

Annual report 2021 BioPorto A/S

April 6, 2022







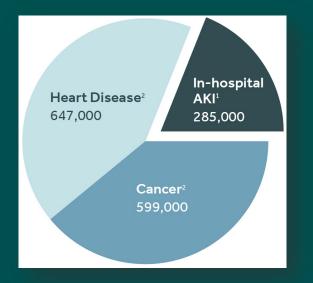
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AKI is the 3rd Leading Cause of Death Annually



Patients at Risk



CARDIAC SURGERY ⁴	
30%	
SEPSIS⁵	
	50%
NEPHROTOXIC MEDICATION ⁶	
14-26%	
MECHANICAL VENTILATION ⁷	
29%	
TRANSPLANT ^{8,9}	
KIDNEY – 37%	STEM CELL – 36%

230%

INCREASE

in AKI hospitalizations

in the US (2000-2014)³

REFERENCES: (1) Brown JR, *BioMed Research International.* 2016;ID4278579. (2) CDC, FastStats: Deaths and Mortality. 2017. (3) Pavkov ME. *MMWR Morb Mortal Wkly Rep.* 2018;67. (4) O'Neal JB, *Crit Care.* 2016;20(1). (5) Alobaidi R, Semin *Nephrol.* 2015;35(1). (6) Perazella MA, *CJASN.* 2018;13. (7) Lombardi R, *CJASN.* 2011;6(7). (8) Alkandari O, *CJASN.* 2018;13(11). (9) Hingorani SR, *Kidney Int.* 2005 Jan;67(1). (10) Susantitaphong P, *CJASN.* 2014;9(6). (11) Kaddourah A, *New England Journal of Medicine* 2017;376.

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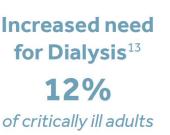
Clinical Burden

IN-HOSPITAL



(
	+	

Increased Length of Stay¹² **7-23 days**



Increased overall Mortality Rate¹³ **25%**

3 YEARS POST-DISCHARGE

- 30% readmission rate¹⁴
- 38% increase in a major cardiac event¹⁵
- 59% of AKI survivors have
 1 or more kidney abnormalities: microalbuminuria, hyperfiltration, decreased GFR, hypertension¹⁶
- Up to 25% progress to CKD¹⁷

Economic Burden AKI costs the US healthcare system \$5.4 - \$24.0 billion* annually¹⁸

REFERENCES: (12) Sutherland SM, CJASN. 2013;8(10). (13) Hoste EA, Intensive Care Med. 2015;41(8). (14) Hessey E, CJASN. 2018;13(5). (15) Odutayo A, JASN. 2016;28. (16) Askenazi DJ, Kidney Int. 2006;69(1). (17) Horne KL, BMJ Open. 2017;7(3). (18) Silver SA, Nephron. 2017;137. (19) Alobaidi R, Semin Nephrol. 2015;35(1). (20) Alshaikh HN, Ann Thorac Surg. 2018;105. *Range for adjusted vs. unadjusted costs (adjusted for demographic factors, hospital differences, comorbidities and procedures).



TOTAL GLOBAL ADDRESSABLE MARKET: ~\$3 BILLION

Estimated Global NGAL Market Opportunity

Based on estimates of NGAL use in these clinical settings:

- ICUs
- Emergency Departments
- Certain outpatient settings





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 costeffectively build the innovation pipeline

- 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



Tracking towards a US Breakthrough

- Global COVID-19 pandemic affected execution of clinical trial
- Promising data from interim analysis of The NGAL Test for pediatrics
- New management team introduced in October 2021
- Revenue growth driven by RUO sales of The NGAL Test in the US and by strong performance in antibody portfolio
- Developed focused strategy on 2022 submission of the FDA De Novo application
- Fully subscribed rights offering in March 2022 strengthened BioPorto's financial position to support activities into first half 2023



