1H24.
- For our first-in-class turbidimetric total NT-proBNP assay, we completed further calibrator adjustments and clinical sample testing across three key clinical analyser platforms. Also, securing access to new patient cohorts supports the regulatory submission plan aiming for 2026 commercial launch.



Both the US and Asia growing by ~30% followed by Europe

Highlights

- Sales growth of 14% (14% organic) vs. 2Q24.
- Another strong quarter for Cystatin C (+31%) with good order book for Q3 from Beckman China.
- Strong momentum in the US continued in Q1 (+157% in 1Q25 vs 1Q24), and 57% adjusted for a changeover in a customer inventory from Europe to the US.



Sales revenues (MNOK)

Sales revenue - geographic split							
MNOK	2Q25	2Q24	YTD25	YTD24	2024		
US	7.2	2.8	11.0	5.7	12.2		
Europe	26.6	28.6	57.2	56.5	116.2		
Asia	9.8	6.9	19.9	14.6	23.7		
Total	43.6	38.3	88.1	76.8	152.1		

Sales revenue - product split

MNOK	2Q25	2Q24	YTD25	YTD24	2024
Cystatin C	17.4	13.3	35.1	28.2	50.6
fCAL [®] turbo	12.8	15.0	27.6	28.7	61.3
Third-party products	6.4	4.6	11.5	9.3	18.3
Other	7.1	5.4	14.0	10.6	21.8
Total	43.6	38.3	88.1	76.8	152.1



Stable cost development

OPEX Capitalised R&D expenses



Operating expenses

MNOK	2Q25	2Q24	YTD25	YTD24	2024
Sales and marketing expenses	7.3	6.4	13.4	12.9	28.1
Administration expenses	6.8	6.6	13.1	12.6	21.7
Research and development expenses	6.7	5.1	11.8	11.2	21.9
Total	20.8	18.1	38.3	36.6	71.7

- Operating expenses ended at 20.8 MNOK in 2Q25 compared to NOK 18.1 million in 2Q24.
- Capitalised R&D expenses was MNOK 2.2 in 2Q25 compared to MNOK 1.4 in 2Q24.



Gross margin heavily influenced by production issues



Gross and EBITDA margin %

- Drop in gross margin related to raw material issues that transferred over to the production process.
- High amount of scrapping and additional work for the operations team resulting in significantly higher unit cost for one of our major products.
- Production stabilised in June and ambition of gross margin in the 55%-60% area is maintained.



Continued EBITDA improvement

EBITDA development (MNOK)



- EBITDA 1H25 of NOK 15.7 million vs 11.6 million in 1H24.
- 2Q EBITDA influenced by weak gross margin.



Solid cash position

2Q25 balance sheet and cash flow

Cash	Capex
MNOK 80.2	MNOK 2.5
MNOK 81.0 in 2Q24	MNOK 1.8 in 2Q24
FCF	Equity ratio
MNOK -1.5	85.4%
MNOK -3.3 in 2Q24	84.2% in 2Q24

Capital priorities

- NOK 4.5 million decrease in cash from 31 December 2024.
- Solid cash position of NOK 80.2 million
- Dividend of NOK 6.2 million paid in 2Q25.
- No interest-bearing debt
- Long-term net working capital/sales assumed at ~30%.





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Product update

TIDEMO

gentian

17-212-2

Cystatin C 31% growth from continued US momentum and stabilised China business



Sales of Cystatin C last 2 years (MNOK)



- US, China and South Korea drove sales in 2Q.
- Orders worth NOK 9.4 million from Asia were shipped during 2Q. Solid orderbook for Q3.
- In the US a total of 31 new Cys C customers added in 2025. Positive impact because of increased market presence and intensified collaboration with the company's distribution partners.



fCAL[®] turbo Sales impacted by order phasing

Sales of fCAL[®] turbo last 2 years (MNOK)





- Sales dropped by 15% YoY to NOK12.8 million for 2Q25.
- 2Q was effected by stocking during previous quarters and phasing of orders.
- Overall, the outlook for 2025 remains unchanged with additional distribution agreements in place expexted to start contributing to sales during H2.



Other products with highest quarterly sales ever delivering 32% YoY growth

Sales of other products last 2 years (MNOK)



- 2Q sales of NOK 7.1 million, up 32% compared to the same quarter 2024.
- fPELA turbo and cCRP had another strong quarter at high double-digit growth in 2Q25 vs. 1Q24
- GCAL at single digit growth in Q2 good progress YTD.



Third-party products Record sales quarter at NOK 6.4 million

Sales of third-party products last 2 years (MNOK)



- Record 2Q sales up by 38% compared to 2Q last year.
- Increased customer base across the region and higher testing levels driving growth.
- Our investments in stronger market presence paying off.



