

BIOARCTIC AB (PUBL)
NASDAQ STOCKHOLM: BIOA B

First Quarter Report January-March 2021

Stockholm, April 21, 2021

Gunilla Osswald, PhD, CEO
Jan Mattsson, CFO

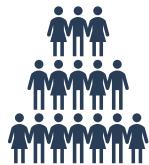


Disclaimer

- This presentation has been prepared and produced by BioArctic AB (publ) (“BioArctic”) solely for the benefit of investment analysis of BioArctic and may not be used for any other purpose. Unless otherwise stated, BioArctic is the source for all data contained in this presentation. Such data is provided as at the date of this presentation and is subject to change without notice.
- This presentation includes forward-looking statements. These forward-looking statements involve known and unknown risks, uncertainties and other factors, which may cause BioArctic’s actual results, performance, achievements or industry results to be materially different from those expressed or implied by these forward-looking statements. Forward-looking statements speak only as of the date of this presentation and BioArctic expressly disclaims any obligation or undertaking to release any update of, or revisions to, any forward-looking statement in this presentation, as a result of any change in BioArctic’s expectations or any change in events, conditions or circumstances on which these forward-looking statements are based.
- This presentation does not constitute or form part of, and should not be construed as, an offer or invitation for the sale of or the subscription of, or a solicitation of any offer to buy or subscribe for, any securities, nor shall it or any part of it or the fact of its distribution form, or be relied on in connection with, any offer, contract, commitment or investment decision relating thereto, nor does it constitute a recommendation regarding the securities of BioArctic.
- The information in this presentation has not been independently verified.
- No regulatory body in Sweden or elsewhere has examined, approved or registered this presentation.

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



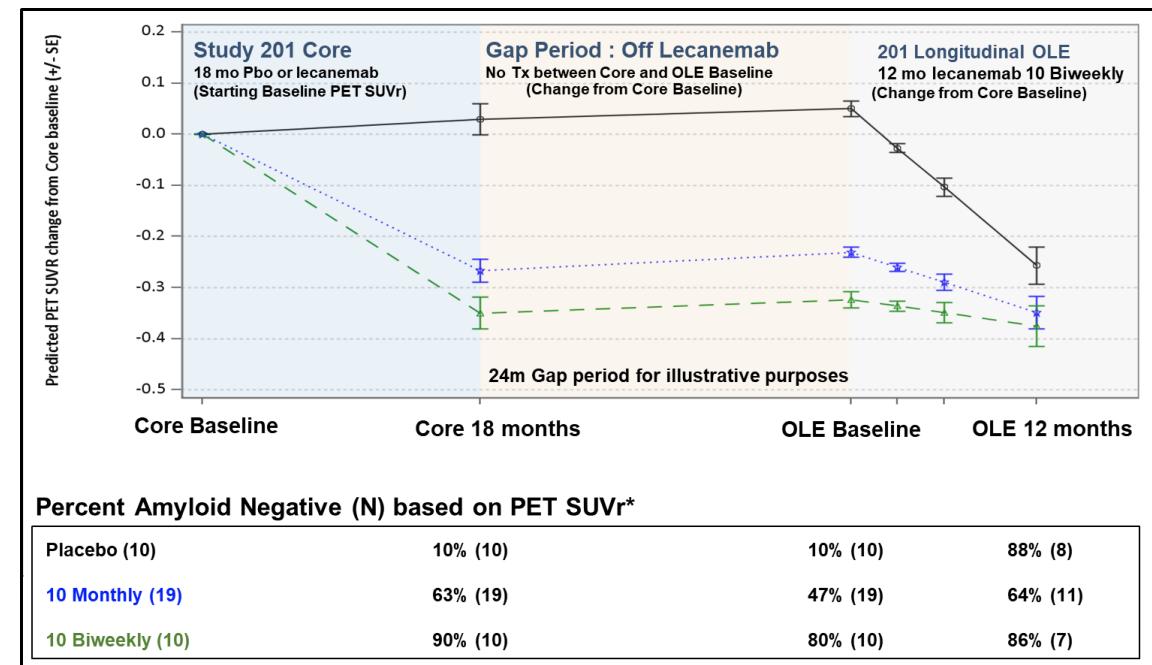
Well-financed with approximately BSEK 1 (MUSD >110¹) in cash, **net profitable** during seven of the last eight years and **valuable collaboration agreements** totaling BSEK 8.9² (BUSD ~1) plus royalties

