

### Forward-looking statements and disclaimer

This report contains forward-looking statements. The words "believe", "expect", "anticipate", "intend" and "plan" and similar expressions identify forward-looking statements. All statements other than statements of historical facts included in this report, including, without limitation, those regarding our financial position, business strategy, plans and objectives of management for future operations (including development plans and objectives relating to our products), are forward-looking statements. Such forward-looking statements involve known and unknown risks, uncertainties and other factors which may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements. Such forwardlooking statements are based on numerous assumptions regarding our present and future business strategies and the environment in which we will operate in the future. The important factors that could cause our actual results, performance or achievements to differ materially from those in the forward-looking statements include, among others, risks associated with product discovery and development, uncertainties related to the outcome of clinical trials. slower than expected rates of patient recruitment, unforeseen safety issues resulting from the administration of our products in patients, uncertainties related to product manufacturing, the lack of market acceptance of our products, our inability to manage growth, the competitive

environment in relation to our business area and markets, our inability to attract and retain suitably qualified personnel, the unenforceability or lack of protection of our patents and proprietary rights, our relationships with affiliated entities, changes and developments in technology which may render our products obsolete, and other factors. Further, certain forward-looking statements are based upon assumptions of future events which may not prove to be accurate. The forward-looking statements in this document speak only as at the date of this report. ExpreS<sup>2</sup>ion Biotech does not undertake any obligation to update or revise forward-looking statements in this report nor to confirm such statements to reflect subsequent events or circumstances after the date made or in relation to actual results, unless required by law.

#### **Definitions**

"ExpreS<sup>2</sup>ion Biotech Holding AB" refers to ExpreS<sup>2</sup>ion Biotech Holding AB with corporate identity number 559033-3729. "The Company" or "ExpreS<sup>2</sup>ion" refers to the group, i.e. ExpreS<sup>2</sup>ion Biotech Holding AB and its fully owned operational subsidiary ExpreS<sup>2</sup>ion Biotechnologies ApS, Denmark.





# A word from our CEO

During the second quarter of 2021, we reported on continued progress in key pipeline areas for ExpreS<sup>2</sup>ion including COVID-19, breast cancer and malaria, while also securing 39 MSEK in additional funding from the exercise of warrants of series TO4. I want to start this CEO comment with thanking everyone who contributed to this funding by exercising TO4 warrants for believing in the future of ExpreS<sup>2</sup>ion. As previously stated, I am proud to say that ExpreS<sup>2</sup>ion's Board of Directors and Executive Management contributed to this funding by exercising all TO4 warrants held

COUGH-1, the clinical phase I/II study for the ABNCoV2 capsid virus-like particle (cVLP) based COVID-19 vaccine, continued to progress during the quarter with an interim safety update published in April. At that time, the first group of 18 healthy volunteers had been satisfactorily administered with the vaccine without any untoward safety signals observed. After the end of the period, we were excited to announce excellent safety and efficacy headline results from the study, including

strong virus neutralization levels also for the Delta and Beta variants. Virus neutralization levels were up to 12 times higher compared to what is seen after COVID-19 infection, compared with up to 4.1 times higher for currently approved COVID-19 vaccines. This positions ABNCoV2 as a strong candidate for world-wide use, either on its own or as a booster vaccine. Bavarian Nordic is now preparing a Phase II booster clinical study, and also a Phase III trial in 2022 pending external funding.

Another important achievement during the quarter was the selection of the lead candidate ES2B-CO01 in our HER2-cVLP breast cancer vaccine project. This was done as a part of the preclinical program, which has been initiated with the expert assistance of Prof. Pier Luigi Lollini at the Interdepartmental Center for Cancer Research, University of Bologna. This collaboration has enabled us to use state-of-the-art in vitro and in vivo models for some of the mechanisms in human breast cancer, and the preclinical program is off to a really good start with preclinical proof of concept data expected in Q1 2022.

"I want to highlight that we are now really starting to see overall positive effects from the ongoing strengthening of our organization in terms of much needed added capacity in key areas. It is truly inspiring to see our growing team perform so well together each day, and this bodes well for the short as well as long term success of the company and our pipeline projects."

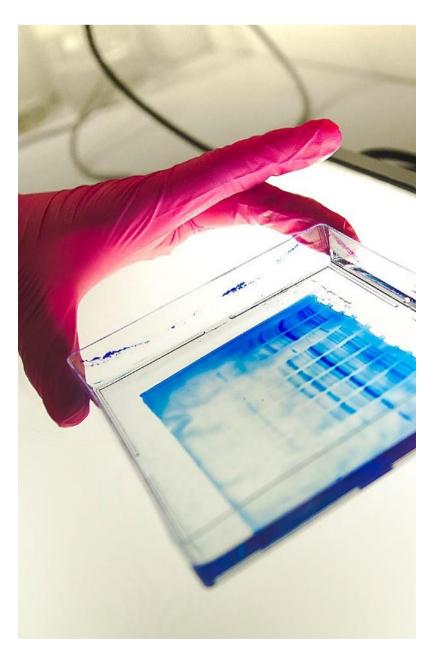
As I stated in the CEO comment for our Q1 interim report, we are seeing increased activity in our malaria projects in 2021 due to COVID-19 research programs taking up less of our key collaboration partners' time. An example of this was the publication of a scientific article in April on the outcome of the VACO63-study, which is the first bloodstage malaria clinical trial. The University of Oxford is running this program, and we are contributing with our ExpreS2™ platform for the production of the recombinant RH5.1 protein. After the end of the period, we were excited to announce that the next step, a Phase Ib clinical trial to assess the safety and immunogenicity of the updated RH5.1/Matrix-M vaccine candidate in adults and infants living in Tanzania, has been initiated.

