BIOARCTIC AB (PUBL) NASDAQ STOCKHOLM: BIOA B

Full Year Report January-December 2020 Stockholm, February 4, 2021

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BioArctic – a unique Swedish biopharma company Improving life for patients with central nervous system disorders



High unmet need for disease-modifying treatments for Alzheimer's and Parkinson's diseases creates **large commercial opportunity**



World-class research and development driven organization with basis in founder's breakthrough discoveries and fruitful collaborations with leading academic researchers and pharma companies generating and developing innovative projects



Attractive and well-balanced project portfolio with projects from discovery through Phase 3 and combination of both proprietary projects with substantial marketing and out-licensing potential and partnered projects generating income



Well-financed with approximately BSEK 1 (MUSD >120¹) in cash, **net profitable** during the last seven years and **valuable collaboration agreements** totaling BSEK 8.7² (BUSD ~1) plus royalties



¹⁾ FX as per December 31, 2020

²⁾ FX as per December 31, 2020

2020 in summary

Lecanemab (BAN2401)

- Phase 2b open-label extension study in early AD:
 - Data presented at several congresses supports the encouraging Phase 2b data
- Phase 3 confirmatory study Clarity AD in early AD:
 - Progressed well and Eisai mitigated potential impact of COVID-19
- Phase 3 AHEAD 3-45 in preclinical (asymptomatic) AD:
 - Started in the US

ABBV-0805

 AbbVie decided to cancel recruitment to Phase 1 MAD study, and instead develop plan to move the project into Phase 2

Other projects

- Project portfolio developed well despite COVID-19 pandemic
- The two new innovative projects in our early portfolio now well underway

General

 Won the Allbright award for the company's leadership and focused work with gender equality





Q4 highlights New data further supports lecanemab (BAN2401)

Lecanemab (BAN2401)

- Lecanemab was assigned as the International Nonproprietary Name (INN) for BAN2401
- Presentations at CTAD congress :
 - Clarity AD Phase 3:
 - Baseline characteristics consistent with Phase 2b, and representative of an early AD population
 - Phase 2b open label extension study
 - Rapid reduction of amyloid in the brain (already at 3 months)
 - Consistent low level of adverse event ARIA-E (<10%)
 - AHEAD 3-45, Phase 3 in preclinical (asymptomatic) AD:
 - Study design and initial screening results

Events after the period

- European patent granted for new antibodies targeting truncated forms of amyloid beta
- Eisai expanded Clarity AD by approximately 200 patients to ensure robust dataset, Eisai still expects 18-month readout in September 2022





Attractive and well-balanced project portfolio combines fullyfinanced partner projects and cutting-edge proprietary projects

	Project	Partner	Discovery	Preclinical	Phase 1	Phase 2	Phase 3
ALZHEIMER'S DISEASE	Lecanemeb (BAN2401) (Clarity AD)	Eisai ¹	Early Alzheimer's disease ⁴				
	Lecanemab (BAN2401) (AHEAD 3-45)	Eisai ¹	Preclinical (asymptomatic) Alzheimer's disease ⁵				
	BAN2401 back-up	Eisai					
	AD1801						
	AD1502						
	AD1503						
	AD2603						
PARKINSON'S DISEASE	ABBV-0805 ²	AbbVie					
	PD1601	AbbVie					
	PD1602	AbbVie					
OTHER CNS DISORDERS	Lecanemab (BAN2401)		Down's syndrome Traumatic brain i	e ³ Injury ³			
	ND3014						
BLOOD-BRAIN BARRIER TECHNOLOGY	BBB technology platform						
DIAGNOSTICS	Imaging and biochemical biomarkers – Alzheimer's disease						
	Imaging and biochemical biomarkers – Parkinson's disease	AbbVie		\			

as of December 31, 2020



¹⁾ Partnered with Eisai for lecanemab (BAN2401) for treatment of Alzheimer's disease. Eisai entered partnership with Biogen regarding lecanemab (BAN2401) in 2014