Q1 highlights and recent news

New data continues to strengthen lecanemab (BAN2401)

Lecanemab

- Eisai expanded Clarity AD to ensure robust dataset.
Enrollment was completed with 1795 early AD patients.
Eisai still expects 18-month readout in September 2022
- Phase 2b manuscript published in Alzheimer's Research & Therapy on April 19
- BioArctic presented data at AD/PD suggesting that lecanemab could help preserve brain function in adults with Down's syndrome with dementia
- Eisai presented updated results at AD/PD from the Phase 2b open label extension study (OLE);
 - Effects of lecanemab on amyloid reduction persist for up to 2 years following lecanemab discontinuation
 - Lecanemab 10 mg/kg biweekly rapidly reduced brain amyloid in Core placebo-treated subjects as early as 3 months in OLE, with continued reduction over 12 months of treatment
 - Lecanemab, dosed at 10 mg/kg biweekly, was shown to reduce brain amyloid to levels below Alzheimer's disease pathology in more than 80 percent of patients who participated in the core and OLE, as early as 12 months into treatment.
- Three different anti-amyloid antibodies have now shown clinical effect and reduction of amyloid in the brain of Alzheimer's disease patients



Q1 highlights (continued...)

ABBV-0805

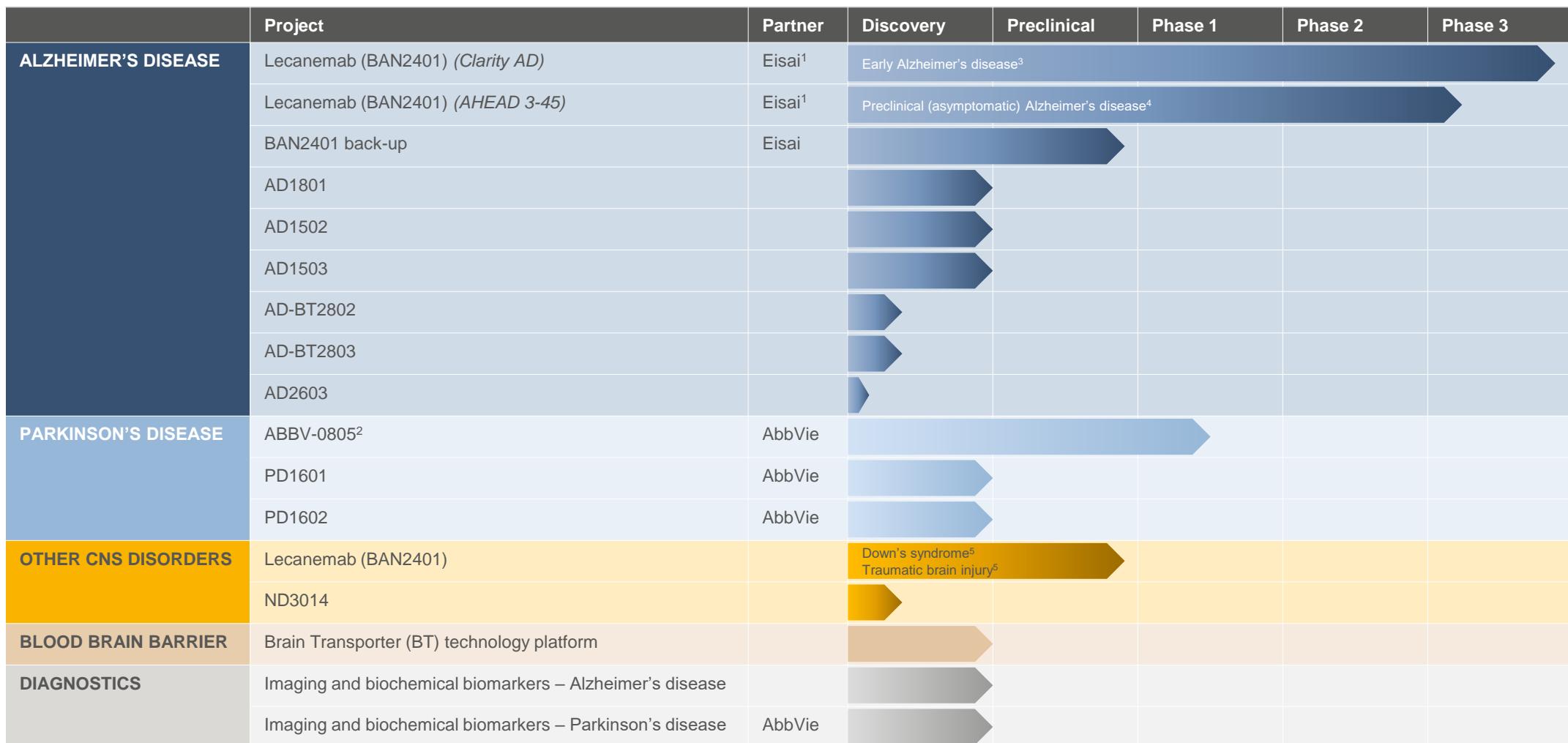
- Important learnings at AD/PD conference in the Parkinson's disease field supporting the design of the Phase 2 program