In addition to the achievements reported during the quarter, I want to highlight that we are now really starting to see overall positive effects from the ongoing strengthening of our organization in terms of much needed added capacity in key

areas. It is truly inspiring to see our growing team perform so well together each day, and this bodes well for the short as well as long term success of the company and our pipeline projects. This also makes me confident in our ability to present a strong news flow also in the coming quarters.

#### Bent U. Frandsen

CEO, ExpreS2ion Biotech Holding AB



# About ExpreS<sup>2</sup>ion

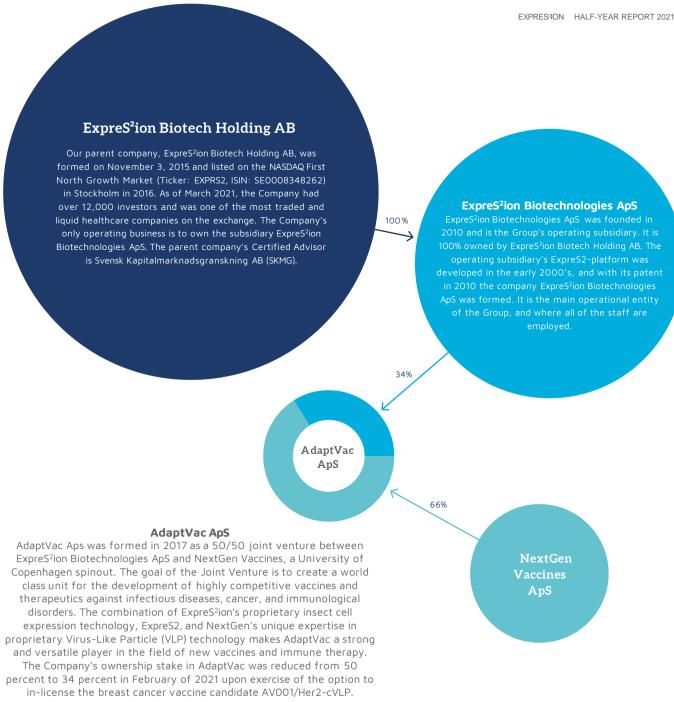
ExpreS<sup>2</sup>ion was founded in 2010 on the realisation that to produce the complex proteins needed for the biological drugs and vaccines of the future, in a safer and more efficient manner, a new protein expression system would be needed.

The ExpreS2 technology platform was developed to be especially well suited for production of the proteins required for the development and production of vaccines and immunotherapy products. The platform is based on insect cells, so called *Drosophila melanogaster* (fruit fly) S2 cells combined with patented expression vectors (the genetic tool researchers employ to commandeer the cell's internal protein production machinery) and especially adapted culture agents and reagents which are needed to make the cells thrive and grow. Among the platform's many advantages are:

- Significantly less costly and timeconsuming than alternative methods,
  which is an important competitive
  advantage, considering time-tomarket and patent expiry. It also
  makes the platform particularly
  valuable for the development of
  diagnostics and vaccines in epidemic or
  pandemic situations where speed is of
  the essence.
- Generates higher yields, i.e. amount of protein per manufacturing batch, compared to competing systems.
- Provides homogeneous manufacturing batches, a requirement in pharmaceutical development. The platform includes the Company's patented expression vectors which were developed, among other things, to make it possible for the cells to generate higher yields.

# Company structure

ExpreS2ion has a streamlined company structure. ExpreS2ion Biotech Holding AB is the Swedish entity listed on Nasdag First North Growth Market since 2016. ExpreS<sup>2</sup>ion Biotechnologies ApS is the operational entity, with offices and labs in the Scion DTU Science park 20 km north of Copenhagen, Denmark, and was established in 2010. AdaptVac ApS is a joint venture established in 2017 together with a group of scientists from the Institute of Immunology and Microbiology at the University of Copenhagen. The scientists own their part of AdaptVac through a joint holding company named NextGen Vaccines ApS.



# **Business** model

The Company's business model is to develop, produce and deliver therapeutic or diagnostic proteins, as well as to generate revenue by outlicensing the ExpreS2 platform to research institutes and pharmaceutical companies who themselves or in cooperation with the Company develop biopharmaceutical drugs and vaccines. This model generates short term revenue for the Company and carries potential future royalties, license fees, and milestone payments through pharmaceutical products developed using the Company's technology.

Under its new strategy the service model above will be complemented by the Company increasingly building its own pipeline of preclinical and later clinical biopharmaceutical drug and vaccine candidates. Under this new model, the Company will carry out its own initial research, preclinical and early clinical development work prior to out-licensing. The recent agreement with Bavarian Nordic, under which Bavarian Nordic assumes all future development costs for the COVID-19 vaccine program and pay certain milestones and royalties, subject

to external funding, is the first example of this new strategy.

The Company believes that the combination of a continued successful service model combined with the creation of an inhouse pipeline of biopharmaceutical drug and vaccine candidates puts it in a good position to balance risk and return and create value for its shareholders

#### Resources

- ExpreS2-platform built for optimal discovery, preclinical development and GMP production of hard-toexpress proteins used in vaccines and vaccine-like treatments
- ES2B CO01 HER-cVLP exclusive global license

A biotech company creating value through pipeline development and service offerings

#### **Our businesses**

- PIPELINE DEVELOPMENT
- Vaccine & immunotherapy candidates
- Partnerships
- Research & commercial licenses
- Driven by ExpreS<sup>2</sup>ion and partners

#### - CRO SERVICES BUSINESS

- Cells, kits, reagents & proteins
- Cell banks, processes & analytics
- Platform research licenses
- Driven by clients

#### Values

- Result oriented
- Entrepreneurial spirit
- Expertise
- Teamwork makes us stronger
- Learning
- Fun
- Quick decision making
- Celebrating / respecting our differences



# Sources of income

With over 100 currently active or former academic and industrial service and license contracts, the Company has built a large network in the international research community since its inception in 2010. Furthermore, the Company is currently a part of an international research consortia which together has been granted more than an estimated EUR 40 million of non-dilutive public funding.

