Q3

Alligator Bioscience AB (publ)
Interim report January - September 2022

Financial Results and Business Update

"I am proud of the work the Alligator team has undertaken to secure this positive momentum through another quarter of progress. We are in a strong position to achieve our next milestones, even earlier than expected, and I am looking forward to adding further to our understanding of the potential of mitazalimab."

Søren Bregenholt
CEO Alligator Bioscience AB (publ)

Significant Events: July – September 2022

Additional data from mitazalimab clinical phase 1b/2 study OPTIMIZE-1 were presented at the AACR Special Conference on Pancreatic Cancer

Alligator announced presentation of data from the Phase 1b dose escalation part of the OPTIMIZE-1 study, which importantly confirmed the pharmacological activity of mitazalimab in combination with mFOLFIRINOX and was found to be safe and well tolerated. Patient enrollment for Phase 2 is now ongoing at sites in Europe, with the primary endpoint to show efficacy in patients with pancreatic cancer as per Response Evaluation Criteria in Solid Tumors (RECIST)-defined overall response rate (ORR).

ALG.APV-527 received FDA green light to initiate clinical Phase 1 in the US

In September, Alligator and Aptevo Therapeutics announced that the US Food and Drug Administration (FDA) had issued a "may proceed" notification for the ALG.APV-527 investigational new drug application (IND), and the companies are now working further to initiate a multi-center Phase 1 trial in the US.

ATOR-1017 900 mg Phase 1 dose escalation study in solid tumors fully enrolled

Data show that ATOR-1017 is safe and well tolerated at doses up to 900 mg with stable disease as the best tumor response, confirming previously announced signs of clinical benefit. This Phase 1 dose escalation study is fully enrolled and has successfully fulfilled its purpose. Alligator will now seek a partner to support the continued development of ATOR-1017.

Alligator intensified presence at leading scientific and medical conferences

As a testament to Alligator's CD40 expertise, senior staff from the company were invited to present at the World Bispecific Summit in Boston and Immuno UK in London. At the former event, Laura von Schantz, VP Discovery, presented "Kick-Starting Tumor Specific T Cell Cross-Priming Using Neo-X-Prime Bispecific Antibodies Targeting CD40", and in London, Peter Ellmark, CSO, presented "Targeting the Myeloid Population In The Tumour Microenvironment".

Financial summary

Figures in brackets refer to the outcome for the corresponding period in the preceding year.

July – September 2022

- Net sales, SEK 5.1 million (3.3)
- Operating profit/loss, SEK -51.4 million (-37.7)
- Profit/loss for the period, SEK -51.4 million (-37.2)
- Earnings per share before and after dilution, SEK -0.23 (-0.45)*
- Cash flow for the period, SEK -45.6 million (-30.4)
- Cash and cash equivalents, SEK 147.4 million (79.3)

Jan – September 2022

- Net sales, SEK 15.6 million (7.7)
- Operating profit/loss, SEK -140.3 million (-104.7)
- Profit/loss for the period, SEK -140.2 million (-104.9)
- Earnings per share before and after dilution, SEK -0.64 (-1.26)*
- Cash flow for the period, SEK -131.0 million (-24.0)
- Cash and cash equivalents, SEK 147.4 million (79.3)

The information was submitted for publication at 8:00 a.m. CET on October 20, 2022. For contact details, see page 9.

* Previous periods have been adjusted, see Note 8, page 20.

CEO Comments

Alligator Bioscience continued to secure the upcoming delivery of key data on its lead assets in Q3 2022 and intensified the promotion of its clinical data to the scientific and investor community. Alligator also continued to strengthen its relationships with its partners, adding significant opportunities to its long-term pipeline and providing further validation of the robustness of its technology platforms.

The Alligator team has worked with dedication through the last months to continue to deliver on our commitment to developing meaningful therapies for patients with hard-to-treat cancer while creating value for our stakeholders and shareholders. In particular, we continued the strong progress with our lead asset, mitazalimab, through clinical development and further underlined the potential of our third-generation proprietary technology, Neo-X-Prime™.

The OPTIMIZE-1 Phase 2 study evaluating the efficacy and safety of mitazalimab in combination with standard-of-care chemotherapy, mFOLFIRINOX, in patients with first-line metastatic pancreatic cancer has experienced a significant acceleration of patient recruitment over the summer. Together with our decision to prioritize mitazalimab, and increase the number of study sites, top line data is now due in Q1 2024, nine months sooner than originally anticipated. This accelerated clinical development is a testament to the operational excellence of the Alligator team and of the interest the medical community has for mitazalimab in this very difficult to treat solid tumor.

In the meantime, OPTIMIZE-1 is progressing at the recommended dosing level of 900 µg/kg and enrollment is ongoing at multiple sites in Europe. We continue to expect interim data collection in Q4 2022 with the publication of these results due towards the end of the year or in the first weeks of January 2023. We are excited to see these first results from the trial and will use the potential learnings from these interim data to optimize the design of the upcoming OPTIMIZE-2 clinical trial.

We have also intensified the promotion of our clinical data to medical conferences and have received significant interest from the scientific community as well as from potential partners.

We reported promising Phase 1 data on our second product, ATOR-1017, a tumor-directed therapy for advanced and metastatic solid cancers. The data show that ATOR-1017 is safe and well tolerated at doses up to 900 mg with stable disease as the best tumor response, confirming previously announced indications of clinical benefit.

These data further differentiate ATOR-1017 from other 4-1BB antibodies, which have not achieved sufficient efficacy or have shown unacceptable side effects. It builds on results presented at the American Society of Clinical Oncology Annual Meeting earlier this year, which confirmed ATOR-1017's mechanism-of-action and showed a strong safety profile and encouraging signs of clinical benefit.

Overall, data clearly indicate ATOR-1017's potential to be best-in-class and to address a significant unmet medical need in patients with advanced malignancies. This Phase 1 study is now fully enrolled and has fulfilled its purpose. We have successfully created a strong foundation for the further clinical development of ATOR-1017, which we believe will be well-positioned to make a clinical difference. Alligator will now look for a partner to support the continued development of ATOR-1017.

This quarter, we also received investigational new drug (IND) authorization from the US Food and Drug Administration for our third drug candidate, ALG.APV-527, which we are developing in partnership with Aptevo Therapeutics. This is an important milestone which allows us to move towards the initiation of a multi-center Phase 1 trial in the US to evaluate ALG.APV-527 in the treatment of 5T4-expressing tumor antigens in multiple solid tumor types. Preclinical data suggest the 4-1BB and tumor-binding antibody has a wide therapeutic window with potential across a range of 5T4-expressing cancers with high unmet need, so we are excited to get clinical evaluation underway as soon as possible.

This strong clinical progress is an important validation of our technology and its ability to produce differentiated drug candidates with the potential to improve treatment options for cancer patients.

During Q3 we have intensified business development discussions on selected assets and technology partnerships, and I look forward to see the results of these efforts in the months to come.

I am proud of the work the Alligator team has undertaken to secure this positive momentum through another quarter of progress. We are in a strong position to achieve our next milestones and I am looking forward to adding further to our understanding of the potential of mitazalimab. We will, of course, continue to keep you up to date on our latest news as we continue to create value also for you, our shareholders and partners.

Søren Bregenholt
CEO Alligator Bioscience AB (publ)

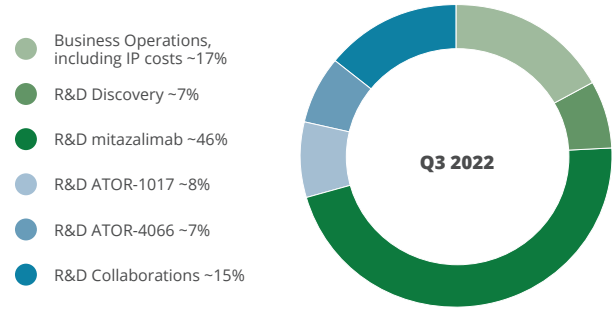
Performance measures Group

	Note	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Result (KSEK)						
Net sales	5	5,108	3,300	15,634	7,694	12,943
Operating profit/loss		-51,419	-37,679	-140,330	-104,692	-141,565
Profit/loss for the period		-51,365	-37,245	-140,152	-104,946	-141,737
R&D costs		-47,064	-21,771	-124,417	-72,056	-110,123
R&D costs as a percentage of operating costs excl. Impairments, %		83%	53%	79%	64%	70%
Capital (KSEK)						
Cash and cash equivalents at end of period		147,411	79,314	147,411	79,314	278,148
Cash flow from operating activities		-42,742	-28,197	-125,289	-93,461	-127,033
Cash flow for the period		-45,555	-30,400	-131,038	-24,006	174,717
Equity at the end of the period		142,256	85,029	142,256	85,029	282,273
Equity ratio at the end of the period, %		69%	65%	69%	65%	85%
Info per share (SEK)						
Average number of shares		220,584,878	83,173,402	220,584,878	83,173,402	89,670,050
Earnings per share before and after dilution*		-0.23	-0.45	-0.64	-1.26	-1.58
Equity per share before and after dilution*		0.64	0.99	0.64	0.99	1.28
Personnel						
Number of employees at end of period		55	46	55	46	46
Average number of employees		52	45	51	44	45
Average number of employees employed within R&D		43	35	42	38	38