2021 Financial highlights

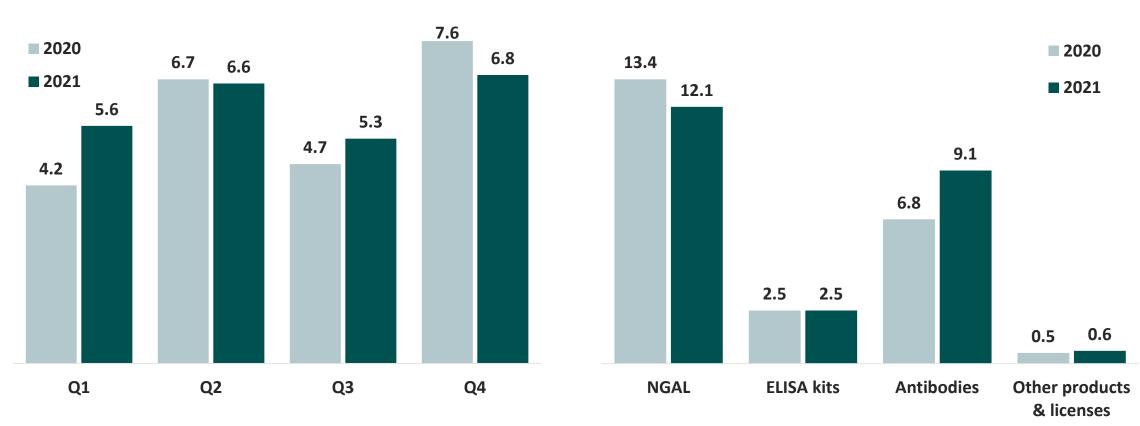


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Revenue growth in 2021 driven by strong performance in antibodies and higher RUO sales of the NGAL Test in the US

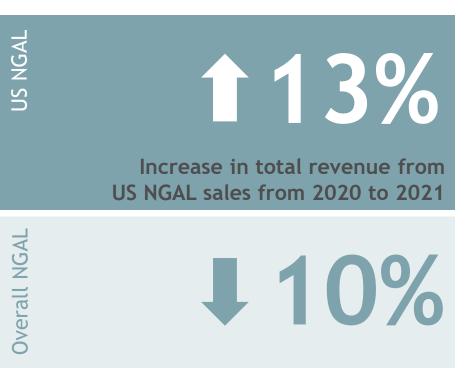


Revenue by Quarter (DKKm)

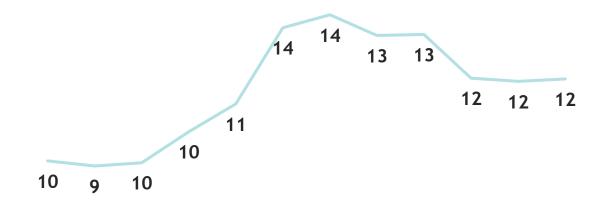
Revenue by Product Category (DKKm)

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2021 NGAL Product sales affected by shift of focus to US



Development in global sales of NGAL from 2020 to 2021 NGAL Product sales by Quarter (LTM, DKKm)

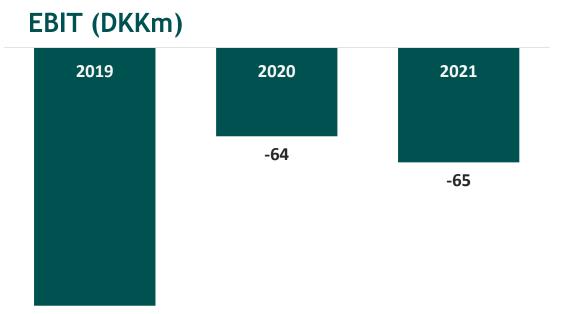








EBIT loss below expectations as commercial activities and clinical trials were postponed due to COVID-19