The Company also sells licenses to use the ExpreS2 platform as a whole or in part, thus allowing its clients to participate in or be entirely responsible for the development of the required proteins. The Company sells ExpreS2 test kits and reagents for application as research tools or diagnostics. The Company may also enter into agreements in which the client accepts a quotation and is charged for the development, production and delivery of research grade proteins, using the ExpreS2 platform.

The Company services both pharmaceutical companies and research institutions. The ExpreS2 platform is equally suited for academic research, analytics and commercial drug development, both in vaccines and other biopharma fields. The Company's clients are not limited to any geographic area

and are located all over the world. Since its foundation in 2010, the Company has worked with more than 100 clients and partners. The agreements with these clients, which in many cases are worldleading universities, research institutions and pharmaceutical companies, have generated significant revenues for the Company over the years. It currently has more than ten major clients. For instance. the Company has out-licensed the ExpreS2 platform for research to Hoffman-La Roche, Imperial College London and Francis Crick Institute among others, and out-licensed the platform for clinical development to the University of Copenhagen and the Jenner Institute of the University of Oxford, among others. Five of the Company's current material transfer agreements (MTAs) relate to the transfer of Company-made SARS-CoV-2 material for various COVID-19 diagnostic and research support purposes.

# **Pipeline**

DISEASE		Project / Target	Discovery	Pre-clinical Pharmacology	cGMP / Tox	Phase 1	Phase 2	Phase 3	Market potential
Corona virus		ABNCoV2/SARS-CoV-2 cVLP				1/11	BN: II		> 30 billion EUR
Breast cancer	The state of the s	ES2B-C001/Her2 cVLP							> 10 billion EUR
Influenza		Hemagglutinin							> 4 billion EUR
Malaria									> 0.4 billion EUR
I: Blood		RH5					lb / lla		
II: Blood		RH5-VLP							
III: Transmissio	n	Pfs 48/45							
IV: Placenta		VAR2CSA			la / Ib				
V: Blood		CYRPA complex							

As of August 2021

# Pipeline



#### **CORONAVIRUS/COVID-19**

ExpreS<sup>2</sup>ion and its joint venture partner AdaptVac are engaged in the development of a unique capsid viruslike particle (cVLP) COVID-19 vaccine, partly sponsored through a Horizon 2020 EU grant award to the PREVENT-nCoV consortium to rapidly advance the vaccine candidate against COVID-19 into the clinical stage. The candidate vaccine is a cVLP applying ExpreS2-produced SARS-CoV-2 antigens, thereby creating a powerful immunogenic vaccine.

AdaptVac and AGC Biologics, a global Contract Development and Manufacturing Organization (CDMO) for biopharmaceuticals have entered into a partnership for the scale-up and cGMP manufacture of the COVID-19 vaccine. In July 2020. AdaptVac and Bayarian Nordic. a fully integrated biotechnology company focused on the development. manufacture and commercialization of life-saving vaccines, have entered into a license agreement which provides Bavarian Nordic the global commercialization rights to the proprietary capsid virus like particle based SARS-CoV-2 subunit vaccine, designated ABNCoV2. For application of our proprietary protein production system ExpreS2, ExpreS2ion and AdaptVac have also entered into a license agreement for this project.

In addition to ExpreS<sup>2</sup>ion and AdaptVac, the PREVENT-nCoV consortium members are Leiden University Medical Center (LUMC), Institute for Tropical Medicine (ITM) at University of Tübingen, The Department of Immunology and Microbiology (ISIM) at University of Copenhagen, the Laboratory of Virology at Wageningen University, and Radboud University Medical Center. We announced the first headline results of the clinical

Phase I/II in August 2021 and demonstrated positive safety and efficacy outcomes. The next step is a Bavarian Nordic sponsored Phase II study to determine potential as a universal booster vaccine.



#### **BREAST CANCER**

Breast cancer is a widespread oncology indication affecting more than 1.3 million people worldwide annually, resulting in more than 450,000 deaths (Tao, 2015: www.ncbi.nlm.nih.gov/pubmed/25 543329). The most common treatment today is based on monoclonal antibodies, where the

dominating therapy Herceptin (trastuzumab) generates annual global sales of USD 7 billion. The target product profile of AdaptVac's lead breast cancer project, AVOO1(HER2-cVLP), is tailored to be highly competitive both in terms of cost and efficacy, thus aiming at a significant market share.

In February 2021, ExpreS<sup>2</sup>ion signed a final patent license agreement with AdaptVac whereby ExpreS<sup>2</sup>ion exclusively licensed in AV001 (renamed ES2B-C001). This gives ExpreS<sup>2</sup>ion full control over and responsibility for driving this valuable asset forward, hereby realising the very significant value of this project.



#### **INFLUENZA**

The international next-generation influenza vaccine consortium INDIGO, led by the University of Amsterdam with ExpreS<sup>2</sup>ion as a participating member, is developing a next-generation influenza vaccine in a large collaboration between public and private R&D organisations from the EU, India, and the United States. The project has been awarded a 10.0 MEUR Horizon 2020 grant from the EU, of which ExpreS<sup>2</sup>ion's participation was directly awarded 0.6 MEUR.

The INDIGO consortium plans to carry out the preclinical and clinical development of the project, which contains two novel influenza vaccine concepts, including the application of a novel potent adjuvant by LiteVax BV, the Netherlands, as well as the use of the ExpreS2 platform for antigen production by ExpreS2ion. The aim is to create an influenza vaccine that meets the requirements of global vaccination, i.e.

to achieve <10% instead of 60% nonresponders, combined with a lower manufacturing cost and better accessibility.



