

Interim Report, Q1 2022

May 11, 2022



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Agenda

- o1. Highlights from Q1 2022
- o2. The NGAL Test Regulatory Studies
- o3. Q1 2022 financial result
- o4. Financial Guidance 2022

Highlights from Q1 2022





Highlights from Q1 2022

- Strong performance of The NGAL Test - US RUO sales up 116% and global sales up 70% compared to Q1 2021
- Completion of pediatric patient enrollment for US FDA submission on-track for end of Q2 2022
- Strategic focus on preparing BioPorto for US breakthrough
- Fully subscribed share offering provides DKKm 93 in net proceeds and runway for strategic execution

Launch an FDA Approved Product in the US

Strategic Focus



Drive NGAL Test Market Adoption & Have a Pipeline of High Medical Value Products

- Complete clinical trials & submit The NGAL Test to FDA
- Grow rest-of-world revenues and NGAL awareness through focused distribution resources and tools
- Build US commercialization team to market the clinical value of the NGAL biomarker, and nurture key relationships at target accounts
- Leverage our antibody library and university relationships to cost-effectively build the innovation pipeline



Strengthen the Company to Scale & Execute

- Strengthen our key supplier relationships and implement scalable manufacturing processes
- Build robustness and ensure Quality Systems are FDA and IVDR audit-ready
- Prepare business processes for efficient and scalable growth



Attract, Develop & Retain the Best and Brightest Employees Aligned with our Values

- Proactively recruit the most qualified talent to drive success
- Embrace flexible work environments enabling the ability to recruit from a larger pool of candidates
- Motivate and incentivize employees to stay & build shareholder value

Pediatric Trial Enrollment On-track to be Completed end-Q2 2022, To Be Followed by FDA Submission



Pediatric study



1 in 4 affected with AKI during hospitalization (ICU)¹

Predict AKI Risk in an Intensive Care Setting

- Urine samples
- Predict Stage 2/3 AKI

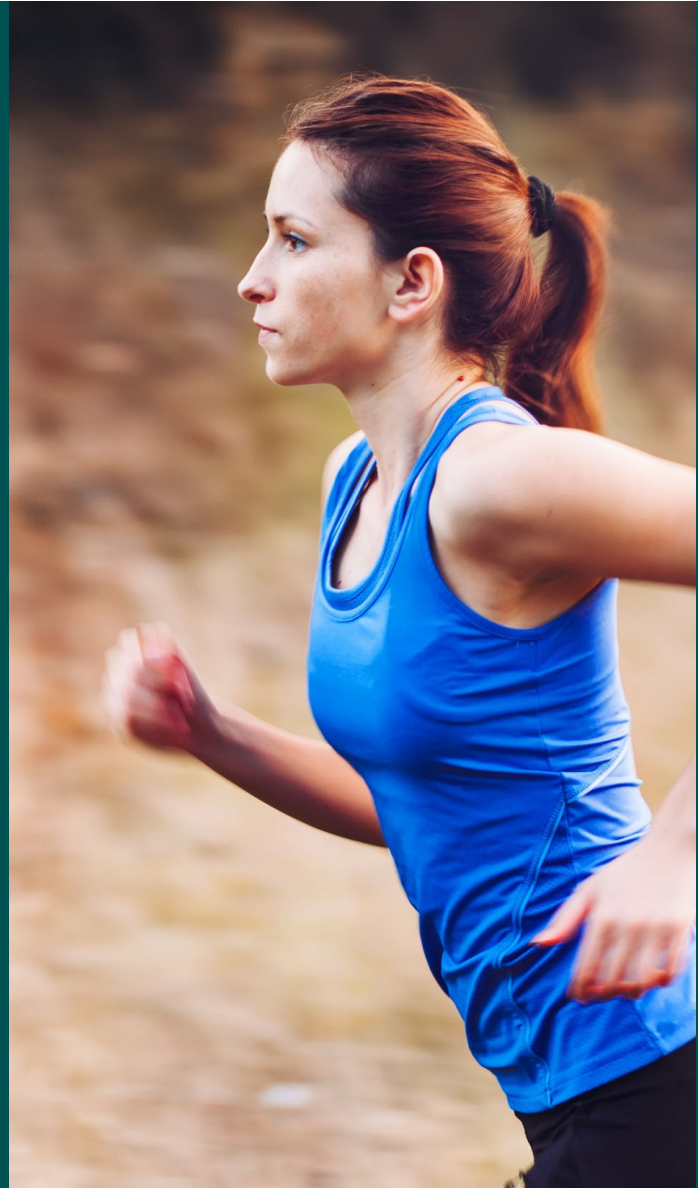
- 15 hospital sites in US clinical trial for The NGAL Test in pediatrics
- BioPorto expects to finalize pediatric trial enrollment in 1H 2022, followed by assembling the FDA submission package & De Novo application with Breakthrough Designation
- FDA targets up to 150 calendar days after submission to respond... ***excludes time for the company to respond to FDA inquiries, which can include more data generation and “stops the clock”***

1) Kaddourah A, et al. *N Engl J Med.* 2017; 376(1):11-20.