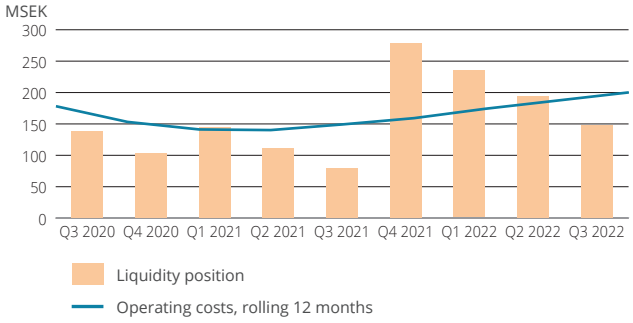
*Effect from dilution is not considered when result is negative and options where call rate is higher than closing rate is not considered.

For definitions and calculations, see the sections later in this report.

Operating costs distributed by function, Parent Company



Operating costs, rolling 12 months and Liquidity (MSEK), Group



Operations

Alligator Bioscience is a clinical stage biotech company developing tumor-directed best-in-class antibodies for hard-to-treat cancers. Our drug candidates have the potential to meet key needs in immuno-oncology by increasing the quantity and quality of tumor specific T cells within the tumor and at the same time remodel the tumor microenvironment making the tumor more inflamed. Alligator's high demands on safety and efficacy of our drug candidates increase their potential to be able to be combined with current standard therapies of cancer, which is highly important for improving treatment results in oncology today.

In 2022, the Company has kept its focus on the two prioritized drug candidates mitazalimab and ATOR-1017. Our technology platforms and pharmaceutical research continue to build long-term value. To drive competitive and time-efficient development, parts of Alligator's work is conducted in collaboration with other biotechnology companies, contract laboratories and leading international research institutions. Our clinical studies are carried out in collaboration with leading specialist physicians and CROs with expertise in oncology clinical development. In summary, the Company has all necessary expertise to pursue successful projects from concept to clinical development.

Alligator's Organization

Alligator's research and development organization is divided into five units: Discovery, CMC (Chemistry, Manufacturing & Control), Non-Clinical Development, Medical Science and Clinical Operations. Members of all these functions collaborate cross-functionally in project teams. The Discovery unit is responsible for early-stage research projects up until a drug candidate has been identified. This normally includes the development and evaluation of treatment concepts, the evaluation of potential drug candidates and early-stage confirmation of efficacy. The CMC unit develops manufacturing processes and is responsible for clinical trial material manufacturing. The Non-Clinical Development unit is responsible for pre-clinical evaluation of safety and efficacy of our molecules, including preparation of the data packages required for clinical trial applications. The Medical Science unit, led by our Chief Medical Officer is responsible for designing all the clinical and regulatory development plans required to show that Alligator's products are safe and effective. The Clinical Operations unit is responsible for timely and excellent implementation of the clinical studies. Alligator will continue to build and shape the organization to match and support its strategy and objectives.

Several Proprietary Technologies

Alligator's technology platforms—FIND® (protein optimization technology), ALLIGATOR-FAB™ and ALLIGATOR-GOLD® (antibody libraries)—are used for the discovery and development of novel drug candidates. These platforms enable efficient generation of novel drug candidates with high potential.

In addition, the Company has bispecific antibody formats for the development of new dual-action antibodies. With the most recent antibody format, RUBY™, Alligator can generate bispecific molecules from any two antibodies, with excellent properties in terms of stability and yield. The format eliminates the need for further optimization, enabling Alligator to quickly move drug candidates from preclinical research to clinical development.

Our 3rd generation proprietary platform technology aims at a more personalized immuno-therapy, using CD40-antibodies that instruct the immune system to recognize and attack cancer cells, based on the tumor mutations unique to the individual patient. These antibodies contain one part that binds to tumors and tumor particles and another part that binds to dendritic cells through the CD40 molecule. This interaction between tumor particles and dendritic cells eventually results in a very efficient education and activation of tumor-specific T cells, that subsequently can recognize and destroy the tumor cells.

Business Model that Creates Value Across the Development Chain

The Company's business model is based on proprietary drug development. To maximize the value of the portfolio, the Company intends to bring molecules from drug discovery and preclinical studies to demonstration of Proof-of-Concept in human clinical Phase 2 trials and beyond. To generate income, limit portfolio risk, and maximize long-term value, the Company seeks strategic global and regional partnerships for certain programs and technologies.

Immuno Oncology Market Overview

Cancer touches all our lives, either directly or through its effect on family and loved ones. With the continued rise of cancer diagnoses around the world, the need for more effective treatments also grows. Alligator's drug candidates are designed with an optimal efficacy-tolerability balance to meet the need for therapies that can safely be combined with current standard cancer treatments, to treat, or possibly even cure, cancers.

Oncology Market Trends

In 2020, 19.3 million new cancer cases were diagnosed globally, with the number expected to rise to 30.2 million by 2040,[1] and the oncology drug market is expected to almost double by 2026 reaching a total of USD 460 billion.[2] A surge of new and innovative treatment methods is expected to emerge in the marketplace, and immunotherapies will play an important role in these treatment options for cancer.

Alligator believes that the demand for novel immuno-therapy drugs will increase along with the global demand for new and more effective oncology therapies.

Immuno-oncology

Most tumors contain immune cells with the potential to attack and destroy cancer cells, and possibly eradicate the entire tumor itself. Cancer cells often activate immunosuppressive strategies to inhibit these types of attacks. Immunotherapies provide several different opportunities to help the immune system defend the body against the cancer. Such strategies could be to educate the immune system to better identify tumor cells, while others aim to enhance the capabilities of the immune system to attack the tumor with full force.

Alligator's innovative assets and technologies target key immune-oncology molecules to educate and activate the immune system to selectively attack tumors without affecting the rest of the body, a core concept that separates us from other competitors in the industry. The main benefit of tumor-directed treatment is the ability to effectively attack the tumor while minimizing the adverse effects caused by stimulating the whole immune system. This allows our candidates to work synergistically with current chemotherapy regimens and other immunotherapeutic drugs in hard-to-treat, metastatic solid tumors.

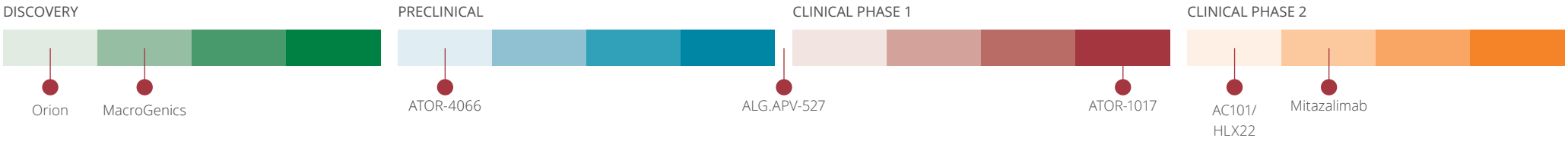
Our lead asset mitazalimab is in a clinical Phase 2 study for the treatment of metastatic pancreatic tumors, a tumor type that is one of the hardest cancers to treat and has one of the lowest five-year survival rates.

Roughly 40,000 people in the United States and about 70,000 in Europe are diagnosed with pancreatic cancer each year. Only 15-20 percent of those diagnosed can be treated by surgery, and there are few treatment options available for the remaining 85 percent, with chemotherapy regimens being the standard of care.[3]

We develop our pipeline programs, from Discovery Phase through clinical Phase 2, with an excellent efficacy-tolerability balance in mind, either alone or in collaboration. These collaborations provide an opportunity of income, and an external validation of our platform, building on our confidence that our candidates will provide meaningful treatment options for people with hard-to-treat cancer, as stand-alone or combination therapies.