#### **MALARIA PROJECTS**

#### Malaria I

#### Blood stage (RH5-1)

Jenner Institute of the University of Oxford is developing the blood-stage Plasmodium falciparum malaria antigen RH5.1 with ExpreS<sup>2</sup>ion as a collaboration partner. The RH5.1 antigen is produced in ExpreS<sup>2</sup>ion's ExpreS2 platform.

#### Malaria II

#### Blood stage (RH5-2)

With the aim to further improve efficacy, Jenner Institute of the University of Oxford is developing a second-generation RH5 vaccine, RH5.2, in the ExpreS2 platform. RH5.2 has been

engineered to retain regions important for red blood cell recognition, which are targeted by neutralising antibodies. Additionally, the RH5.2 protein will be displayed on the surface of a hepatitis B derived virus-like particle (VLP) in order to maximise the induction of high titre antibodies. The project is founded by the Wellcome Trust.

#### Malaria III

#### Transmission (Pfs48/45)

The goal for a transmission-blocking vaccine is to prevent the transfer to mosquitos feeding on persons infected with malaria, thus effectively hindering further spread of the disease. Thereby a transmission-block- ing vaccine does not give direct protection from the disease. but it stops the disease from spreading and could therefore lead to eradication of malaria. During the last decade, the inability to produce the fulllength Pfs48/45 antigen has been a major roadblock for researchers aiming to create a transmission-blocking malaria vaccine. However, this challenge was overcome by ExpreS<sup>2</sup>ion and Jenner Institute at the University of Oxford.

This vaccine is developed by the Horizon 2020-funded OptiMalVax grant consortium, led by Jenner Institute at the University of Oxford with ExpreS²ion as a member. The objective of the consortium is to create a combination malaria vaccine, and its clinical program will include trials to assess the pre-

erythrocytic, blood-stage and mosquitostage components of the combi- nation vaccine, including this transmission vaccine

#### Malaria IV

#### Placenta borne (VAR2CSA)

ExpreS<sup>2</sup>ion is a part of the PlacMalVac project that started in 2013 as an international consortium project with the aim to develop a vaccine against placental malaria. The project is based on the antigen VAR2CSA, which enable parasite accumulation in the placenta and was discovered by Professor Ali Salanti and others at the University of Copenhagen.

#### Malaria V

#### Blood-stage (PfRipr complex)

An international research team, including scientists from ExpreS<sup>2</sup>ion and led by the Walter and Eliza Hall Institute of Medical Research (WEHI), is developing a next generation malaria vaccine that is targeting a recently discovered molecular 'key' that the deadly malaria parasite uses to enter human blood cells. The malaria 'key' was first described in a Nature article, published December 2018 from the group. It is a com- plex of three parasite proteins called Rh5, CyRPA and Ripr, where the three proteins work together to unlock and enter the cell. This central role in the infection of human blood cells makes the complex a new and promising target for vaccine development. The vaccine is based on a patent co-owned by WEHI and ExpreS<sup>2</sup>ion.

### Significant events

#### Second quarter of 2021

On April 12, ExpreS<sup>2</sup>ion announced that the first group of volunteers in the clinical Phase I/II study, COUGH-1, had been satisfactorily administered with the ABNCoV2 capsid virus-like particle (cVLP) based COVID-19 vaccine. The clinical study ran as planned with no untoward safety signal in 18 healthy volunteers.

On April 21. ExpreS<sup>2</sup>ion announced that its protein production platform ExpreS2™ has contributed to a scientific article published in the journal Med. The article highlights the outcome of the VACO63-study, a Phase I/IIa clinical trial to assess the safety. immunogenicity and efficacy of the bloodstage Plasmodium falciparum malaria vaccine candidate RH5.1/ASO1B. In conclusion the RH5.1/ASO1B vaccine is safe. well tolerated, and immunogenic in healthy adults. A significantly reduced blood-stage parasite growth rate was observed in vaccinees following controlled human malaria infection, a defining milestone for the blood-stage malaria vaccine field.

On April 26, ExpreS<sup>2</sup>ion announced that Gitte L. Pedersen would not seek re-election to the Board of Directors at the Annual General Meeting on May 26, 2021, and that two new members, Karin Garre and Sara Sande, were proposed for election. Thus, it was proposed to re-elect Jakob Knudsen, Martin Roland Jensen and Allan Rosetzsky and to elect Karin Garre and Sara Sande as new directors and to re-elect Martin Roland lensen as chairman of the board

On April 28, ExpreS<sup>2</sup>ion announced the outcome of the exercise of warrants of series TO4, which were issued in connection with the Company's rights issue of units in 2020. In total, 5,324,670 warrants of series TO4 were exercised, corresponding to approximately 97.6 percent of the total number of outstanding warrants of series TO4, for subscription of 1,774,890 shares at an exercise price of SEK 22.00 per share. ExpreS<sup>2</sup>ion received approximately SEK 39.0 million before issuing costs through the exercise of the warrants of TO4.

On May 11, ExpreS<sup>2</sup>ion announced the selection of its lead candidate HER2-cVLP breast cancer vaccine, and that the project runs according to plan.

On May 26, ExpreS<sup>2</sup>ion held the 2021 Annual General Meeting (AGM), during which resolutions were passed related to the adoption of the income statement and balance sheet, allocation of profit, discharge from liability, election of the Board of Directors, Auditor and remuneration, changes to the articles of association, security issuance authorization, and incentive programs. Due to the corona pandemic, the AGM was carried out through

postal voting only, without physical presence.