US and Asia driving growth in Q2

Sales by region last 2 years (MNOK)







R&D update and summary

R&D Update

- The early-stage development project, in partnership with a global IVD player, is progressing towards the end of the proof-of-concept phase and getting ready to move to optimization.
- Additionally, Gentian has started exploration of a next-generation technology platform. Early work shows promising results with promise of detection capabilities substantially below those achievable with traditional turbidimetric methods, potentially enabling entry into biomarker markets formerly not available for clinical chemistry analyser platforms.
- We are re-visiting the pipeline candidate list and the decision for our next product development target will be made during H2 2025.

R&D Spend break-down

MNOK	2Q25	2Q24	YTD25	YTD24
Technical and Clinical support	2.6	2.2	4.7	4.8
Pipeline development	4.1	3.0	7.1	6.4
Capitalised development expenses	2.2	1.4	4.2	3.9
Total	8.9	6.6	16.0	15.1

Technical and clinical support relates to sending on products that are developed and on the market.

Pipeline development are expenses on products under development.



Gentian's NT-pro-BNP late-stage development

Selected total NT-proBNP option advancing towards final validation phase



About NT-proBNP

Measuring NT-proBNP levels support diagnosis of heart failure. The Gentian assay will be the first test of its kind available on high-throughput analyzers which should increase laboratory productivity and reduce overall costs. Unpredictable and individual variation in glycosylation of NTproBNP creates the opportunity for clinical differentiation using the Gentian NT-proBNP assay, especially in underserved patient subgroups. The company is currently investigating the scope for clinical evidence generation.

2Q highlights:

- Following the strategic decision to align the product with a total NT-proBNP format, the team completed further calibrator adjustments and clinical sample testing across three key clinical analyser platforms. This refined calibration has yielded enhanced precision at low analyte concentrations.
- Some instrument alignment challenges were resolved.
- Key reagent and calibrator components now demonstrate long-term stability in both real-time and accelerated testing conditions.
- Gentian also advanced its clinical validation plan, securing access to new patient cohorts to support the regulatory submission. With the preparation of the IVDR dossier progressing on schedule, the project remains well-positioned to enter its final validation phase in the second half of the year.
- Our aim to introduce the assay as a research-use-only product in the second half of 2025 and full commercial launch in 2026 remain unchanged (depending on regulatory review timelines).





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Q&A



Appendix

P&L highlights

MNOK	2Q25	2Q24	YTD25	YTD24	2024
Sales	43.6	38.3	88.1	76.8	152.1
Cost of goods sold	-24.2	-16.6	-40.4	-34.8	-69.3
Gross profit	19.3	21.7	47.2	42.0	82.8
Other revenues	0.9	1.0	1.8	1.7	4.6
R&D expenses	-6.7	-5.1	-11.8	-11.2	-21.9
Sales and marketing expenses	-7.3	-6.4	-13.4	-12.9	-28.1
Administrative expenses	-6.8	-6.6	-13.1	-12.6	-21.7
Operating profit	-0.5	4.5	11.2	7.1	15.7
Net financial items	0.2	0.2	-1.4	1.8	-2.5
Net profit (loss)	-2.0	4.7	5.8	8.9	45.3



Balance sheet highlights

MNOK	2Q25	2Q24	2024
Inventory	51.7	41.2	45.9
Accounts- and other receivables	35.5	23.5	31.3
Cash and cash equivalents	80.2	81.0	84.7
Total non-current assets	63.8	41.1	67.7
Total assets	231.2	186.9	229.7
Total paid-in equity	318.9	315.7	316.3
Total retained equity	-121.4	-158.4	-122.2
Total equity	197.5	157.4	194.1
Total non-current liabilities	3.2	7.8	5.5
Total current liabilities	30.6	21.6	30.1
Total equity and liabilities	231.2	186.9	229.7



Cash flow highlights

MNOK	2Q25	2Q24	YTD25	YTD24	2024
Operating activities	0.9	-1.5	7.5	1.1	13.5
Investing activities	-2.5	-1.8	-4.5	-5.0	-11.0
Financing activities	-7.4	-1.3	-8.7	-2.5	-5.0
Changes in cash and cash equivalent	-9.0	-4.6	-5.6	-6.4	-2.4
Cash and cash equivalent at the beginning of period	88.7	85.6	84.7	87.6	87.6
Cash and cash equivalent at the end of period	80.2	81.0	80.2	81.0	84.7



Achieved 26% p.a. sales growth last six years



Partnerships prove viability of go-to-market model

SIEMENS ... Healthineers Global distribution agreement for GCAL[®], initial roll-out in Europe