[1] International Agency for Research on Cancer (IARC), Cancer Tomorrow. 30 March 2022.
[2] Database GlobalData (Pharma Intelligence Center – Drug Sales), September 2021.
[3] Database GlobalData (Pancreatic Cancer – Opportunity Analysis and Forecasts to 2029), December 2020.

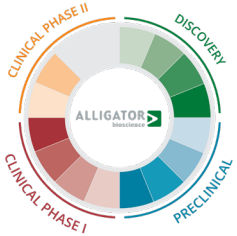
PIPELINE PROJECTS



Pipeline Projects – Internal Programs

Alligator's competitive project portfolio consists of the two clinical-stage assets, mitazalimab and ATOR-1017, and ATOR-4066, a pre-clinical program developed using Alligator's proprietary technology platform Neo-X-Prime™ – as well as several programs developed in collaboration with partners.

Mitazalimab



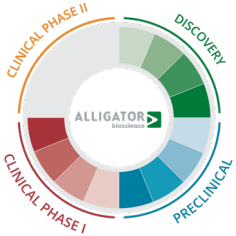
Alligator's most advanced drug candidate mitazalimab, a potential game changer in the treatment of solid tumors, entered a Phase 2 clinical trial in pancreatic cancer, with the first patient dosed in the OPTIMIZE-1 study in Q3 2021. This clinical trial is designed to evaluate the safety and efficacy of mitazalimab in combination with mFOLFIRINOX, the most efficacious standard of care chemotherapy for the treatment of advanced pancreatic cancer.

The clinical trial has been designed on the principal that mFOLFIRINOX efficiently kills tumor cells, leading to an increased release of tumor antigens which, when mFOLFIRINOX is used as standalone therapy, only triggers minimal immune response, leading to limited overall efficacy. The use of mitazalimab in combination with mFOLFIRINOX allows for activation of CD40, a receptor on dendritic cells, leading to significantly improved tumor antigen-presentation and subsequent activation of tumor-specific T cells that attack the cancer. The combination of mitazalimab with mFOLFIRINOX is therefore expected to significantly boost the immune response secondary to the release of tumor antigens, hence triggering powerful attack on the solid tumors.

Mitazalimab has previously undergone two Phase 1 clinical trials, one conducted by Alligator, and one conducted by Janssen Biotech Inc., both of which showed strong evidence of efficacy and proof-of-mechanism, as well as a manageable safety profile.

Subsequent to the 450 µg/kg dose cohort of mitazalimab in combination with mFOLFIRINOX showing good safety, Alligator announced in Q1 2022 that also 900 µg/kg was safe in combination with mFOLFIRINOX. The study is progressing better than initially anticipated with the recruitment of patients in the 900 µg/kg dose level having accelerated over the summer 2022. As a consequence, Alligator announced that the full Phase 2 data would become available in Q1 2024, 9 months earlier than initially expected. In the meantime, Alligator is still expecting an interim efficacy readout for OPTIMIZE-1 with data becoming available towards the end of Q4 2022 and a potential read-out in the first weeks of January at the latest.

ATOR-1017



ATOR-1017 is Alligator's second most advanced program and is in the final stages of a Phase 1 dose-escalation study. The study is designed to assess the safety and tolerability of ATOR-1017 in patients with advanced, solid cancers, and to establish a recommended Phase 2 dose for future studies.

ATOR-1017 is a 4-1BB agonist with a unique profile, most importantly through its ability to enhance the immune activating effect in tumors. This creates opportunities for a powerful, tumor-directed immune activation, which can increase the therapeutic effect and reduce adverse side effects for patients.

Clinical data generated to date have shown a favorable pharmacokinetic profile and proof-of-mechanism biomarker responses. In Q3 2022, Alligator announced the Phase 1 dose escalation study fully enrolled, and that is successfully had fulfilled it's purpose, where doses of up to 900 mg were shown safe and well tolerated, with stable disease as best tumor response.

ATOR-4066

ATOR-4066 is a bispecific antibody created to elicit powerful, patient-specific anti-tumor effects, developed using Alligator's technology platform, Neo-X-Prime™.

ATOR-4066 targets two molecules: the same CD40 receptor on dendritic cells that mitazalimab targets, as well as carcinoembryonic antigen (CEA), a protein located on tumor cells. Thanks to its bispecificity, ATOR-4066 has an exceptional ability to induce cross-priming of tumor specific T cells, resulting in very efficient tumor killing.

Early data for ATOR-4066 have shown significantly higher preclinical anti-tumor efficacy compared to a corresponding monospecific CD40 antibody. Preclinical studies have also shown that ATOR-4066 results in prolonged T cell-mediated anti-tumor response.

In 2022, Alligator aims to initiate IND-enabling development of ATOR-4066.

Collaborations and Out-Licensing Agreements

Aptevo Therapeutics, Inc.

ALG.APV-527 is a bispecific antibody that targets the 4-1BB and 5T4 molecules, designed for the treatment of metastatic cancer. In 2017, Aptevo Therapeutics and Alligator Bioscience AB signed a co-development agreement. Under the agreement, both companies will equally own and finance the development.

The original molecules involved in the tumor-binding function and the immunomodulatory function of ALG.APV-527 were developed using Alligator's patented ALLIGATOR-GOLD® antibody library. The bispecific molecule was further developed and improved with Aptevo's technology platform ADAPTIR™. By combining a tumor-binding function with an immunomodulatory function in the same molecule, the drug candidate selectively targets the tumor and stimulates the antitumor-specific immune cells that are present in the tumor.

In September 2022, US FDA cleared the IND for 527 allowing A&A to initiate phase clinical trials in the US.

Orion Corporation

In 2021, Alligator entered a research collaboration and license agreement with Orion Corporation, a global pharmaceutical company based in Finland. The aim of the collaboration is to discover new bispecific antibody cancer therapeutics against immuno-oncology targets. The agreement covers an option to develop three bispecific antibodies.

Under the agreement, Alligator will employ its proprietary phage display libraries and its RUBY™ bispecific platform. During the initial research period of the collaboration, Alligator will receive an upfront payment and reimbursement of research costs and other fees.

As part of the agreement, Alligator is eligible for development, approval, and sales milestone payments of up to EUR 469 million. Should Orion exercise its option to continue development and commercialization of the resulting product candidates, Alligator will be eligible to receive additional royalty payments.

MacroGenics, Inc.

In 2021, Alligator entered a joint research collaboration with US-based MacroGenics, Inc., a Nasdaq-listed biopharmaceutical company focused on developing and commercializing innovative monoclonal antibody-based therapeutics for the treatment of cancer. The research collaboration utilizes Alligator's proprietary myeloid engaging Neo-X-Prime™ platform to develop bispecific antibodies against two undisclosed targets.

Under the joint research collaboration agreement, which covers activities from candidate drug generation up until IND-enabling studies, each company will be responsible for its own costs. The parties may continue further development of the resulting bispecific molecule under a separate co-development collaboration and licensing agreement.

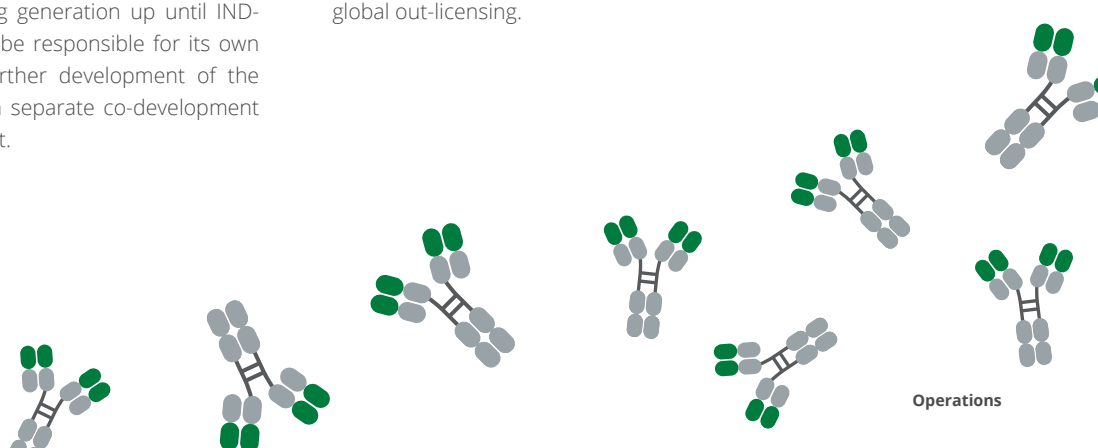
Biotheus

In 2019, an agreement was concluded with Chinese company Biotheus, where Biotheus obtained the Chinese rights (Greater China, Hong Kong, Taiwan and Macao) to an antibody from the ALLIGATOR-GOLD® antibody library. The agreement gives Alligator the right to total initial upfront payments, as well as milestone and option payments of potentially USD 142 million. To date, Alligator has received upfront payments of about SEK 10 million.