²⁾ AbbVie in-licensed BAN0805 in late 2018 and develops the antibody with the designation ABBV-0805

³⁾ Dementia and cognitive impairment associated with Down's syndrome and with traumatic brain injury

⁴⁾ Mild cognitive impairment due to Alzheimer's disease and mild Alzheimer's disease

⁵⁾ Normal cognitive function with intermediate or elevated levels of amyloid in the brain

Long-standing and extensive partnerships

Alzheimer's disease

Partner track record

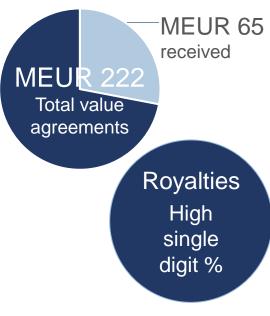


Discovered and developed world's best-selling medicine for symptoms in Alzheimer's



Industry-leading pipeline in dementia area

Collaboration and license



BioArctic retains rights to lecanemab in other indications and option to market in the Nordics

Parkinson's disease

Partner track record

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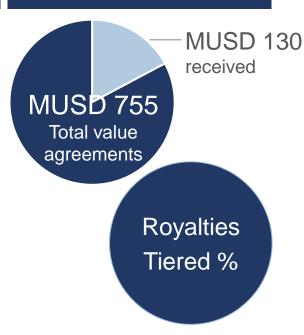
World's all-time best-selling medicine (BUSD 20)



Approved product for symptoms associated with Parkinson's disease



Collaboration and license

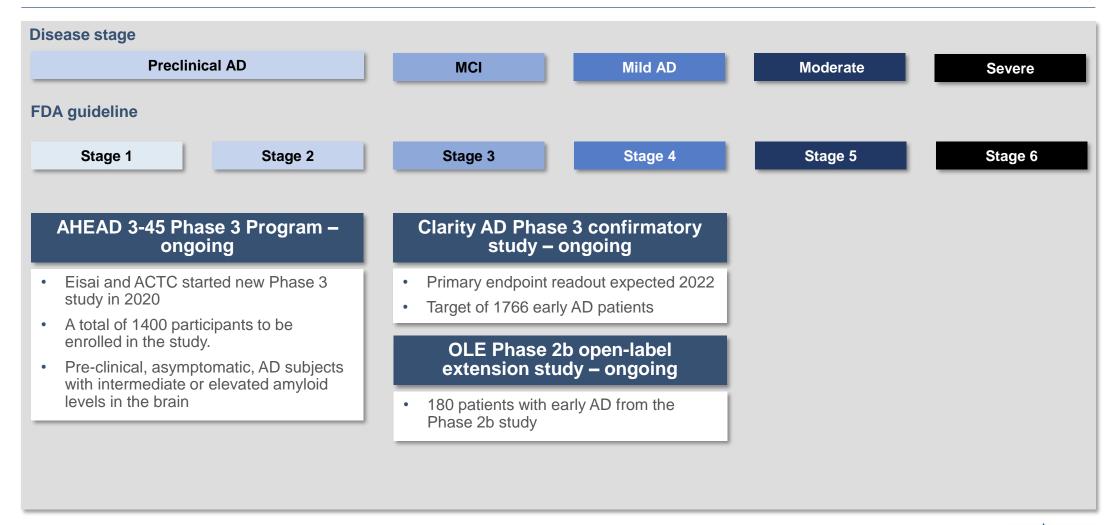


 AbbVie global rights to alphasynuclein portfolio for all indications

Sources: Eisai, AbbVie and BioArctic corporate information



Broad lecanemab clinical program – driven by BioArctic's partner Eisai





Early-stage portfolio continues to develop well



Discovery stage programs

- 4 fully-owned disease modifying antibody projects in Alzheimer's disease
- Each project has a different mechanism from the others