Other

- European patent granted for new antibodies targeting truncated forms of amyloid beta
- Patent portfolio expanded and now includes more than 230 granted patents and 60 pending patent applications within 13 patent families
- Project portfolio expanded with two new Alzheimer's disease projects coupled with our Brain Transporter technology
- Presented BioArctic and our important research in Alzheimer's disease to the Swedish royal family



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



as of March 31, 2021

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

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

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

Long-standing and extensive partnerships

Alzheimer's disease

Partner track record

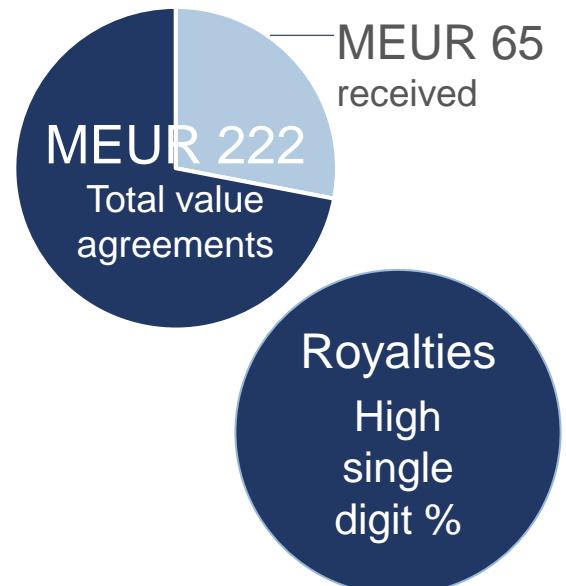


Discovered and developed world's best-selling medicine for symptoms in Alzheimer's



