BIOARCTIC AB (PUBL) NASDAQ STOCKHOLM: BIOA B

First Quarter Report January-March 2020 Stockholm, April 22, 2020

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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 BSEK >1 (MUSD >100) in cash, **net profitable** during the last seven years and **valuable collaboration agreements** totaling BSEK 9.6 (BUSD ~1) plus royalties



Operations progressing according to plan



BAN2401

- 3 clinical studies underway or in planning
- Including Phase 3
 confirmatory study
 ("Clarity AD")
 underway

 Eisai

Discovery stage programs

 4 fully-owned disease modifying programs

Parkinson's disease



ABBV-0805

 Second stage of Phase 1 study in patients with Parkinson's disease now underway by partner

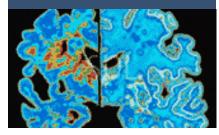


Discovery stage projects

 Progressing according to plan in research collaboration with



Other CNS disorders



Neurodegeneration research

- Working on new indications and new targets
- New research program
 potential in various
 neurodegenerative
 disorders

Blood-brain barrier technology



Blood-brain barrier technology platform

Continuing development

Diagnostic tools



Diagnostic tools

 Continuing development of imaging and biochemical biomarkers



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	BAN2401	Eisai, Biogen ¹					
	BAN2401 back-up	Eisai					
	AD1801						
	AD1502			•			
	AD1503			•			
	AD2603						
PARKINSON'S DISEASE	ABBV-0805 ²	AbbVie					
	PD1601	AbbVie					
	PD1602	AbbVie					
OTHER CNS DISORDERS	BAN2401 Down's syndrome ³ Traumatic brain injury						
	ND3014						
BLOOD-BRAIN BARRIER TECHNOLOGY	BBB technology platform						
DIAGNOSTIC TOOLS	Imaging and biochemical biomarkers – Alzheimer's disease						
	Imaging and biochemical biomarkers – Parkinson's disease	AbbVie					

as of March 31, 2020

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

Long-standing and extensive partnerships

Alzheimer's disease

Partner track record

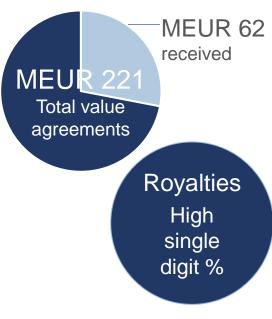


Discovered and developed world's best-selling medicine for symptoms in Alzheimer's



Industry-leading pipeline in dementia area

Collaboration and license



 BioArctic retains rights to BAN2401 in other indications and option to market in the Nordics

Parkinson's disease

Partner track record

abbvie

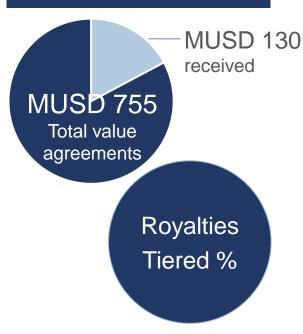
World's all-time best-selling medicine (BUSD 20)



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



BAN2401: potential disease modifying antibody for Alzheimer's disease with positive Phase 2b results now in Phase 3

No existing disease-modifying treatment TODAY >30 million people with Alzheimer's

BAN2401 has positive Phase 2b results

- Large trial: 856 early Alzheimer's patients
- Consistent effects on clinical outcomes, imaging and neurodegenerative biomarkers
- Effect increase over time
- Good safety profile

BAN2401 unique profile

Unique and targeted binding profile

 Highly selective for toxic forms of misfolded Abeta (oligomers/protofibrils)

Unique clinical fingerprint

- Rapid onset of clinical effect
- Consistent effects
- No titration required due to low frequency of ARIA–E

Broad clinical program

- **Confirmatory Phase 3 study** ("Clarity AD") ongoing
 - Primary endpoint final readout expected 2022
- Phase 2b open label extension study ongoing
- Phase 3 prevention program ("AHEAD 3-45" comprised of 2 groups "A3" and "A45") in even earlier stages of AD
 - Preparing to start 2020