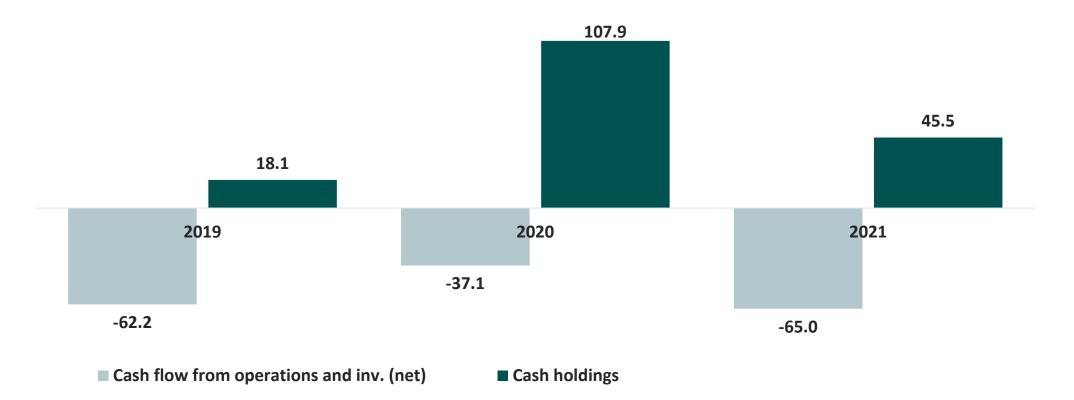


• 2021 EBIT was favorably impacted on a non-cash basis by approx. DKK 4 million for the forfeiture of warrants from former employees and management



Cash position of DKK 45 million end-2021

Cash flows and cash holdings (DKKm)



Fully subscribed DKK 100.4 million rights issue in March 2022



- 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

Future Capital Market Intentions



An Approach to Access US Markets: Funding Onward Growth & Expansion

	2022-Q2		2022-Q3	2022-Q4	2022-Q4 2023			
Financing				Targeted Cross-Bo Offering Including t			U.S. Nasdaq Listing	
Pediatric NGAL	Finalize Clinical Trial Data Collection	Compile Results	FDA Review of De Novo Application (150 days + any inquiries) ¹			Cor	Commercialization	
Expand NGAL Population & Indications ²			Develop Stu	dy Protocol, Sites & Contracts	Prepare Pre-sub	FDA Reviews Pre-sub	Clinical Trial & Analytical Testing	

- Our objective includes building a dynamic presence in the US... the world's largest and leading IVD market
- To be Defined: capital structure, exchange listing(s), legal entities, etc.

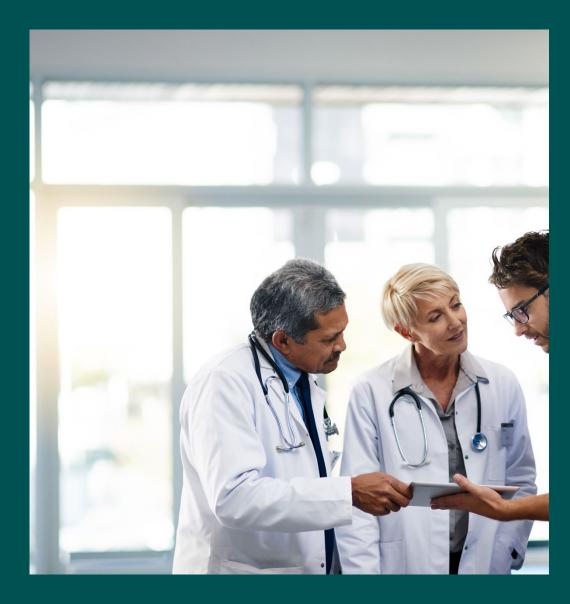
• Timing and decision-making depend on a range of factors, including in particular the timing of the potential FDA De Novo application, pre-submissions and related reviews

¹FDA 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"

²Includes adult population, expanded claims, other NGAL indications, other lab instruments, etc.

2022 Outlook & Strategy







Financial Outlook for 2022



BioPorto's performance and outlook 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.

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



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Financial Calendar 2022

April 28, 2022 May 11, 2022 August 17, 2022 November 9, 2022

Annual General Meeting Q1 2022 Results Q2 2022 Results Q3 2022 Results

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