Abclon

Through its subsidiary Atlas Therapeutics AB, Alligator holds a participating interest in the clinical Biosynergy (AC101/HLX22) project, run by the listed Korean company AbClon. The drug candidate is now being further developed by Chinese company Shanghai Henlius, which increased its rights to encompass a global license for development and commercialization in 2018. The phase 2 study had the first patient dosed in Q4 2021.

Alligator incurs no cost for this project and is entitled to 35% of AbClon's revenue from out-licensing to Shanghai Henlius. In previous financial years, Alligator received two milestone payments totaling USD 3 million in conjunction with regional and global out-licensing.



The Alligator Share

Number of shares, stock option program and share saving program

The total number of outstanding shares in the Company at the end of the quarter was 221,534,728 (220,584,878), of which 220,584,878 are ordinary shares with one vote per share and 949,850 are series C shares with one-tenth of a vote per share. The number of votes in the company amounts to 220,679,863 votes

Share saving program LTI 2021

At the annual general meeting 2021 it was resolved to implement a long-term incentive program by way of a performance-based share saving program for employees in the company ("LTI 2021"). For each ordinary share acquired by the participant on Nasdaq Stockholm, so called saving shares, the participant has a right to receive so called matching shares. In addition, given that a requirement related to the development of the company's share price from the day of the annual general meeting 2021 up until 30 September 2024 has been achieved, the participant has a right to receive further shares in the company free of charge, so called performance shares. After recalculation due to a completed rights issue in 2021, each saving share entitles to 1.0947 matching shares. The thresholds for the receipt of one, two or four performance shares per saving share amounts to SEK 15.74 for receipt of one performance share, SEK 31.65 for receipt of two performance shares and SEK 52.89 for receipt of four performance shares.

The maximum number of ordinary shares that can be issued in relation to LTI 2021 amount to 949,850, whereby 722,759 for the deliverance of matching shares and performance shares to participants and 227,091 to hedge payments of future social security contributions, which corresponds to a dilution of approximately 0.4 per cent of the company's share capital and votes.

Warrant programs, LTI 2022 I/II

At the annual general meeting 2022 it was resolved to implement a long-term incentive program by way of a warrant program for employees in the company ("LTI 2022-I"). In case all warrants issued within the warrant program LTI 2022-I are utilized for subscription of new ordinary shares, a total of 3,700,000 new ordinary shares will be issued, which corresponds to a dilution of approximately 1.65 per cent of the company's ordinary shares after full dilution.

The annual general meeting 2022 also resolved to adopt a warrants program for certain board members of the company, (LTI 2022-II"). In case all warrants issued within this program are utilized for subscription of new ordinary shares, a total of 600,000 new ordinary shares will be issued, which corresponds to a dilution of approximately 0.27 per cent of the company's ordinary shares after full dilution.

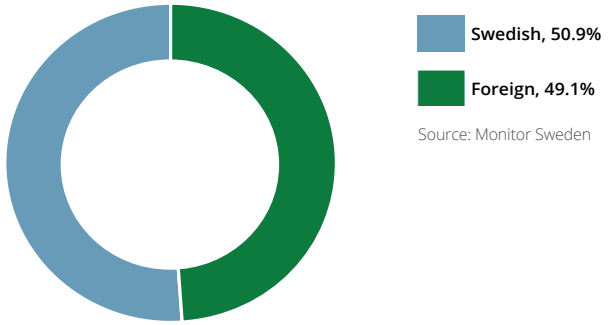
Each warrant in LTI 2022-I/II entitle to subscription of one ordinary share in the company. Subscription of shares by virtue of the warrants may be effected as from 1 June 2025 up to and including 30 June 2025. The subscription price per share for above warrant programs, was calculated to SEK 3,38 which corresponds to 200 per cent of the volume weighted average price during 10 trading days immediately after the annual general meeting 2022. All warrants will be transferred to the participants at fair market value.

In case the existing share saving program as well as both warrant programs are exercised in full, a total of 5,249,850 new shares will be issued, which corresponds to a total dilution of approximately 2,32 percent.

The Alligator share in brief September 30, 2022

Listed on:	Nasdaq Stockholm Small Cap
Number of shares:	221,534,728 (220,584,878 ordinary shares and 949,850 C shares)
Average turnover per day:	Approximately 159,000 (preceeding quarter: approx. 155,000)
Number of shareholders:	Approx. 8,600 (preceeding quarter: approx. 8,600)
Market capitalization:	MSEK 331 (preceeding quarter: approx. MSEK 322)
Ticker:	ATORX
ISIN:	SE0000767188

Swedish and foreign ownership



Source: Monitor Sweden

Largest Shareholders, Sep 30, 2022	No of Shares	%
Allegro Investment, Inc.	55,643,092	25.1
Omentum SA	13,609,162	6.1
Lars Spånberg	9,641,572	4.4
4 AP-fund	6,819,547	3.1
Magnus Petersson	6,228,326	2.8
Sunstone Capital	5,758,485	2.6
Avanza pension	5,549,979	2.5
Nordnet pension	5,136,673	2.3
Roxette Photo NV	4,804,788	2.2
Mikael Lönn	4,326,547	2.0
Remaining share holders	103,066,707	46.9
Total number of shares	**220,584,878**	**100**

The Company's owner structure is updated regularly on the Company's website: www.alligatorbioscience.com
Source: Modular Finance AB. Compiled and processed data from various sources, including Euroclear, Morningstar and the Swedish Financial Supervisory Authority (Finansinspektionen).

Other information

Review
This report has been reviewed by the Company's auditor.

Employees
The number of employees in the Group at the end of the quarter was 55 (46). Of these, 17 (12) were men and 38 (34) were women. Of the total number of employees at the end of the quarter 46 (38) were employed within research and development.

Future report dates
Alligator intends to publish its financial reports according to the following:

- Year-end Report: February 10, 2023
- Annual Report: March 2023
- Q1 Interim Report: April 26, 2023

Risks and uncertainties
During the course of its business operations, the Group is exposed to various financial risks, such as market risk (comprising foreign exchange risk, interest-rate risk and price risk), credit risk and liquidity risk. The aim of the Group's overall risk management is to achieve minimal adverse effects in terms of earnings and financial position. The Group's business risks, risk management and financial risks are described in detail in the Annual report for 2021.

The impact of Covid-19 on the Group's risks
The Covid-19 pandemic has affected the way we work, but currently we do not forsee any negative long-term effects on our operations due to the pandemic.

The impact of the Ukraine's crisis on the Group's risks
The situation in Ukraine is foremost a humanitarian tragedy that is causing great human suffering. The Russian invasion of Ukraine has worsened the political security situation in the rest of the world and created great uncertainty in the financial markets, which may affect the company's ability to finance clinical trials in the future. The company has no direct business in, nor does it conduct any clinical studies in Ukraine or Russia, but see a risk that the company eventually will suffer from increased raw material and energy prices, which are likely to translate into increased prices for goods and services.

Cyber security
Cyber attacks have become a significant threat in society and for Alligator Bioscience, which is dependent on IT support in its daily operations. The company has ongoing work to ensure that the company is well prepared to counter cyber-attacks and other types of intrusion.

Statement of financial position
The Company works continuously to secure the financing of the operation. This include both business development for new partnering agreements, with an upfront payment upon signing, as well as other options. As the company within the next 12 months has additional financing needs that have not yet been secured, the Board is continuously working on evaluating various financing options to ensure continued operation. It is the Board's assessment that the company has good prospects of securing future financing, for example, through a new share issue, however, the absence of assurance at the time of submission of this report means that there is a significant uncertainty factor regarding the company's ability to continue operation.

Forward-looking information
Even though the board and management believe the expectations in this report are justified, no guarantees can be given that they will turn out to be correct. Accordingly, the actual outcome may differ significantly from the assumptions stated in the forward- looking information depending on, among other factors, changes in the economy or market, changes in legal or regulatory demands, political decisions and changes in exchange rates.

Parent Company
Both Group management functions and all operating activities are carried out in the Parent Company. For additional details, refer to the information provided for the Group since the subsidiaries do not conduct their own operations.

Notes to the reader
Figures in brackets refer to the outcome for the corresponding period in the preceding year for figures related to the income statement and cash flow. For figures related to the financial position and personnel, figures in brackets refer to December 31, 2021. Unless otherwise stated, all amounts stated are rounded correctly, which may mean that some totals do not tally exactly. "Dollar" means US dollars unless otherwise stated.