#### After the end of the period

On July 21, ExpreS2ion announced that University of Oxford had initiated the VACO80-study, a Phase Ib clinical trial to assess the safety and immunogenicity of the blood-stage Plasmodium falciparum malaria vaccine candidate RH5.1/Matrix-M in adults and infants living in Tanzania. The RH5.1 blood-stage malaria protein vaccine has previously been administered to 67 healthy UK adults, with various doses, and was found to be safe and well tolerated. The study is estimated to be completed in H2 2023. The primary aim of the new Phase Ib trial is to assess the safety and immunogenicity of the RH5.1/Matrix-M formulation in a malaria-endemic population for the first time.

On August 9, ExpreS2ion announced that COUGH-1, the COVID-19 Phase I/II clinical trial to evaluate the ABNCoV2 vaccine, as headline results met its safety and efficacy endpoints with excellent virus neutralization levels of up to 12 times higher compared to the levels achieved after COVID-19 infection. This was significantly higher than the virus neutralization levels reported for leading mRNA COVID-19 vaccines reaching only up to 4.1 times higher than the levels achieved after COVID-19 infection. High efficacy was reported in all groups receiving ABNCoV2,

including the lowest dose ranges and nonadjuvanted formulations. Importantly, high virus neutralization levels were shown also for relevant COVID-19 variants such as the dominant Delta and the escape Beta variant.

# 2021 - 2023 Outlook

			2021					2022			2023
are in	CORONAV	IRUS (ABNC	oV2)			 					
	Phase I/II trial, COUGH-1 initiated	▼ COUGH-1 initial safety results (Q2)	COUGH-1 full safety & efficacy results (Q3)	BN Phase II trial initiation (Q3)	BN Phase II trial readout	BN Phase trial initi pending funding		BN Phase initial readout	Ready to market (subject regulate approve	launch to ory	
Thurs	BREAST CA	ANCER (ES2	B-C001)			 					
	Executed in-licensing (Feb 2021)	Preclinical animal sinitiated	tudies		Preclinica proof-of- results		GMP manuf batch	facturing & tox	Filing of clinical trial application	Initiation of first human clinical trial	Outlicensing window opens pending human data
	INFLUENZA	Α									
		in pre	n INDIGO progres clinical animal es in (H2)		pport further at of one or more n 2021						
*	MALARIA										
	<b>Ø</b>	Phase IIa resu the Rh5.1 vacc published in 2	cine	Additional pha malaria endem launched durin alternative adj	nic region in Africa ng 2021, with					Rh5 phase I readout	trial

# Summary of Q2 interim results



### Second quarter (April – June 2021)

- Operating income amounted to 4,793 (4,016) KSEK.
- Profit/loss after financial items amounted to -10,136 (-4,953) KSEK.
- Profit/loss for the period amounted to -9,068 (-4,239) KSEK.
- Net income per share\* amounted to -0.32 (-0.27) SEK.



#### First half (January - June 2021)

- Operating income amounted to 6,724 (6,845) KSEK.
- Profit/loss after financial items amounted to -22,264 (-10,888) KSEK.
- Profit/loss for the period amounted to -20,223 (-9,473) KSEK.
- Net income per share\* amounted to -0.72 (-0.64) SEK.

#### **Key financials**

Q2 2021	Q2 2020	% Change
4,793	4,016	19%
-10,136	-4,953	105%
-9,068	-4,239	114%
-0.32	-0.27	18%
108,289	12,764	748%
128,398	28,291	354%
88%	15%	73%

H1 2021	H1 2020	% Change
6,724	6,845	-2%
-22,264	-10,888	104%
-20,223	-9,473	113%
-0.72	-0.64	13%
108,289	12,764	748%
128,398	28,291	354%
88%	15%	73%

FY 2020
15,263
-34,923
-31,713
-1.83
106,832
118,858
80%

Figures in parenthesis are the numbers from the same period in 2020.

<sup>\*</sup>The Group's net income per share: The net income for the period divided with the average number of shares for the period. For the period January to June 2021, the average number of shares amounted to 28,177,050. As of 30/6/2021, the total number of shares in ExpreS<sup>2</sup>ion Biotech Holding AB was 29,383,191.

<sup>\*\*</sup>Equity ratio: Shareholder's equity divided by total capital.

### Financial overview

#### Development in figures for Q2 2021

#### Revenue

Total operating income during the second quarter of 2021 amounted to KSEK 4,793 (4,016). The revenues in the second quarter of 2021 were 19% higher compared to the same period last year. Thus far in 2021, there has been a significant shift in the mix of revenues coming from grant projects, which were very strong in 2020, to client driven projects which have partially recovered from the 2020 decline related to COVID-19. The key drivers include follow-on projects related to development of the COVID-19 vaccine, and royalty and direct revenues from the sale of ExpreS<sup>2</sup>ion-produced proteins. This is a reversal of the trend experienced in 2020.

#### Net result

The consolidated net loss for the second quarter of 2021 amounted to KSEK -9,068 (-4,239). The lower result is primarily due to the combination of (1) higher R&D, raw materials and consumables costs related to the transition to a pipeline driven strategy, and specifically the Company's investment in the ES2B-CO01 breast cancer therapeutic vaccine candidate, and (2) changes to how the Company accrues for incentive compensation, specifically warrants and bonuses, which impacted personnel costs. The personnel cost changes do not have a cash impact and are consistent with changes made earlier this year when we published the 2020 annual report. These factors more than offset higher operating income, a one-time dividend from the company's joint venture with Nextgen Vaccines Aps, AdaptVac Aps, and a higher tax deduction.

#### Cash and cash equivalents

As of June 30, 2021, ExpreS<sup>2</sup>ion's cash and cash equivalents amounted to KSEK 108,289 (12,764). During the quarter, cash increased by SEK 26.3 million, primarily reflecting SEK 39 million raised through a warrant subscription in April. Other key drivers of cash in the quarter include (1) an operating loss of SEK 10.4 million, (2) SEK 3.3 million in costs of issuing shares related to the warrant subscription, and (3) SEK +2.5 million in adjustments for items not included in the cash flow primarily related to depreciation and warrant fair value charges.