Broad BAN2401 clinical program

Clarity AD Phase 3 confirmatory study

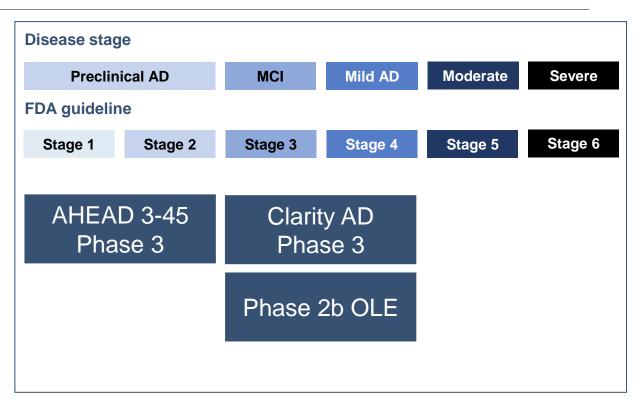
- Started May 2019, MEUR 15 milestone to BioArctic
- Study ongoing with target of 1566 early AD patients
- Primary endpoint readout expected 2022

Phase 2b OLE open-label extension study

- Ongoing with approx. 200 patients with early AD
- Baseline data presented at CTAD December 2019 showed maintenance of benefit after BAN2401 treatment conclusion

AHEAD 3-45 Program

- Eisai and ACTC planning to start A3 and A45 prevention study in 2020
- A45: ~1000 subjects with preclinical AD, little to no cognitive impairment and elevated levels of amyloid in the brain
- A3: ~400 subjects with early preclinical AD, cognitively normal with intermediate amyloid levels in the brain
- Biomarkers on amyloid, tau and neurodegeneration
- Clinical evaluation scale PACC5¹ for A45



Clinical program driven by:





ABBV-0805: potential disease modifying antibody for Parkinson's disease with strong preclinical results now in Phase 1

High unmet medical need

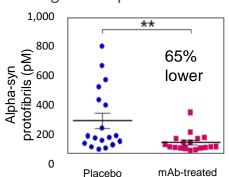
No existing disease-modifying treatment

- 2nd most common neurodegenerative disease
- 6.2 million people with Parkinson's¹
- Younger patient group, still at working age

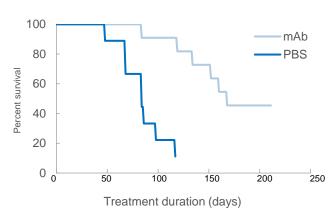
1) Dorsey and Bloem, JAMA Neurology 2018;75:9-10

Preclinical proof of concept

Reduction of neurotoxic alpha-synuclein oligomers/protofibrils



Delays disease progression and increases lifespan



Unique profile

Unique and targeted binding profile

 Highly selective for toxic forms of misfolded alpha-synuclein (oligomers/protofibrils)

Built on genetic and pathology rationale

- Alpha-synuclein mutations lead to Parkinson's
- Alpha-synuclein oligomers/protofibrils are elevated in Parkinson's

ABBV-0805 in clinical development

- Phase 1 with ABBV-0805 ongoing by AbbVie
- BioArctic delivers follow-up antibodies in the continued collaboration with AbbVie



Continued progress in collaboration with AbbVie on alpha-synuclein

Collaboration highlights

- ABBV-0805 targeting disease modification in Parkinson's disease
- Potential to expand to earlier stage Parkinson's disease patients and other diseases where alphasynuclein plays a role
- AbbVie is responsible for clinical development
- BioArctic delivers followup antibodies in the continued collaboration with AbbVie

ABBV-0805 advancing in clinical trials

December 2018

January 2019 February 2019 March 2019 2020

Alpha-synuclein antibody portfolio licensed by AbbVie

Milestone of 50 MUSD for the license ABBV-0805 INDapplication approved by the US FDA AbbVie started Phase 1 with ABBV-0805

Phase 1 study ongoing

- Aim to evaluate safety and tolerability
- Second stage of Phase 1 study in patients with Parkinson's disease now underway



Drug Discovery Today



feature

How partnership should work to bring innovative medicines to patients

Janice M. Twombly¹, jan@rhythmofbusiness.com, Johanna Fälting², Marco Giorgetti³, Anna C. Maroney³ and Gunilla Osswald²

Scientists increasingly find themselves working in bilateral drug development alliances. Alliances are conceptually simple, but operationally challenging, resulting in the value-eroding misalignment and delays that alliances often experience. This case study of an exemplary collaboration between a small biotech and a global biopharmaceutical company is based on 15 interviews and a lessons-learned workshop conducted with the principal alliance team members. We outline five repeatable practices identified as contributing to their success that other alliance teams can follow.