Registered trademarks
FIND® and ALLIGATOR-GOLD® are Alligator Bioscience AB proprietary trademarks which are registered in Sweden and other countries.

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Financial statements

Unless otherwise stated in this Interim report, numbers refer to the Group. Due to the nature of the business, there can be large fluctuations in revenue which are not seasonal or regular but are mainly linked to when milestones generating a payment are reached in out-licensed research projects. Like revenue, expenses can also fluctuate between periods. Among other factors, this fluctuation in expenses is influenced by the current phase of the various projects since certain phases generate higher costs. Figures in brackets refer to the outcome for the corresponding period in the preceding year for figures related to the income statement and cash flow. For figures related to the financial position and personnel, figures in brackets refer to December 31, 2021. Unless stated otherwise, all amounts are in SEK thousand (KSEK). All amounts stated are rounded, which may mean that some totals do not tally exactly.

Consolidated
Income Statement

Net Sales
Sales for the period pertain primarily to the collaboration and licence agreement with Orion Corporation. In the same period prior year sales reffered primarily to the collaboration and licence agreement with Orion Corporation and to the joint research agreement with BioArctic AB.

Other operating income
Other operating income for the quarter for both year 2022 and 2021 comprises primarily of exchange gains in the company's operations.

Operating costs
The company's costs are higher compared to the same period previous year, and pertain mainly to costs related to the clinical projects mitazalimab and its study OPTIMIZE-1 and ALG.APV-527. External costs for mitazalimab amounted to SEK 18,478 thousand (3,706) during the third quarter of the year and the increased costs are mainly related to the increased number of patients in the study. In September 2022, an IND (Investigational New Drug) -application for ALG.APV-527 was approved, which enables the initiation of clinical studies which generates higher costs. The personnel costs in the third quarter is higher than last year due to an increased number of employees.

Net financial items
Pertains to unrealized exchange gains and losses as a result of liquidity positions in USD, EUR and GBP.

All amounts KSEK unless specified	Note	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Operating income						
Net sales	5	5,108	3,300	15,634	7,694	12,943
Other operating income	5	130	50	597	367	2,183
Total operating income		**5,238**	**3,350**	**16,231**	**8,061**	**15,126**
Operating costs						
Other external costs		-39,446	-24,539	-100,347	-62,171	-86,982
Personnel costs		-15,416	-13,414	-49,661	-41,705	-57,814
Depreciation of tangible assets and intangible assets		-1,285	-2,941	-5,375	-8,549	-11,144
Other operatings expenses		-511	-134	-1,178	-328	-751
Total operating costs		**-56,658**	**-41,029**	**-156,561**	**-112,753**	**-156,691**
Operating profit/loss		**-51,419**	**-37,679**	**-140,330**	**-104,692**	**-141,565**
Financial items						
Other interest income and similar income statement items		26	-	151	-9	-2
Interest expense and similar income statement items		28	434	27	-245	-169
Net financial items		**54**	**434**	**178**	**-253**	**-171**
Profit/loss before tax		**-51,365**	**-37,245**	**-140,152**	**-104,946**	**-141,736**
Tax on profit for the period		-	-	-	-	-
Profit for the period attributable to Parent Company share-holders		**-51,365**	**-37,245**	**-140,152**	**-104,946**	**-141,736**
Earnings per share						
Earnings per share before and after dilution, SEK		**-0.23**	**-0.45**	**-0.64**	**-1.26**	**-1.58**

Consolidated
Statement of Comprehensive Income

All amounts KSEK	Note	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Profit/loss for the period		-51,365	-37,245	-140,152	-104,946	-141,736
Other comprehensive income		-	-	-	-	-
Comprehensive income for the period		**-51,365**	**-37,245**	**-140,152**	**-104,946**	**-141,736**

Consolidated
Statement of Financial Position

ASSETS

Participations in development projects
The Group's participations in development projects refers to cooperation with the South Korean company AbClon Inc. for the Biosynergy project. Biosynergy is outlicensed to the Chinese company Shanghai Henlius, which is now further developing the drug candidate. At the end of the period, participations in development projects amounted to SEK 17,949 thousand (17,949).

Right of use assets
At the end of the period, right of use assets amounted to SEK 19,801 thousand (10,456). Right of use assets pertain to leases for offices and laboratories, machines and vehicles. Rights of use assets are higher compared to the previous period and year due to an extension in one of our leasing contracts relating to office rent. The contract should expire on December 31, 2022 but has been extended for another 3 years. Alligator entered into a lease agreement with Medicon Village for office premises valid from October 2024 with an agreement period of 5 years. The new agreement is estimated to increase the right of use assets by SEK 42,281 thousand, based on the use of the agreement period without extension, and replaces the current agreement with Medicon Village regarding office premises.

Cash and cash equivalents
Consolidated cash and cash equivalents, which consist of bank balances, totaled SEK 147,411 thousand (278,148).

The Group plans to use its liquidity for operating activities. A portion of the Group's liquidity is invested in USD, EUR and GBP foreign currency accounts. In accordance with the Group's Financial Policy, inflows of foreign currencies exceeding the expected requirements for the coming 18 months are to be converted to SEK at the time of payment. Besides this, no further hedging has taken place.

All amounts in KSEK	Note	2022-09-30	2021-09-30	2021-12-31
ASSETS				
Fixed assets				
Intangible assets				
Participations in development projects	3	17,949	17,949	17,949
Patents		-	27	17
Softwares		103	234	201
Tangible assets				
Improvements in leased premises		152	761	608
Right of use assets		19,801	13,096	10,456
Equipment, machinery and computers		1,831	5,378	4,355
Total fixed assets		**39,835**	**37,445**	**33,587**
Current assets				
Current receivables				
Accounts receivable	6	6,639	-	7,446
Other receivables	6	4,678	8,991	7,044
Prepayments and accrued income		6,492	6,073	6,975
Cash and cash equivalents	6	147,411	79,314	278,148
Total current assets		**165,219**	**94,377**	**299,613**
TOTAL ASSETS		**205,054**	**131,822**	**333,200**

Consolidated

Statement of Financial Position

EQUITY AND LIABILITIES

Equity

Equity at the end of the period amounted to SEK 142,256 thousand (282,273), corresponding to an equity ratio of 69 (85) %. During the year the number of shares and votes in the Company have increased due to directed issue and repurchase of 949,850 series C shares, which were resolved upon by the board of directors on 22 March 2022 pursuant to the authorization granted by the annual general meeting on 1 June 2021.

As of 30 September 2022, the number of shares in Alligator Bioscience AB amounts to 221,534,728 shares, of which 220,584,878 are ordinary shares with one vote per share and 949,850 are series C shares with one-tenth of a vote per share. The number of votes in the company amounts to 220,679,863 votes.

Equity per share before and after dilution

At the end of the period, equity per outstanding share amounted to SEK 0.64 (1.28), before and after dilution. Since the subscription price for issued options has not been reached, these are not taken into account (not "in-the-money"). C shares are not taken into account either.

Lease liabilities and loans

At the end of the period long- and short-term lease liabilities amounted to SEK 18,945 thousand (9,736). Lease liabilities pertain to leases for offices and laboratories, machines and vehicles. Lease liabilities are higher compared to the previous period due to an extension in one of our leasing contracts relating to office rent. The contract should expire on December 31, 2022 but has been extended for another 3 years. Alligator entered into a lease agreement with Medicon Village for office premises valid from October 2024 with an agreement period of 5 years. The new agreement is estimated to increase the right of use assets by SEK 42,281 thousand, based on the use of the agreement period without extension, and replaces the current agreement with Medicon Village regarding office premises.

Accrued expenses and deferred income

At the end of the period, accrued expenses and deferred income amounted to SEK 38,323 thousand (29,586). Expenses pertains to accrued expenses for clinical activities, personnel and other expenses.