Figures in parenthesis are the numbers from the same period in 2020.

#### Development in figures for the first six months of 2021

#### Revenue

Total operating income during the first six months of 2021 amounted to KSEK 6,724 (6,845). The revenues in the first six months of 2021 were 2% lower compared to the same period last year. As mentioned to the left, net sales from client projects have recovered significantly from the significant headwinds faced in 2020 as COVID-19 resulted in project delays. Offsetting is much lower other operating income, driven by grants, which benefitted from a large COVID-19 related grant in 2020.

#### Net result

The consolidated net result for the first six months of 2021 amounted to KSEK -20,223 (-9,473). The decline is driven by investment in pipeline projects, notably the pre-clinical development of the ES2B-C001 breast cancer vaccine, as well as the personnel related factors described for the second quarter. These were partially offset by a one-time dividend payment from the company's joint venture with Nextgen Vaccines ApS, AdaptVac Aps, a larger income tax deduction and lower depreciation.

### Income statement - group

KSEK	Q2 2021	Q2 2020	% change	H12021	H12020	% change	FY 2020
Operating income							
Net sales	4,584	1,545	197%	6,288	3,768	67%	5,259
Other operating income	209	2,471	-92%	436	3,077	-86%	10,004
Total operating income	4,793	4,016	19%	6,724	6,845	-2%	15,263
Operating costs							
Rawmaterials & consumables	-3,356	-2,315	45%	-6,273	-3,254	93%	-6,102
Other external costs	-3,503	-2,635	33%	-9,456	-5,214	81%	-21,450
Personnel costs	-7,909	-3,178	149%	-12,172	-6,875	77%	-15,990
Depreciation of tangible & intangible fixed assets	-432	-734	-41%	-1,174	-1,452	-19%	-2,91
Total operating costs	-15,200	-8,862	72%	-29,075	-16,795	73%	-46,459
Operating profit/loss	-10,407	-4,846	115%	-22,351	-9,950	125%	-31,196
Result from financial investments							
Result in jointly governed companies	463	143	224%	670	-197	-440%	-194
Other interest income & similar items	-34	-9	275%	-34	-9	273%	-8
Interest expense & similar items	-159	-241	-34%	-550	-732	-25%	-3,52
Total result from financial investments	271	-107	-352%	87	-938	-109%	-3,727
Profit/loss after financial items	-10,136	-4,953	105%	-22,264	-10,888	104%	-34,923
Income tax on the result for the period	1,068	714	50%	2,041	1,415	44%	3,210
Profit/loss for the period	-9,068	-4,239	114%	-20,223	-9,473	113%	-31,713

### Balance sheet - group

KSEK	Q2 2021	YE 2020	% change	Q2 2020
Assets				
Concessions, patents, licenses, trademarkets				
and similar intellectual rights	3,363	3,907	-14%	4,863
Goodwill	0	194	-100%	505
Total non-current intangible assets	3,363	4,101	-18%	5,368
Plants and machinery	1,419	1,294	10%	1,381
Total non-current tangible assets	1,419	1,294	10%	1,381
Interest in jointly governed companies	23	34	-33%	35
Other long-term receivables	1,066	966	10%	957
Total non-current financial assets	1,089	1,000	9%	992
Total non-current assets	5,871	6,395	-8%	7,741
Accounts receivable	3,100	525	490%	1,024
Tax receivables	4,737	2,788	70%	3,304
Other receivables	5,739	1,791	220%	2,347
Prepaid expenses and accrued income	662	527	26%	1,111
Total receivables	14,238	5,631	153%	7,786
Cash and bank	108,289	106,832	1%	12,764
Total current assets	122,527	112,463	9%	20,550
TOTAL ASSETS	128,398	118,858	8%	28,291

кѕек	Q2 2021	YE 2020	% change	Q2 2020
Equity and liabilities				
Share capital	3,265	3,068	6%	1,789
Other capital contributions	130,445	122,921	6%	12,106
Other equity including net loss for the period	-20,652	-31,441	-34%	-9,697
Total equity	113,058	94,548	20%	4,198
Provision for taxes	709	827	-14%	1,032
Total provisions	709	827	-14%	1,032
Other long-term liabilities	4,270	5,272	-19%	6,358
Total long-term liabilities	4,270	5,272	-19%	6,358
Liabilities to credit institutions	1,966	1,889	4%	1,221
Accounts payable	1,739	2,078	-16%	1,401
Other liabilities	6,656	14,244	-53%	14,081
Total short-term liabilities	10,361	18,211	-43%	16,703
TOTAL EQUITY AND LIABILITIES	128,398	118,858	8%	28,291

