Q4 2021 Financial Report February 24, 2022

Proteins for life

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EXPRES²ION BIOTECHNOLOGIES

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Investment Highlights

Key player in advanced protein sciences, with deep pipeline of novel vaccines addressing highvalue markets



High-potential pipeline of key focus, backed up by Contract Research Organization (CRO) business, that has generated SEK 60 million since IPO in 2016



Vaccine development platform with track record and partner validation. +500 proteins produced while posting +90% success rate



Global vaccine market rapidly growing, from USD 33bn (2019) to USD 187bn (2021), corresponding to 460% growth



ExpreS²ion is advancing towards key catalysts during 2022, further de-risking the company's pipeline. COVID-19 phase II readout in Q1'22, initiation of phase III in H1'22

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ABNCoV2 COVID-19 Vaccine (I)

Phase I/IIa study confirms safety and tolerability and excellent efficacy profile



- Dose response: increased titers with higher vaccine doses up to 25 mg, • reaching a plateau at higher doses
- Up to 12 times higher compared to the levels achieved after COVID-19 ٠ infection - significantly higher than the virus neutralization levels reported for leading mRNA COVID-19 vaccines
- for Comirnaty[™])





No reduction in neutralization capacity against Alpha or Delta A 2.2-fold reduction is seen against Beta (compared to >10-fold reported





ABNCoV2 COVID-19 Vaccine (II) Bavarian Nordic carries on with the Phase II study, and granted DKK 800m funding for Phase III

Phase II study design

- Evaluate ABNCoV2 as a booster vaccine in individuals with existing immunity. The study will also assess neutralizing immune responses against cirulating variants of SARS-CoV2
- Enrolling a total of up to 210 health adults

N = 90

N = 90

N = 30

100 µg

50 µg

100 µg

- Individuals (n=180) with existing immunity against SARS-CoV-2, acquired through previous disease or from prior immunization with approved COVID-19 vaccines (mRNA and Adeno)
- Individuals (n=30) with no prior vaccination or disease.

- First of three groups in the phase II study
- One week post vaccination, a 2-34-fold increase in the levels of • neutralizing antibodies was observed against the original (Wuhan) variant and peaked at two weeks with a 2-40-fold increase depending on the initial antibody levels.



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Phase II

Seropositive

Seronegative

No existing

immunity

infected or fully vaccinated

Previously



Phase II study topline results (1/3 groups)



Results from the two other study groups in the phase II study are expected during the first quarter of 2022

Bavarian Nordic is also preparing for a Phase III study of ABNCoV2, expected to be initiated in the first half of 2022



Partnership with Bavarian Nordic ABNCoV2 is already out-licensed with near-term revenue streams supporting ExpreS²ion

AdaptVac receive from Bavarian Nordic

- EUR 4 million upfront (paid in July 2020) ٠
- Up to EUR 136 million in development and sales milestones
- Single- to double-digit-% royalties of Bavarian revenues





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ExpreS²ion receive from AdaptVac

- 34% ownership of AdaptVac
- Up to EUR 2 million in commercial milestone payments
- Lower double-digit percentage of AdaptVac royalties

ES2B-C001 Overcomes Herceptin Resistance The soft agar human cancer cell growth inhibition assay provides in vitro evidence



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Note that this data was generated for AdaptVac's predecessor vaccine candidate (HER2-VLP very similar to ES2B-C001) Source: Palladini, A. et al. (2018), "Virus-like particle display of HER2 induces potent anti-cancer responses", Oncolmmunology, pub. Vol 7, no 3



Both Herceptin (trastuzumab) and ES2B-COO1 inhibited growth in the trastuzumab-sensitive cells

Only ES2B-COO1 inhibited growth in the trastuzumab-resistant cells; cells were

unresponsive to Herceptin



ES2B-CO01 has demonstrated animal proof-of-concept

Effectively inhibited tumor development



- Two weeks after the inoculation of tumor cells, the first vaccine • administration was given. Repeated every 2nd week during the study
- ES2B-CO01 formulated in an adjuvant effectively inhibited tumor ٠ **development**, whereas the control group progressively expanded with tumor development
- administered
- tumors