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



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

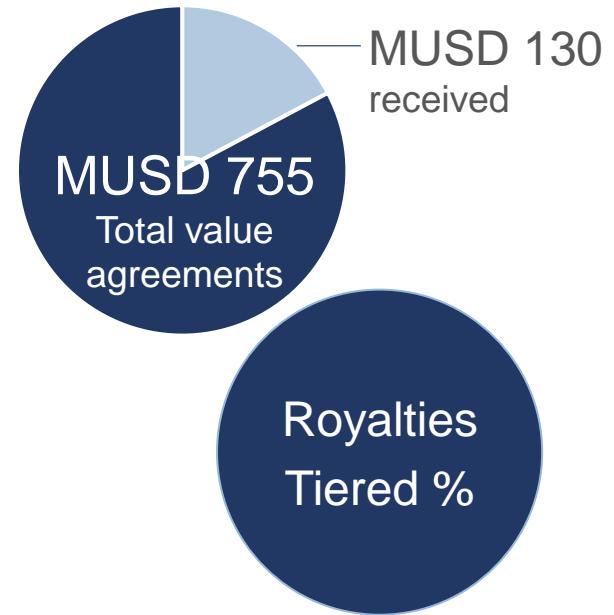


10 different indications in immunology

Approved product for symptoms associated with Parkinson's disease



Collaboration and license



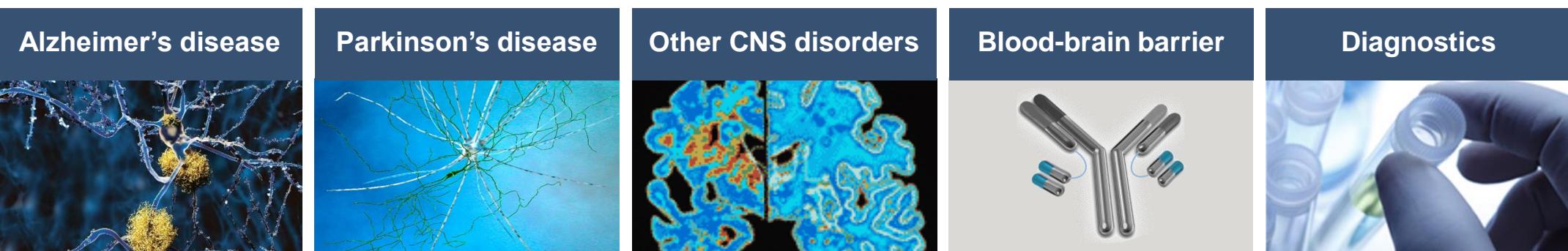
- AbbVie global rights to alpha-synuclein portfolio for all indications

Sources: Eisai, AbbVie and BioArctic corporate information

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

Disease stage	Preclinical AD	MCI	Mild AD	Moderate	Severe	
FDA guideline	Stage 1	Stage 2	Stage 3	Stage 4	Stage 5	Stage 6
AHEAD 3-45 Phase 3 Program – ongoing		Clarity AD Phase 3 confirmatory study – ongoing				
<ul style="list-style-type: none">• Eisai and ACTC started new Phase 3 study in 2020• A total of 1,400 participants to be enrolled in the study.• Pre-clinical, asymptomatic, AD subjects with intermediate or elevated amyloid levels in the brain		<ul style="list-style-type: none">• Primary endpoint readout expected 2022• 1,795 early AD patients				
OLE Phase 2b open-label extension study – ongoing		<ul style="list-style-type: none">• 180 patients with early AD from the Phase 2b study				

Early-stage portfolio continues to develop well



Alzheimer's disease

Discovery stage programs

- Expanded early-stage portfolio with 2 new AD+BT projects
- 6 fully-owned disease modifying antibody projects in Alzheimer's disease
- BAN2401 backup in collaboration with Eisai

Parkinson's disease

Discovery stage projects

- Preclinical stage alpha-synuclein projects in research collaboration with

abbvie

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

Brain Transporter (BT)

- Continued development of our Brain Transporter (BT) technology platform
- Collaboration with Uppsala University under Vinnova grant