### Cash flow statement - group

KSEK	Q2 2021	Q2 2020	% change	H12021	H12020	% change	FY 2020
Operating profit/loss	-10,407	-4846	115%	-22,351	-9950	125%	-31,196
Adjustments for items not included in the cash flow	2,502	734	241%	3,411	1,452	135%	3,201
Received interest	-34	-9	278%	-34	-9	278%	0
Interest paid	-201	-124	62%	-594	-584	2%	-3,137
Income tax received	-2	-4	-45%	-5	-4	30%	2,046
Cash flow from operating activities before changes in working capital	-8,142	-4,249	92%	-19,573	-9,095	115%	-29,086
Decrease(+)/increase(-) of current receivables	-4,014	804	-599%	-6,731	-827	714%	-336
Decrease(+)/increase(-) of current liabilities	3,160	7,382	-57%	-8,075	6,319	-228%	11,247
Cash flow from operating activities	-8,996	3,937	-329%	-34,379	-3,603	854%	-18,175
Investments in jointly governed companies	463	141	228%	681	-193	-453%	-194
Investments in intangible non-current assets	0	0	n/a	0	0	n/a	0
Investments in tangible non-current assets	-392	-304	29%	-512	-534	-4%	-885
Cash flow from investing activities	71	-163	-144%	169	-727	-123%	-1,079
Leasing agreement	-154	75	-305%	-308	-79	290%	-415
Bridge loan	-339	3,167	-111%	-669	3,167	-121%	3,172
Payment for warrants	0	0	n/a	0	0	n/a	2,656
Issuance of new shares	39,048	0	n/a	39,048	9,600	307%	140,527
Costs of issuing shares	-3,347	-137	2343%	-3,347	-721	364%	-22,558
Cash flow from financing activities	35,208	3,105	1034%	34,724	11,967	190%	123,382
Cash flow for the period	26,282	6,879	282%	5 13	7,637	-93%	104,128
Cash and cash equivalents at the beginning of the period	82,807	6,194	1237%	106,832	5,418	1872%	5,418
Exchange difference cash and cash equivalents	-800	-309	159%	944	-291	-424%	-2,714
Cash and cash equivalents at the end of the period	108,289	12,764	748%	108,289	12,764	748%	106,832

### Changes in shareholders equity - group

#### H12021

кѕек	Share capital	Other capital contributions	Other equity including net profit for the period	Total equity
Opening balance as of January 1st, 2021	3,067	178,042	-86,561	94,548
Issuance of new shares	198	38,850		39,049
Payments for warrants				0
Issuing expenses		-3,348		-3,348
Vesting of share-based compensation		2,244		2,244
Exchange difference for the period			789	789
Profit-loss for the period			-20,223	-20,223
Total equity as of June 30th, 2021	3,265	215,789	-105,996	113,058

#### FY 2020

		Other capital	Other equity including net profit for the	
KSEK	Share capital	contributions	period	Total equity
Opening balance as of January 1st, 2020	1,512	50,100	-52,691	-1,079
Issuance of new shares	1,212	129,715		130,927
Payments for warrants	226	12,030		12,256
Issuing expenses		-22,558		-22,558
Conversion of debt	117	8,471		8,588
Vesting of share-based compensation		284		284
Exchange difference for the period			-2,157	-2,157
Profit-loss for the period			-31,713	-31,713
Total equity as of December 31st, 2020	3,067	178,042	-86,561	94,548

### Income statement - parent

KSEK	Q2 2021	Q2 2020	% change	H12021	H12020	% change	FY 2020
Operating income							
Net sales	167	167	0%	167	167	0%	335
Total operating income	167	167	0%	167	167	0%	335
Operating costs							
Other external costs	-2,122	-766	177%	-2,548	-1,243	105%	-2,675
Personnel costs	-643	-19	3284%	-693	-113	513%	-363
Total operating costs	-2,765	-785	252%	-3,241	-1,356	139%	-3,038
Operating profit/loss	-2,598	-618	320%	-3,074	-1,189	159%	-2,703
Result from financial investments							
Other interest income & similar items	0	130	-100%	0	166	-100%	390
Interest expense & similar items	-52	-84	-38%	-171	-310	-45%	-2,584
Total result from financial investments	-52	46	-213%	-171	-144	19%	-2,194
Profit/loss after financial items	-2,650	-572	363%	-3,245	-1,333	143%	-4,897
Income tax on the result for the period	0	0	n/a	0	0	n/a	0
Profit/loss for the period	-2,650	-572	363%	-3,245	-1,333	143%	-4,897

### Balance sheet - parent

	_			
KSEK	Q2 2021	YE 2020	% change	Q2 2020
Assets				
Shares in group companies	165,887	165,887	0%	45,053
Receivables from group companies	0	0	n/a	10,611
Other non-current receivables	0	0	n/a	50
Total financial non-current assets	165,887	165,887	0%	55,714
Total non-current assets	165,887	165,887	0%	55,714
Tax receivables	40	32	0	41
Other receivables	195	397	-51%	1,147
Prepaid expenses and accrued income	60	60	0%	620
Total receivables	295	489	-40%	1,808
Cash and bank	44,363	5,069	775%	3,723
Total current assets	44,658	5,558	703%	5,531
TOTAL ASSETS	210,545	171,445	23%	61,245

KSEK	Q2 2021	YE 2020	% change	Q2 2020
NOLIN	Q2 2021	12 2020	70 Change	Q2 2020
Equity and liabilities				
Share capital	3,265	3,067	6%	1,789
Restricted equity	3,265	3,067	6%	1,789
Share premium fund and retained earnings	204,414	169,964	20%	57,125
Profit/loss for the period	-3,620	-3,672	-1%	-1,332
Unrestricted equity	200,794	166,292	21%	55,793
Total equity	204,059	169,359	20%	57,582
Payables to group companies	5,804	1,801	222%	3,500
Other liabilities	682	285	139%	163
Total short-term liabilities	6,486	2,086	211%	3,663
TOTAL EQUITY AND LIABILITIES	210,545	171,445	23%	61,245

### Changes in shareholders equity - parent

#### H12021

кѕек	Share capital	Other capital contributions	Other equity including net profit for the period	Total equity
Opening balance as of January 1st, 2021	3,067	171,502	-5,210	169,359
Appropriation of retained earnings according to the AGM				0
Issuance of new shares	198	38,850		39,049
Payments for warrants				0
Issuing expenses		-3,348		-3,348
Conversion of debt				0
Vesting of share-based compensation		2,244		2,244
Profit-loss for the period			-3,245	-3,245
Total equity as of June 30th, 2021	3,265	209,249	-8,455	204,059