Prevented tumor development with 95% efficiency

At mouse age 6-8 weeks, 2 vaccinations with 2 weeks interval were

Two vaccinations prevented tumor development with 95% efficiency as

compared to a control group, where all mice spontaneously developed

Advancing Towards Key Catalysts

		2022		2023
and the second	CORONAVIRUS (ABNCoV2) Solution (Q3'21)	BN Phase II BN Phase III study readout study initiatio H1 2022 H1 2022	BN Phase III on initial readout H2 2022	BN ready for market launch (subject to
Here to	BREAST CANCER (ES2B-COC)1)		regulatory approval)
	 Executed in-licensing (Feb 2021) Preclinical animal studies initiated (Q2) 	Preclinical animalGNproof-of-conceptmaresults H1 2022ba	1P anufacturing tch	Preclinical safety studies readout
	INFLUENZA			
	Advance/support further development of one or more candidates in 2022			cGMP/Preclinical safety studies initiation 2023
	MALARIA			
~ V \	 Phase IIa results from the ORH5 A Rh5 vaccine published in mala 2021 launce alterrest 	Additional phase I study in a ria endemic region in Africa thed during 2021, with native adjuvant	Pfs 48/45 phase I study initiation 2022	RH5-VLP phase initiation 2023

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	2024			
)				
Filing of clinical study application H2 2023	Initiation of first human clinical study 2024	Outlicensing window opens pending human data		
e I RH5 phase I study readout H2 2023				

Management Team

Experienced team driving pipeline-focused business



- MSc. In Finance/Strategic Management, Copenhagen Business School, Denmark
- >25 years industry finance, business dev and management experience



accenture





Keith Alexander, CFO

- MBA, The Wharton School and the University of Pennsylvania, USA
- >20 years of equity research, corporate strategy, asset management and consulting experience

Danske Bank

J.P.Morgan





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ASTION

Dr. Mette Thorn, VP Preclinical Development

• PhD in Immunology, and a MSc in Chem Eng., Tech. Univ of Denmark • 20 years industrial research experience

Dr. Mattis F. Ranthe, CMO

- PhD in Cardiovascular Epidemiology (2013), and a Medical
- Diploma (MD) (2006) from the University of Copenhagen,
- 10 years of broad clinical experience

Q4 2021 Financial Results



Operating income



SEK `000s	2021	
4Q	4,430	
January – December	13,730	

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Other operating income, SEK '000s

Operating costs



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YTD Operating costs, SEK '000s

FINANCE

Profit / loss for the period



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SEK millions





2021 Cash development

Cash development including SKAT balance, SEK millions



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Shareholder Composition

During 2021 No. of shareholders grows from ~6,000 to ~14,000

No. of shareholders



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No. of shares outstanding

Thank you!

Contact:

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Our Technology



ExpreS² Platform for Complex Proteins Enables unique non-viral approach to protein and vaccine production



Schneider I (1972). "Cell Lines Derived from Late Embryonic Stages of Drosophila melanogaster". J. Embryol. Exp. Morphol. 27: 363–365 Note: ExpreS²ion Biotech founders invented an Improved Vector System derived from S2 cells; granted patent until 2032 (US); glyco-engineered S2 cells pending patents until 2040.



Reducing risks in discovery manufacturing

- Fast & high level protein expression
- Robust; high batch-to-batch consistency
- Superior success rates in early research

Proprietary process and expertise has established ExpreS²ion as the leader in specialty protein production

- \checkmark 20+ years of experience
- \checkmark Over 90% success rate, over 350 proteins expressed
- \checkmark Go-to source for challenging proteins
- \checkmark Rapid delivery (3-6 months) of high-quality, uniform proteins with exceptional yields

Virus Like Particle (VLP) Technology VLP technology has proven track record in cancer vaccine applications (HPV)

Our ExpreS² platform produces the complex surface proteins (antigens), which are critical to immune system recognition and response

AdaptVac's proprietary viruslike particles technology¹

securely attaches our proteins to the surface of a spherical shell (capsid), mimicking a virus to elicit an immune response





AdaptVac ApS is a joint venture established in 2017 owned by ExpreS²ion Biotechnologies (34%) and NextGen Vaccines (66%), a spinout of the Department of Immunology and Microbiology, the University of Copenhagen, led by Professor Ali Salanti



High immunogenic potential

other VLP approaches can only support fragments

High density display on surface (180 attachment

Directional attachment (vs random orientation in