Parkinson's disease

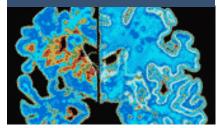


Discovery stage projects

 Preclinical stage alpha-synuclein projects in research collaboration with



Other CNS disorders



Neurodegeneration research

- Lecanemab in indications other than Alzheimer's disease
- Research project in neurodegeneration ("ND") with potential in various CNS disorders

Blood-brain barrier technology



Blood-brain barrier technology platform

- Continued development with expanded and enhanced capabilities
- Collaboration with Uppsala University under Vinnova grant

Diagnostics



Diagnostics

 Continued development of imaging and biochemical biomarkers







Net revenues and operating profit/loss Q4 2020







- Net revenues were 8 MSEK (26) for the fourth quarter
- The change is mainly related to lower activity for BioArctic in the Parkinson's program, according to plan
- Lower currency exchange losses, travel costs and costs for consultants

■ Other external ■ Personnel ■ Depreciations ■ Other operating

- Project expenses in total decreased to 13 MSEK (21) in the fourth quarter, while project expenses in our own projects increased in Q4 compared to the previous year due to higher activity
- Operating loss was -30 MSEK (-21) for the fourth quarter
- The change is mainly attributable to lower activity for BioArctic in the Parkinson's program, according to plan

Operating expenses are expected to be in the range of 180 - 220 MSEK for the financial year January - December 2021

2020

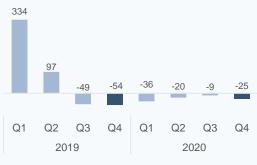


Cash and net result Q4 2020



• Cash balance amounted to 1,000 MSEK (1,113) at the end of the

Cash Flow From Operating Activities (MSEK)



 Operating cash flow amounted to -25 MSEK (-54) during Q4

Net Result (MSEK)



- Net result for the period was -13 MSEK (-17)
- The change is mainly attributable to lower activity for BioArctic in the Parkinson's program, according to plan

In summary, BioArctic continues to have a strong financial position

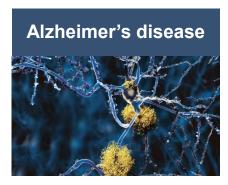


fourth quarter





Upcoming news flow



Lecanemab (Eisai)

- To present data at international congresses
- Phase 3 confirmatory study in early AD results 2022
- Phase 2b open label extension study results
- Phase 3 study in preclinical asymptomatic AD

Discovery stage programs

 Advance into preclinical development

Parkinson's disease



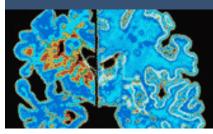
ABBV-0805 (AbbVie)

 Complete Phase 1 and start Phase 2

Discovery stage projects

• Development in AbbVie collaboration

Other CNS disorders



Neurodegeneration research

- New project development
- New indications and new targets

Blood-brain barrier technology



Blood-brain barrier technology platform

 Continue development of platform

Diagnostics

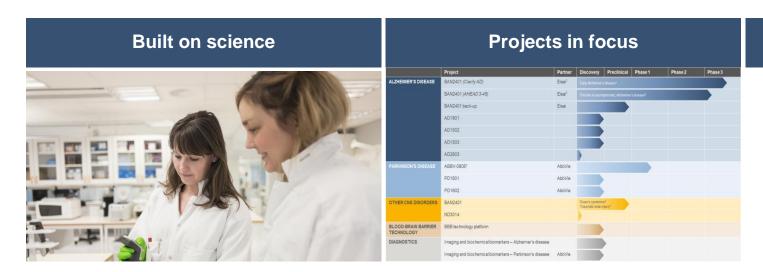


Diagnostics

 Continue development of imaging and biochemical biomarkers



BioArctic: With Patients in Mind



Great science Great projects

Value-driven leadership



Great people









NEXT REPORT & IR CONTACT

- **Next Report:** Q1 2021 Jan-Mar 2021 on April 21, 2021
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