Q1 2022 Financial Results



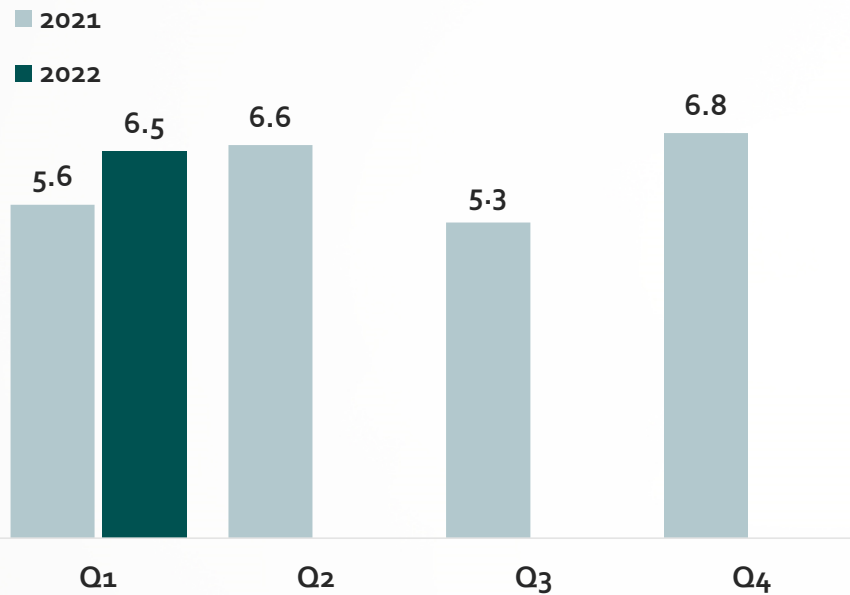
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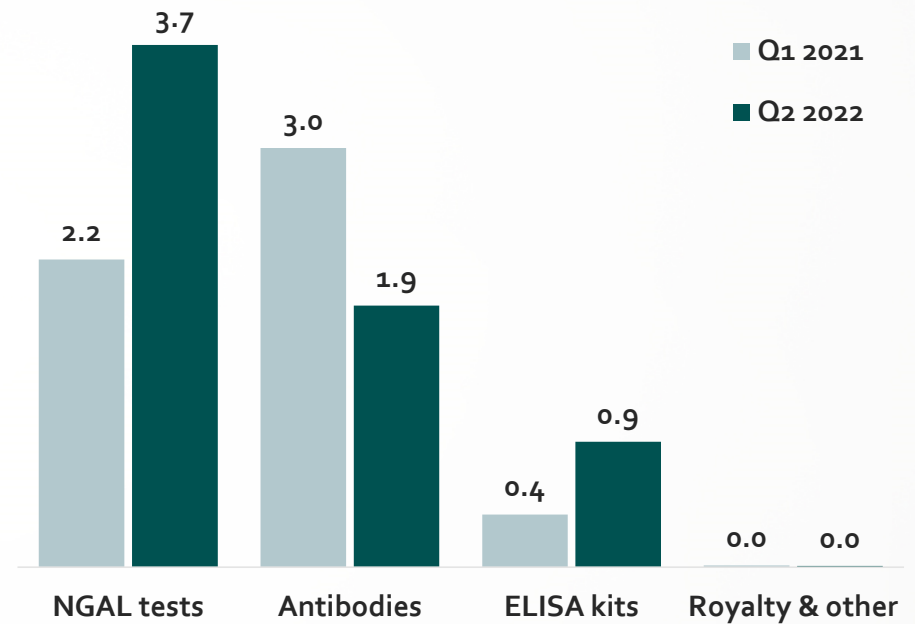


70% Increase The NGAL Test Sales Drives 17% Revenue Growth in Q1 2022 vs. Q1 2021

Revenue by Quarter (DKKm)



Revenue by Product Category (DKKm)