Long-standing commercial partnership for Cystatin C



Partnership for fCAL[®]turbo initiated through Bühlmann Laboratories



Dedicated and experienced management team





Njaal

Kind

CEO Matti Heinonen

CFO & COO CGO Markus Jaquemar



CSO Dr. Alexandra Havelka



СТО

Dr. Frank

Frantzen

VP R&D Dr. Torsten Knüttel



VP QA & RA Anne-Mette Horsrud Akre

20+ years of relevant industry experience across management positions

Track record from leading global diagnostics companies in across all phases













Board of directors

Hilja Ibert	Kari E. Krogstad	Kjersti Grimsrud	Runar Vatne	Christian Åbyholm
Chair of the Board	Board member	Board member	Board member	Board member
Dr. Hilja Ibert has more than 25 years' experience from the international diagnostic industry, including VP International Diagnostic Solutions at Hologic and senior positions within Becton Dickinson and bioMerieux. She was previously the CEO for miDiagnostics in Belgium. In 2018, she was appointed CEO of Gentian Diagnostics ASA, a position she served until May 2024. She is currently a board member in Gradientech and VitaDx.	Kari Krogstad has more than 25 years of experience from the biomedical industry, from commercial leadership roles within the pharma, biotech and medtech sectors. Ms. Krogstad has held her current role as President and CEO at Medistim ASA since 2009. She was previously General Manager at Invitrogen Dynal. Ms. Krogstad holds a Cand. Scient. degree in Molecular Biology from the University of Oslo as well as a Business degree from IHM Business School.	Kjersti Grimsrud is currently President and COO of Infusion care at Convatec plc, where she has spent the last 5 years. She has over 30 years' experience in MedTech and IVD companies with roles in science, operations and commercial in Axis-Shield ASA and Alere Inc./Abbott, where she last held the position of VP Commercial EME (Europe Middle East) and International (APAC). Ms Grimsrud served as a board member of Biotec Pharmacon (now ArcticZymes technologies) from 2011 to 2015. Ms. Grimsrud holds a master's degree in biotechnology from the Norwegian University of Science and Technology in Trondheim.	Mr. Vatne is the principal and owner of Vatne Capital, a family office investing in financial assets and real estate. He has extensive experience from the real estate sector, primarily from Søylen Eiendom, a leading Oslo based real estate company which he co-founded in 2004. Prior to Søylen Eiendom, Mr. Vatne was a Partner and stock broker in Pareto Securities. Mr. Vatne served as board member of Gentian Diagnostics from November 2019 to May 2022. Mr. Vatne and companies controlled by him currently own 15.12% of the outstanding shares in Gentian Diagnostics ASA.	Christian Åbyholm is a partner in Kvantia AS where he joined in 2007. Prior to joining Kvantia AS, Christian worked as Head of Department within Mergers and Acquisitions in Norsk Hydro and as Senior Vice President in business development in Aker RGI. Christian has also worked in London as an Associate in Equity Research in Morgan Stanley where he was part of the number one European Paper and Packaging team ranked by Institutional Investor. Prior to that, Christian worked as an Analyst in Merrill Lynch's Investment Banking division. Mr. Åbyholm is a CFA charter holder and has an MBA from IMD and a Siviløkonom degree from Norwegian School of Economics and Business Administration. In addition, Christian has
Dr. Ibert holds a PhD degree in Nutrition Science from the University of Bonn, Germany.				completed first two years of law school at University of Oslo.

Caaby AS, a wholly owned company by Mr. Åbyholm owns 173,500 shares in the company. Kvantia AS and its subsidiaries (Victoria India Fund AS and Obligasjon 2 AS) own 1,992,208 shares in the company. In addition, Christian Åbyholm is Chairman in INSR ASA and Norda ASA, which both own 614,215 shares in the company. The combined shareholding corresponds to 22% of the outstanding shares in Gentian Diagnostics ASA.



Top 20 shareholders

Shareholder	No of shares	%
Vatne Equity AS	2 110 224	13.68 %
Kvantia AS	1 803 368	11.69 %
Carpe Diem Afseth AS	797 516	5.17 %
Norda ASA	716 099	4.64 %
DNB Carnegie Investment Bank AB	681 000	4.42 %
Safrino AS	649 700	4.21 %
Insr ASA	614 251	3.98 %
J.P. Morgan SE	523 631	3.40 %
DNB Bank ASA, Meglerkonto Innland	447 536	2.90 %
Verdipapirfondet Delphi Norge	384 572	2.49 %
Verdipapirfondet DNB Smb	341 338	2.21 %
Portia AS	300 000	1.95 %
Krefting, Johan Henrik	298 000	1.93 %
Intertrade Shipping AS	257 716	1.67 %
Lioness AS	220 000	1.43 %
Marstal AS	212 407	1.38 %
Sp Capital 22 AS	200 000	1.30 %
Silvercoin Industries AS	187 455	1.22 %
Caaby AS	173 500	1.12 %
T.D. Veen AS	164 967	1.07 %
Other Shareholders	4 339 070	28.13 %
Total Shares	15 422 350	100.00%