All amounts in KSEK	Note	2022-09-30	2021-09-30	2021-12-31
EQUITY AND LIABILITIES				
Equity				
Share capital		88,614	34,267	88,234
Other capital contributions		911,914	731,765	911,831
Retained earnings and profit/loss for the period		-858,271	-681,003	-717,792
Equity attributable to Parent Company shareholders		**142,256**	**85,029**	**282,273**
Non-current provisions and liabilities				
Lease liabilities	6	11,877	4,222	3,511
Other long-term liabilities	6	-	-	-
Total non-current provisions and liabilities		**11,877**	**4,222**	**3,511**
Current liabilities				
Accounts payable	6	4,362	7,078	9,367
Other liabilities		1,167	1,103	2,237
Lease liabilities	6	7,068	7,295	6,225
Accrued expenses and deferred income	6	38,323	27,096	29,586
Total current liabilities		**50,920**	**42,571**	**47,416**
TOTAL EQUITY AND LIABILITIES		**205,054**	**131,823**	**333,200**

Consolidated

Statement of Changes in Equity, in summary

All amounts in KSEK		2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Opening balance		**193,224**	**122,275**	**282,273**	**115,244**	**115,244**
New capital issue		-	-	380	85,666	359,570
Transaction costs		370	-	-343	-10,931	-50,801
Treasury shares*		-	-	-380	-	-
Warrants**		-	-	426	-	-
Effect of share-based payments personnel		27	-	52	-8	-3
Profit/loss for the period		-51,365	-37,245	-140,152	-104,946	-141,736
Other comprehensive income in the period		-	-	-	-	-
Closing balance		**142,256**	**85,029**	**142,256**	**85,029**	**282,273**

*The item refers to the repurchase of 949,850 C shares that the Board, with the support of authorized members of the Annual General Meeting on June 1, 2021, decided on March 22, 2022.

**The item refers to cash compensation for issuing warrants. For more information on the Warrant Program, see page 8.

Consolidated
Statement of Cash Flows

Investments

No investments were made during the third quarter compared to SEK 45 thousand in the same period last year. Investments during the year consisted of laboratory equipment SEK 293 (45) thousand.

Cash flow for the period

Cash flow for the quarter totaled SEK -45,555 thousand (-30,400) and relates mainly to costs from operating activities.

All amounts in KSEK	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Operating activities					
Operating profit/loss	-51,419	-37,679	-140,330	-104,692	-141,565
Adjustments for items not generating cash flow					
Depreciation and impairments	1,285	2,941	5,375	8,549	11,144
Effect from warrant program	27	-	80	-8	4
Other items, no impact on cash flow	3	27	180	-18	65
Interest received	-	-	-	-	-
Interest paid	-	-53	-126	-192	-235
Tax paid	-	-	-	-	-
Cash flow from operating activities before changes in working capital	**-50,105**	**-34,764**	**-134,821**	**-96,361**	**-130,587**
Changes in working capital					
Change in operating receivables	-1,862	-2,987	567	-8,061	-13,589
Change in operating liabilities	9,225	9,554	8,965	10,961	17,144
Cash flow from operating activities	**-42,742**	**-28,197**	**-125,289**	**-93,461**	**-127,033**
Investing activities					
Acquisition of tangible assets	-	-45	-293	-45	-45
Cash flow from investing activities	**-**	**-45**	**-293**	**-45**	**-45**
Financing activities					
Amortization of leasing liabilities	-3,183	-2,083	-5,435	-5,010	-6,672
Amortization of installment purchase	-	-75	-104	-226	-301
New share issue*	380	-	380	85,666	342,665
Transaction costs*	-10	-	-343	-10,931	-33,897
Option premiums received	-	-	426	-	-
Purchase of treasury shares	-	-	-380	-	-
Cash flow from financing activities	**-2,813**	**-2,159**	**-5,456**	**69,499**	**301,795**
Cash flow for the period	**-45,555**	**-30,400**	**-131,038**	**-24,006**	**174,718**
Cash and cash equivalents at beginning of period	**192,913**	**109,705**	**278,148**	**103,342**	**103,342**
Exchange rate differences in cash and cash equivalents	53	9	300	-22	60
Cash and cash equivalents at end of period	**147,411**	**79,314**	**147,411**	**79,314**	**278,148**

*Refers to the share issue of C shares that was carried out in Q1 2022.

Parent Company
Income Statement

All amounts in KSEK	Note	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Operating income						
Net sales	5	5,108	3,300	15,634	7,694	12,943
Other operating income	5	130	50	597	367	2,183
Total operating income		**5,238**	**3,350**	**16,231**	**8,061**	**15,126**
Operating costs						
Other external costs		-39,660	-26,359	-102,450	-67,036	-93,279
Personnel costs		-15,416	-13,414	-49,661	-41,705	-57,814
Depreciation and impairment of tangible assets and intangible assets		-1,083	-1,190	-3,389	-3,865	-5,084
Other operatings expenses		-511	-134	-1,178	-328	-751
Total operating costs		**-56,671**	**-41,097**	**-156,678**	**-112,934**	**-156,928**
Operating profit/loss		**-51,432**	**-37,747**	**-140,447**	**-104,874**	**-141,802**
Results from financial items						
Other interest income and similar income statement items		26	-	151	-9	-2
Interest expense and similar income statement items		27	-34	146	-71	39
Net financial items		**53**	**-34**	**297**	**-80**	**37**
Profit/loss after financial items		**-51,379**	**-37,782**	**-140,150**	**-104,954**	**-141,765**
Result before tax		**-51,379**	**-37,782**	**-140,150**	**-104,954**	**-141,765**
Tax on profit for the year		-	-	-	-	-
Profit/loss for the period		**-51,379**	**-37,782**	**-140,150**	**-104,954**	**-141,765**

Parent Company
Statement of Comprehensive Income

All amounts in KSEK	Note	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Profit/loss for the period		-51,379	-37,782	-140,150	-104,954	-141,765
Other comprehensive income		-	-	-	-	-
Profit/loss for the year		**-51,379**	**-37,782**	**-140,150**	**-104,954**	**-141,765**

Parent Company
Balance Sheet

ASSETS

All amounts in KSEK	Note	2022-09-30	2021-09-30	2021-12-31
ASSETS				
Fixed assets				
Intangible assets				
Patents		-	27	17
Software		103	234	201
Total intangible assets		**103**	**261**	**219**
Tangible assets				
Improvements in leased premises		152	761	608
Equipment, machinery and computers		1,831	5,378	4,355
Total tangible assets		**1,983**	**6,138**	**4,963**
Financial assets				
Participations in Group companies	3	20,294	20,294	20,294
Total financial assets		**20,294**	**20,294**	**20,294**
Total fixed assets		**22,379**	**26,694**	**25,475**
Current assets				
Current receivables				
Accounts receivables		6,639	-	7,446
Receivables from Group companies		438	438	438
Other receivables		4,677	8,991	7,044
Prepayments and accrued income		8,313	7,894	8,796
Total current receivables		**20,067**	**17,323**	**23,724**
Cash and bank deposits		146,130	78,450	277,288
Total current assets		**166,196**	**95,773**	**301,012**
TOTAL ASSETS		**188,576**	**122,467**	**326,488**

Parent Company
Balance Sheet

EQUITY AND LIABILITIES

All amounts in KSEK	Note	2022-09-30	2021-09-30	2021-12-31
EQUITY AND LIABILITIES				
Equity				
Restricted equity				
Share capital		88,614	34,267	88,234
Total restricted equity		**88,614**	**34,267**	**88,234**
Non-restricted equity				
Share premium reserve		911,488	731,765	911,831
Retained earnings		-715,941	-573,888	-573,877
Profit/loss for the period		-140,150	-104,954	-141,765
Total non-restricted equity		**55,397**	**52,923**	**196,190**
Total equity		**144,011**	**87,190**	**284,424**
Non-current provisions and liabilities				
Other long-term liabilities		-	178	143
Total non-current provisions and liabilities		**-**	**178**	**143**
Current liabilities				
Accounts payable		4,362	7,078	9,367
Other liabilities		1,167	925	2,095
Accrued expenses and deferred income		39,036	27,096	30,459
Total current liabilities		**44,565**	**35,098**	**41,921**
TOTAL EQUITY AND LIABILITIES		**188,576**	**122,467**	**326,488**

Notes

Note 1 General information

This Interim report covers the Swedish Parent Company Alligator Bioscience AB (publ), corporate registration number 556597-8201, and its subsidiaries Atlas Therapeutics AB, corporate registration number 556815-2424, and A Bioscience Incentive AB, corporate registration number 559056-3663. Group's business operations are mainly carried out in the Parent Company.

The Parent Company is a Swedish public limited liability company registered and domiciled in the Municipality of Lund. The office is located at Medicon Village, SE-223 81 Lund.

Note 2 Accounting policies

This Year End report for the Group has been prepared in accordance with IAS 34 Interim Financial Reporting and applicable regulations in the Swedish Annual Accounts Act (ÅRL). The interim report for the Parent Company has been prepared in accordance with the Swedish Annual Accounts Act (ÅRL) and the Swedish Financial Reporting Board's recommendation RFR 2 Accounting for Legal Entities.

The accounting policies and calculation methods used in this report are the same as those described in the Annual report for 2021.