Diagnostics

Diagnostics

- Continued development of imaging and biochemical biomarkers



Financial Summary

Net revenues and operating profit/loss Q1 2021



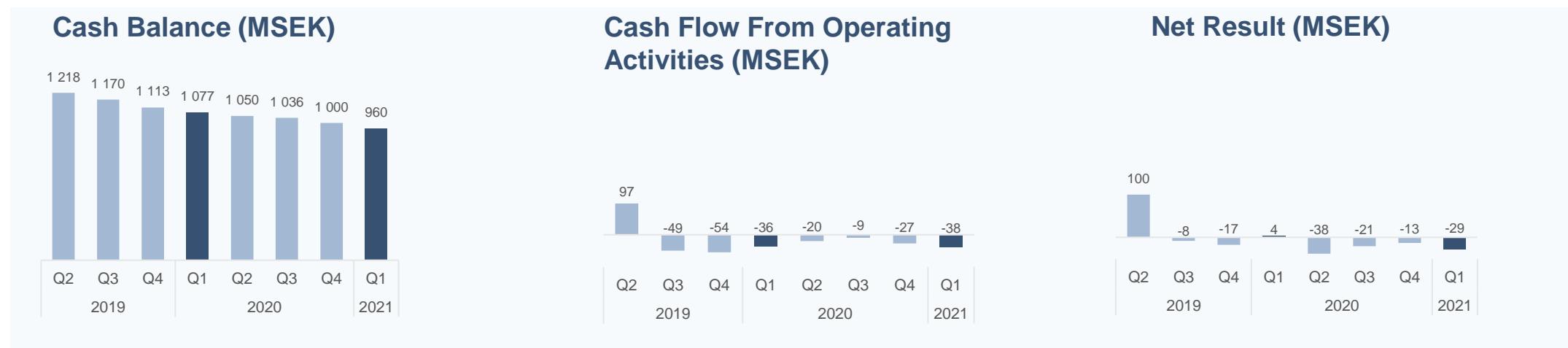
- Net revenues were 7 MSEK (36) for the first quarter
- The change is mainly related to a one-off revenue adjustment of 23 MSEK in Q1 2020, and lower activity for BioArctic in the Parkinson's program according to plan

- Total costs in the quarter were in line with the same period previous year
- Project expenses in total increased to 11 MSEK (10) in the first quarter, mainly explained by increase in our own projects and expanded portfolio

- Operating loss was -29 MSEK (4) for the first quarter

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

Cash and net result Q1 2021



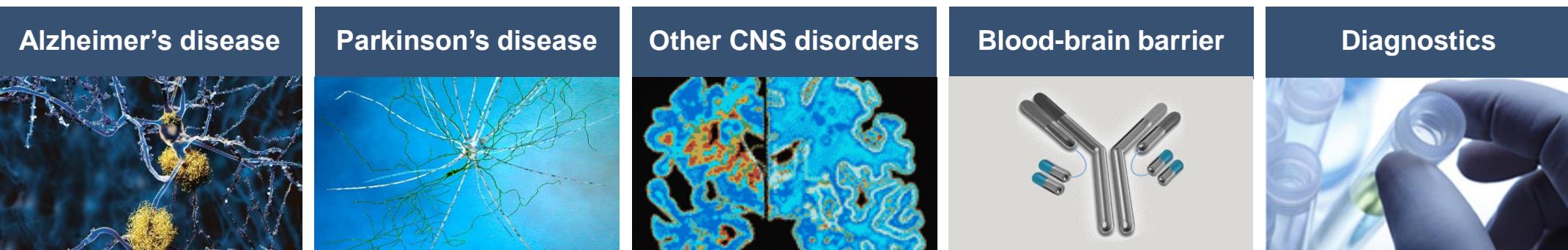
- Cash balance amounted to 960 MSEK at the end of the first quarter
- Operating cash flow amounted to -38 MSEK (-36) during Q1
- Net result for the period was -29 MSEK (4)

In summary, BioArctic continues to have a strong financial position



Upcoming news and closing remarks

Upcoming news flow



Lecanemab (Eisai)

- Data presented 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

ABBV-0805 (AbbVie)

- Complete Phase 1 and start Phase 2

Discovery stage projects

- Development in AbbVie collaboration

Neurodegeneration research

- New project development
- New indications and new targets

Brain Transporter (BT) technology platform

- Continue development of platform

Diagnostics

- Continue development of imaging and biochemical biomarkers

BioArctic: With Patients in Mind

Great science



Great projects



Great partners



Great people



GUNILLA OSSWALD, CEO



JAN MATTSSON, CFO



**OSKAR BOSSON, VP
COMMUNICATIONS & IR**



**NEXT REPORT & IR
CONTACT**

- **Next Report:**
Q2 2021 Jan-Jun 2021 on July 9, 2021
- **Contact:**
Oskar Bosson,
VP Communications & IR
+46 704 10 71 80
ir@bioarctic.se

To subscribe to financial reports/press releases and for more information, please visit www.bioarctic.com