#### FY 2020

KSEK	Share capital	Other capital contributions	Other equity including net profit for the period	Total equity
Opening balance as of January 1st, 2020	1,512	45,615	-2,368	44,759
Appropriation of retained earnings according to the AGM	0	-2,055	2,055	0
Appropriation of retained earnings relating to misstatement in 2019	0	0	0	0
Issuance of new shares	1,212	129,715	0	130,927
Payments for warrants	226	12,030	0	12,256
Issuing expenses	0	-22,558	0	-22,558
Conversion of debt	117	8,471	0	8,588
Vesting of share-based compensation	0	284	0	284
Profit-loss for the period	0	0	-4,897	-4,897
Total equity as of December 31st, 2020	3,067	171,502	-5,210	169,359

# Shareholder information

ExpreS<sup>2</sup>ion Biotech Holding AB's share was listed at Nasdaq First North Growth Market on July 29, 2016. The trading name of the share is EXPRS2 and the ISIN-code is SE0008348262. As of June 30, 2020, the number of shares in ExpreS<sup>2</sup>ion Biotech Holding AB amounted to 29,383,191. The average amount of shares in the second quarter of 2021 amounted to 28,739,550. The Company has one class of shares. Each share carries equal rights to share in the Company's assets and earnings.

#### **Certified Advisor**

Svensk Kapitalmarknadsgranskning AB

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#### List of largest shareholders

Name	Number of shares held	Share of votes and capital
Summary, shareholders over 5%	0	0.00%
Remaining shareholders under 5%	29,383,191	100.00%
Total June 30, 2021	29,383,191	100.00%

## Warrants

As of June 30<sup>th</sup>, 2021, the Company had three series of warrants issued, where two series are part of incentive programs and one series is part of the rights issue of units carried out during the fourth quarter 2020. These series are identified as T02, T05 and T06.

#### TO2 (2019/2022)

On May 23, 2019, the Annual General Meeting resolved to implement an incentive program for all employees and issue a maximum of 680,100 warrants, of which 612,084 were subscribed for and allocated to the employees.

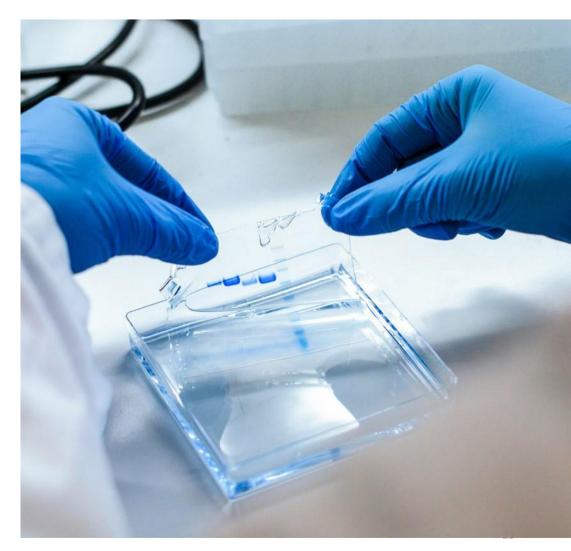
#### TO4 (2020/2021-1) and TO5 (2020/2021-2)

On September 23, 2020, the Extraordinary General Meeting resolved to approve the board resolution from August 25, 2020 regarding a rights issue of units consisting of shares and two series of warrants of 5,455,297 warrants each. All shares and warrants issued in the rights issue of units were subscribed and paid for by the subscribers. On April 28, ExpreS<sup>2</sup>ion announced the outcome of the exercise of warrants of series TO4. In total, 5,324,670 warrants of series TO4 were exercised, corresponding to approximately 97.6

percent of the total number of outstanding warrants of series TO4, for subscription of 1,774,890 shares at an exercise price of SEK 22.00 per share. ExpreS²ion received approximately SEK 39.0 million before issuing costs through the exercise of the warrants of TO4. The warrants of series TO5 will have a subscription period in September, through which the company hopes to raise up to SEK 45 million.

#### T06 (2020/2024)

On September 23, 2020, the Extraordinary General Meeting resolved to implement an incentive program for management and key persons and issue a maximum of 1,000,000 warrants. All warrants were subscribed for by the Company's subsidiary ExpreS<sup>2</sup>ion Biotechnologies ApS. 998,000 warrants have subsequently been transferred to selected employees and 2,000 warrants are still held by the subsidiary.



## Other matters

#### Employees

As of June 30, 2021, there were a total of 22 employees, corresponding to 20 full-time equivalents (FTE's).

#### Operational risks and uncertainties

The risks and uncertainties that ExpreS²ion's operations are exposed to are summarized in terms of pharmaceutical development, competition, technology development, patents, government requirements, capital requirements, currencies, and interest rates. During the current period, no significant changes regarding risk or uncertainty factors have occurred. For more detailed reporting of risks and uncertainties refer to the Company's annual report for the fiscal year of 2020.

#### Auditor review

This interim report has not been formally reviewed by the Company's auditor.

#### Accounting principles

ExpreS<sup>2</sup>ion Biotech Holding AB applies the Swedish Annual Accounts Act and Swedish Accounting Standards Board's general standard BFNAR 2012:1 (K3) when preparing its financial statements.

#### For more information, please contact

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#### Financial calendar

November 18, 2021	Interim report, Q3 2021
February 24, 2022	Year-end report, 2021



# Declaration of The Board of Directors and CEO

The Board of Directors and CEO assure that the interim report presents a true and fair view of ExpreS<sup>2</sup>ion Biotech Holding AB's business, operations, position and results.

Hørsholm, Denmark August 19, 2021

ExpreS<sup>2</sup>ion Biotech Holding AB c/o Mindpark, Rönnowsgatan 8c, S-252 25 Helsingborg

Board of Directors and CEO