Note 3 Effects of changed estimates and judgments

Significant estimates and judgments are described in Note 3 and Note 19 of the Annual report for 2021. There have been no changes to the company's estimates and judgments since the Annual report for 2021 was prepared.

Note 4 Segment reporting

The company conducts only one business activity, namely research and development in the field of immunotherapy, and the chief operating decision-maker is thus only responsible for regularly making decisions on and allocating resources to one entity. Accordingly, the company comprises only one operating segment, which corresponds to the Group as a whole, and no separate segment reporting is provided.

Note 5 Consolidated Net Sales

A breakdown of the Group's revenue regarding license revenue as follows:

All amounts in KSEK	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Licensing income	-	2,549	-	4,643	4,643
Reimbursement for development work	5,108	751	15,635	3,051	8,300
Total	**5,108**	**3,300**	**15,635**	**7,694**	**12,943**

A breakdown of the Group's other operating income is as follows:

All amounts in KSEK	2022 Jul-Sep	2021 Jul-Sep	2022 Jan-Sep	2021 Jan-Sep	2021 Jan-Dec
Swedish government grants received	-	6	179	6	384
EU grants received	-	-	6	-	1,251
Operational exchange rate gains	121	44	403	361	547
Other	9	-	9	-	-
Total	**130**	**50**	**596**	**367**	**2,183**

Note 6 Financial instruments

Cash and cash equivalents for the Group at September 30, 2022 consisted of bank balances amounting to SEK 147,411 thousand (278,148). For financial assets and liabilities, the reported value as below is considered a reasonable approximation of fair value.

All amounts in KSEK	2022-09-30	2021-09-30	2021-12-31
Financial assets valued at amortized cost			
Accounts receivable	6,639	-	7,446
Other receivables	-	924	1,823
Liquid assets - Bank accounts	147,411	79,314	278,148
Total financial assets	**154,050**	**80,238**	**287,417**
Financial liabilities valued at amortized cost			
Long-term lease liabilities	11,877	4,222	3,511
Other long-term liabilities	-	-	-
Accounts payable	4,362	7,078	9,367
Short-term lease liabilities	7,068	7,295	6,225
Other short-term liabilities	-	178	143
Accrued expenses	33,238	21,457	24,038
Total financial liabilities	**56,545**	**40,230**	**43,285**

Note 7 Related party transactions

The Company had no related party transactions during the first quarter 2022. Until August 31 2021, Alligator had a consulting agreement with former board member Carl Borrebaeck through the company Ocean Capital AB pertaining to expert assistance with the evaluation of early-phase research projects and new antibodies. These related party transactions corresponded to an expense of SEK 480 thousand for the first nine months of the year 2021.

Since 2020 and up until 29 October 2021, Gayle Mills was the Company's Chief Business Officer on a consultant basis in accordance with a consultancy agreement between Alligator and Gayle Mills, and received remuneration based on hours worked. These related party transactions corresponded to an expense of SEK 985 thousand for the first nine months of 2021 and SEK 1,054 thousand during 2021.

Note 8 Correction of error

For the financial year 2021 (comparison year), an error has been noted in the average number of shares before and after dilution. We have stated the number of shares as of the balance sheet date instead of the average number of shares before and after dilution and the comparison year has been adjusted in this interim report. The effect of the adjustment means that earnings per share before and after dilution change from SEK -0.43 to SEK -0.45 for Q3 2021 and from SEK -0.64 to SEK -1.58 for 2021 year to date.

All amounts KSEK unless specified	2021 Jul-Sep	2021 Jul-Sep Restated	2021 Jan-Sep	2021 Jan-Sep Restated	2021 Jan-Dec	2021 Jan-Dec Restated
Profit/loss for the period	-37,245	-37,245	-104,946	-104,946	-141,736	-141,736
Average number of shares before dilution	85,666,338	83,173,402	85,666,338	83,173,402	220,584,878	89,670,050
Earnings per share before dilution, SEK	**-0.43**	**-0.45**	**-1.23**	**-1.26**	**-0.64**	**-1.58**
Average number of shares after dilution	85,666,338	83,173,402	85,666,338	83,173,402	220,740,173	89,670,050
Earnings per share after dilution, SEK	**-0.43**	**-0.45**	**-1.23**	**-1.26**	**-0.64**	**-1.58**

Financial definitions

Equity per share after dilution
Equity divided by the total number of shares at the end of the period and any outstanding options where the Company's share price on the reporting date is at least equal to the conversion price of the option.

Equity per share before dilution
Equity divided by the number of shares at the end of the period.

R&D costs
The Company's direct costs for research and development. Refers to costs for personnel, materials and external services.

R&D costs as a percentage of operating costs excluding impairments
R&D costs as a percentage of operating costs excluding impairments.

Average number of shares before and after dilution
Average number of outstanding shares during the period. The number of shares after dilution also takes account of outstanding options where the Company's share price on the reporting date is at least equal to the conversion price of the option.

Average number of employees
Average number of employees at the beginning and end of the period.

Average number of employees within R&D
Average number of employees within the Company's R&D departments at the beginning and end of the period.

Cash flow from operating activities
Cash flow before investing and financing activities.

Cash and cash equivalents, including securities
Cash and cash equivalents consists of bank balances, interest funds and publicly traded corporate bonds.

Cash flow for the period
Net change in cash and cash equivalents excluding the impact of unrealized foreign exchange gains and losses.

Earnings per share before and after dilution
Earnings divided by the weighted average number of shares during the period before and after dilution respectively. If the result is negative, the number of shares before dilution is also used for the calculation after dilution.

Operating costs excluding impairments
Other external costs, personnel costs and depreciation (excluding impairments of tangible and intangible assets).

Operating profit/loss
Profit/loss before financial items and taxes.

Equity ratio
Equity as a percentage of total assets.

Total assets
Total of the Company's assets.

Calculation of Performance Measures

Alligator presents certain financial performance measures in this report, including measures that are not defined under IFRS. The Company believes that these performance measures are an important complement because they allow for a better evaluation of the Company's economic trends. These financial performance measures should not be viewed in isolation or be considered to replace the performance indicators that have been prepared in accordance with IFRS. In addition, such performance measures as Alligator has defined them should not be compared with other performance measures with similar names used by other companies. This is because the above-mentioned performance measures are not always defined in the same manner, and other companies may calculate them differently to Alligator.

Below is shown the calculation of key figures, for the mandatory earnings per share according to IFRS and also for performance measures that are not defined under IFRS or where the calculation is not shown in another table in this report.

The Company's business operation is to conduct research and development which is why "R&D costs/Operating costs excluding impairment in percent" is an essential indicator as a measure of efficiency, and how much of the Company's costs relate to R&D.

The Company does not have a steady flow of income, with income generated irregularly in connection with the signing of license agreements and achievement of milestones. Therefore, the Company monitors performance indicators such as equity ratio and equity per share in order to assess the Company's solvency and financial stability. These are monitored along with the cash position and the various measures of cash flows shown in the consolidated statement of cash flow.

All amounts KSEK unless specified	**2022 Jul-Sep**	**2021 Jul-Sep**	**2022 Jan-Sep**	**2021 Jan-Sep**	**2021 Jan-Dec**
Profit/loss for the period	-51,365	-37,245	-140,152	-104,946	-141,736
Average number of shares before dilution	220,584,878	83,173,402	220,584,878	83,173,402	89,670,050
Earnings per share before dilution, SEK	**-0.23**	**-0.45**	**-0.64**	**-1.26**	**-1.58**
Average number of shares after dilution	220,584,878	83,173,402	220,584,878	83,173,402	89,670,050
Earnings per share after dilution, SEK	**-0.23**	**-0.45**	**-0.64**	**-1.26**	**-1.58**
Operating costs	-56,658	-41,029	-156,561	-112,753	-156,691
Operating costs excluding impairments	**-56,658**	**-41,029**	**-156,561**	**-112,753**	**-156,691**
Administrative expenses	8,309	16,316	26,770	32,148	35,423
Depreciation	1,285	2,941	5,375	8,549	11,144
Research and development costs	**-47,064**	**-21,771**	**-124,417**	**-72,056**	**-110,123**
R&D costs / Operating costs excluding impairments %	**83%**	**53%**	**79%**	**64%**	**70%**
Equity	142,256	85,029	142,256	85,029	282,273
Average number of shares before dilution	220,584,878	85,666,338	220,584,878	85,666,338	220,584,878
Equity per share before dilution, SEK	**0.64**	**0.99**	**0.64**	**0.99**	**1.28**
Average number of shares after dilution	220,584,878	85,666,338	220,584,878	85,666,338	220,740,173
Equity per share after dilution, SEK	**0.64**	**0.99**	**0.64**	**0.99**	**1.28**
Equity	142,256	85,029	142,256	85,029	282,273
Total assets	205,054	131,822	205,054	131,822	333,200
Equity ratio, %	**69%**	**65%**	**69%**	**65%**	**85%**
Cash and cash equivalents at end of period	**147,411**	**79,314**	**147,411**	**79,314**	**278,148**

For definitions, see the section "Financial definitions" on page 21.