Strong Performance of The NGAL Test Sales in the US and ROW

US NGAL

↑ 116%

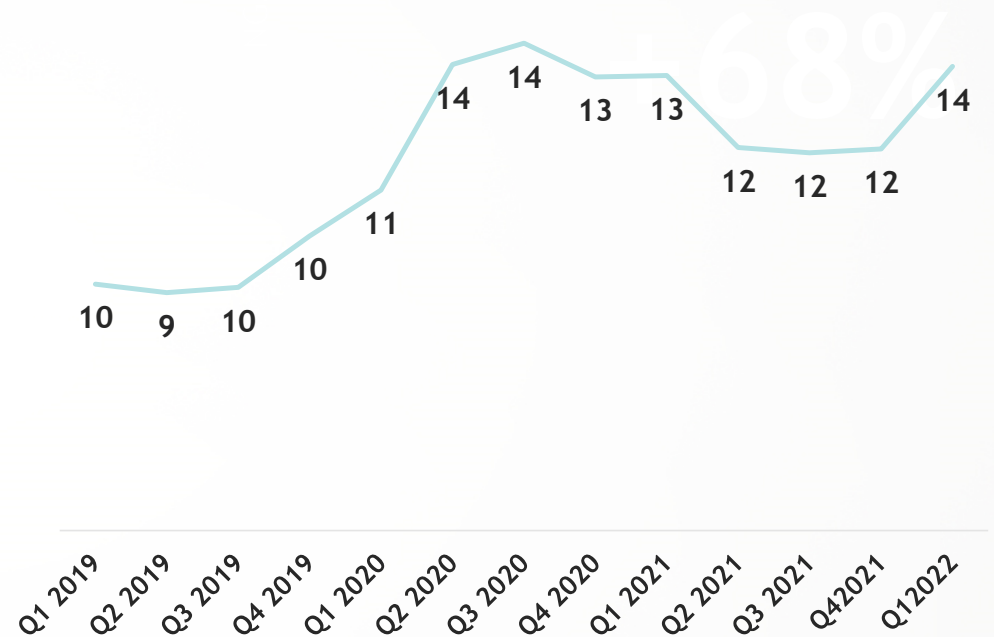
Increase in total revenue from US NGAL sales from Q1 2021 to Q1 2022

Overall NGAL

↑ 70%

Development in global sales of NGAL from Q1 2021 to Q1 2022

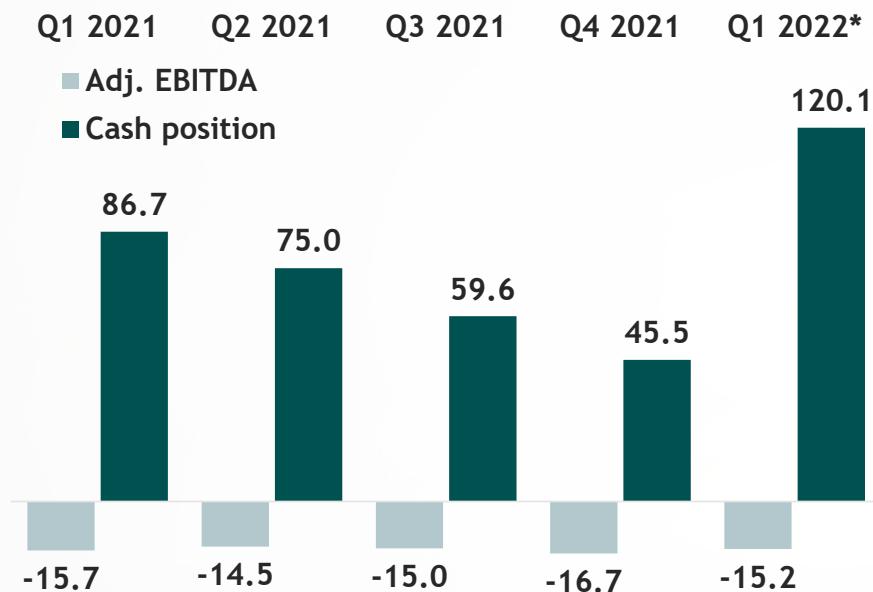
NGAL Product sales by Quarter (LTM, DKKm)





DKKm 0.5 Improvement in Adjusted EBITDA Loss; Cash Position Reflects Share Issue Proceeds

Adj. EBITDA and Cash Position (DKKm)



- Cash burn from operations and investments reduced to DKKm 17.1 (Q1 2021: DKKm 21.3) reflecting working capital management
- Cash position end-Q1 2022 of DKKm 27 strengthened with DKKm 93.1 net proceeds from share offering closed April 1, 2022

* Pro forma DKKm 93.1 net proceeds from share issue that closed April 1, 2022



Fully Subscribed Rights Issue

- 66.9 million new shares at a price of DKK 1.50 per share
- Advance subscription commitments and guarantees from institutional investors and three largest shareholders
- Proceeds + existing cash will be used to strengthen BioPorto's capital resources and advance implementation of its strategic priorities, including:
 - the clinical trial and application to the FDA for approval of The NGAL Test for assessment of AKI in children under the age of 22 (pediatrics) in the U.S.
 - operational and quality improvements to prepare for future scale and IVDR implementation
 - general corporate purposes
 - investments in U.S. organization and marketing to prepare for The NGAL Test launch

Financial Guidance 2022





Guidance for 2022: On-track for Execution on Both Strategic & Financial Targets

Revenue	Approx. DKK 24-26 million	EBIT loss	Approx. DKK 95-100 million	Adj. EBITDA loss	Approx. DKK 76-81 million
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BioPorto's performance and guidance for 2022 is based on certain assumptions described in the annual report and continues to be subject to uncertainty due to COVID-19, including continued opening of societies and the normalization of access to hospitals, research laboratories, and regulatory bodies.

Financial Calendar 2022

August 17, 2022 Q2 2022 Results

November 9, 2022 Q3 2022 Results

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