Early stage portfolio continues to develop well



Discovery stage programs

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

Parkinson's disease

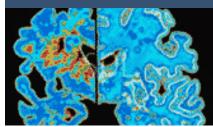


Discovery stage projects

 Preclinical stage alpha-synuclein projects in research collaboration with



Other CNS disorders



Neurodegeneration research

- BAN2401 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

Diagnostic tools



Diagnostic tools

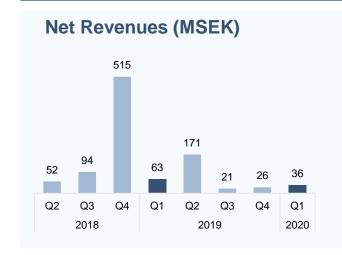
 Continued development of imaging and biochemical biomarkers

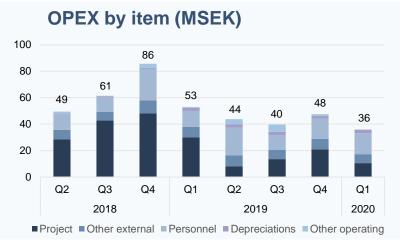






Net revenues and operating profit Q1 2020







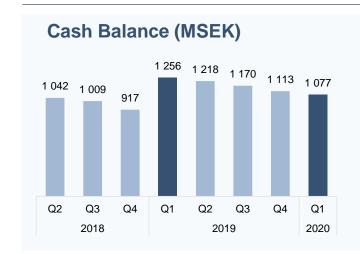
- Net revenues were 36.4 MSEK (63.4) for the first quarter
- The change is primarily due to lower activity in the Parkinson's program as planned
- Project expenses decreased to 10.5 MSEK (30.0) due to lower external expenses in the Parkinson's project versus previous year
- The personnel expenses increased to 16.0 MSEK (12.0) primarily attributable to an increase in the number of employees

- Operating profit was 3.8 MSEK (17.3) for the first quarter
- The change is attributable to lower activity in the Parkinson's program as planned

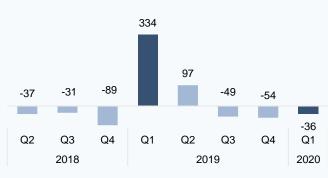
Operating expenses are expected to be in the range of 180 - 230 MSEK for the financial year January - December 2020



Cash related and net profit Q1 2020











 Cash balance amounted to 1,077.3 MSEK (1,255.6) at the end of the first quarter Operating cash flow amounted to -36.3 MSEK (333.6) during Q1 Net result for the period was 3.6 MSEK (13.6)

To sum up, BioArctic showed positive net result for the first quarter and continues to have a strong financial position







Upcoming news flow



BAN2401 (Eisai)

- To present data at international congresses
- Phase 3 confirmatory study results 2022
- Phase 2b open label extension study results
- Phase 3 prevention study to start 2020

Discovery stage programs

 Advance into preclinical development

Parkinson's disease



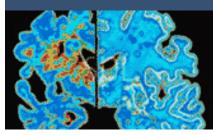
ABBV-0805 (AbbVie)

- Complete Phase 1 and start Phase 2
- Publications on the collaboration

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

Diagnostic tools



Diagnostic tools

 Continue development of imaging and biochemical biomarkers



BioArctic: With Patients in Mind

Built on science



Projects in focus



Value-driven leadership





GUNILLA OSSWALD, CEO



NEXT REPORT & IR CONTACT

- Next Report: Interim report Jan-Jun 2020 on July 10, 2020
- Contact: +46 8 695 69 30 <u>ir@bioarctic.se</u>

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