The declaration of the Board of Directors and the CEO

Anders Ekblom

Hans-Peter Ostler

Eva Sjökvist Saers

Denise Goode

Veronica Wallin

Laura von Schantz

Graham Dixon

Staffan Encrantz

Søren Bregenholt

The Board and the CEO declare that this Interim report provides a true and fair overview of the Company and the Group's operations, positions and earnings and describes the material risks and uncertainty factors faced by the Parent Company and the companies within the Group.

Lund, October 20, 2022

Anders Ekblom
Chairman of the Board

Hans-Peter Ostler
Vice chairman of the Board

Eva Sjökvist Saers
Board member

Graham Dixon
Board member

Veronica Wallin
Board member

Laura von Schantz
Board member

Denise Goode
Board member

Staffan Encrantz
Board member

Søren Bregenholt
CEO

THIS IS THE TRANSLATION FROM THE SWEDISH ORIGINAL

Review report

Alligator Bioscience AB (publ), corporate identity number 556597-8201

To the Board of Directors of Alligator Bioscience AB (publ)

Introduction
We have reviewed the condensed interim report for Alligator Bioscience AB (publ) as at September 30, 2022 and for the nine months period then ended. The Board of Directors and the Managing Director are responsible for the preparation and presentation of this interim report in accordance with IAS 34 and the Swedish Annual Accounts Act. Our responsibility is to express a conclusion on this interim report based on our review.

Scope of review
We conducted our review in accordance with the International Standard on Review Engagements, ISRE 2410 Review of Interim Financial Statements Performed by the Independent Auditor of the Entity. A review consists of making inquiries, primarily of persons responsible for financial and accounting matters, and applying analytical and other review procedures. A review is substantially less in scope than an audit conducted in accordance with International Standards on Auditing and other generally accepted auditing standards in Sweden. The procedures performed in a review do not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.

The procedures performed in a review do not enable us to obtain assurance that we would become aware of all significant matters that might be identified in an audit. Accordingly, we do not express an audit opinion.

Conclusion
Based on our review, nothing has come to our attention that causes us to believe that the interim report is not prepared, in all material aspects, in accordance with IAS 34 and the Swedish Annual Accounts Act regarding the Group, and in accordance with the Swedish Annual Accounts Act regarding the Parent Company.

Significant uncertainty factors regarding the assumption of going concern
We would like to draw attention to the information provided in the interim report, under the section ”Financial position”, page 9, where it appears that the group’s continued operations are dependent on new financing to ensure continued operations. Should the measures that the Board of Directors plans to implement not be able to be implemented, there is a significant uncertainty factor regarding the company’s ability to continue operations. Our statement is not modified in this regard.

Malmö, 20 October 2022
Ernst & Young AB

Peter Gunnarsson
Authorized Public Accountant

Glossary

Agonist. A compound which binds to a receptor and stimulates its activity.

Antigen. Substance which triggers a reaction in the immune system, such as a bacteria or virus.

Antibody. Proteins used by the body's immune defenses to detect and identify xenobiotic material.

Bispecific antibodies. Antibody-based products which bind to two different targets and thus have dual functions.

Cancer. A disease in which cells divide in an uncontrolled manner and invade neighboring tissue. Cancer can also spread (metastasize) to other parts of the body through the blood and the lymphatic system.

Checkpoint inhibitor. An antibody with the ability to break the immune system's tolerance to something dangerous, for example a cancer tumor. Immune-inhibiting signals can be blocked through binding to a specific receptor such as CTLA-4 or PD-1.

Clinical study. The examination of healthy volunteers or patients to study the safety and efficacy of a potential drug or treatment method.

CRO (Clinical Research Organization). Company specialized in performing contract research and clinical studies on behalf of other pharma or biotech companies.

CTA (Clinical Trial Authorization). Application to start clinical trials in humans which is submitted to a regulatory authority.

CTLA-4 (Cytotoxic T-lymphocyte-Associated protein-4). An immune-inhibiting molecule expressed in and on the surface of T cells, primarily regulatory T cells.

Dendritic cell. A type of cell which detects xenobiotic substances. A key role of dendritic cells is their ability to stimulate T cells in the immune system.

Discovery. This research phase usually encompasses the development and evaluation of treatment concepts, the evaluation of potential drug candidates, and early efficacy studies.

Drug candidate. A specific compound usually designated before or during the preclinical phase. The drug candidate is the compound that is then studied in humans in clinical studies.

EMA. The European Medicines Agency.

Experimental model. A model of a disease or other injury to resemble a similar condition in humans.

FDA. The US Food and Drug Administration.

GMP (Good Manufacturing Practice). Quality assurance methodology designed to ensure that products are manufactured in a standardized manner, such that quality requirements are satisfied.

Immuno-oncology. Field of oncology in which cancer is treated by activating the immune system.

INN (International Nonproprietary Name). Generic name on a drug substance. The INN is selected by the World Health Organization (WHO) since 1953.

Lead. A potential drug candidate which binds to the actual target molecule/s.

Ligand. Binds to a receptor. Could be a drug, hormone or a transmitter substance.

Lymphocyte. A type of white blood cells.

Macrophages. A type of white blood cell of the immune system that engulfs and digests cellular debris and foreign materia such as bacteria.

Milestone payment. Financial consideration received in the course of a project/program when a specified objective is reached.

Mitazalimab. Generic name (INN) for ADC-1013.

Monospecific antibodies. Antibody-based product which bind only to one target, such as a receptor.

NK cells. NK cells (Natural Killer) are lymphocytes with the ability to activate several different cells in the immune system, such as macrophages.

Oncology. Term for the field of medicine concerned with the diagnosis, prevention and treatment of tumor diseases.

Patent. Exclusive rights to a discovery or invention.

PD-1 (Programmed Death-1). Immune-inhibiting receptor on the surface of certain cells, for example tumor cells.

PD-L1 (Programmed Death-Ligand-1). The ligand that binds to PD-1, helping the cancer evade the body's immune defense.

Phase 1,2 and 3. The various stages of studies on the efficacy of a pharmaceutical in humans. See also "clinical study." Phase 1 examines the safety on healthy human subjects, Phase 2 examines efficacy in patients with the relevant disease and Phase 3 is a large-scale study that verifies previously achieved results. In the development of new pharmaceuticals, different doses are trialed and safety is evaluated in patients with relevant disease. Phase 2 is often divided into Phase 2a and Phase 2b. In Phase 2a, which is open, different doses of the pharmaceutical are tested without comparison against placebo and focusing on safety and the pharmaceutical's metabolism in the body. Phase 2b is 'blind', and tests the efficacy of selected dose(es) against placebo.

Pharmacokinetics. The study of the turnover of substances in the body, for example how the amount of the substance is changed by absorption, distribution, metabolism and excretion.

Pharmacology. The study of how substances interact with living organisms to bring about a functional change.

Preclinical. The stage of drug development before the drug candidate is tested in humans. It includes the final optimization of the drug candidate, the production of materials for future clinical studies and the compilation of a data package for an application to start clinical studies.

Proof of concept studies. Studies carried out to provide support for dosages and administration paths in subsequent clinical studies.

R&D. Research & Development

Receptor. A receptor on a cell which picks up chemical signals.

RECIST. Response Evaluation Criteria in Solid Tumors - simple and pragmatic methodology to evaluate the activity and efficacy of new cancer therapeutics in solid tumors, using validated and consistent criteria to assess changes in tumor burden.

Sponsor. The person, company, institution or organization responsible for initiating, organizing or financing a clinical study.

T cell. A type of white blood cell which is important to the specific immune defense.

Tumor-associated antigen (TAA). A protein expressed to a much higher degree on the surface of tumor cells than healthy cells.

Tumor cell. A cell that divides relentlessly.

Tumor necrotic factor receptor superfamily (TNFR-SF). A group of immune-modulating target proteins related to the tumor necrosis factor protein. The name 'tumor necrosis factor' was derived from the fact that the first function detected for the protein was its ability to kill some types of tumor cells, though it was later discovered to have an immune-regulatory function.

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