

### Q2 Interim report Aug-Oct 2025

### Positioned for growth in Pharma Services and in the USA

#### Q2 (August – October 2025) (comparison figures same period 2024)

- Net sales amounted to SEK 2,675 (2,312) thousand for the quarter, corresponding to an increase of 16%. In local currency, net sales increased by 27%.
- Cash flow from operating activities amounted to SEK -18,111 (- 21,053) thousand.
- At the end of the quarter, cash and cash equivalents amounted to SEK 100,664 thousand (SEK 16,305 thousand previous quarter).

#### Q1 and Q2 (May – October 2025) (comparison figures same period 2024)

- Net sales amounted to SEK 5,271 (4,027) thousand for the period, corresponding to an increase of 31%. In local currency, net sales increased by 45%.
- Cash flow from operating activities during the period amounted to SEK -35,306 (-50,096) thousand.

### Significant events during Q2 (August – October 2025)

- A rights issue in August raised capital of SEK 80.1 million prior to issue costs.
- A directed share issue in August generated approximately SEK 42.2 million for Biovica, prior to issue costs.

### Significant events after the end of the quarter

- Two new studies with DiviTum TKa presented at SABCS 2025
- New commercial agreement signed with NCI Center in the USA.

Selected Key Performance Indicators:	Aug-Oct	Aug-Oct	May-Oct	May-Oct	Full year
SEK 000s	2025	2024	2025	2024	24/25
Net sales	2,675	2,312	5,271	4,027	8,619
Operating profit (loss)	-14,964	-19,753	-34,298	-43,316	-85,839
Profit (loss) for the period	-15,111	-21,152	-32,886	-44,041	-87,624
Earnings per share, before dilution, SEK	-0.06	-0.23	-0.18	-0.50	-0.95
Cash and cash equivalents at the end of the period	100,664	61,883	100,664	61,883	24,415
Cash flow from operating activities	-18,111	-21,053	-35,306	-50,096	-85,367
Cash flow for the period	84,429	-3,231	76,359	-17,466	-54,730
Equity	123,804	86,879	123,804	86,879	43,206
Equity per share, SEK	0.48	0.96	0.69	0.99	0.44
Equity ratio (%)	89%	80%	89%	80%	67%

### **Financial Targets**

- In connection with this interim report, the Board of Directors has decided to refrain from communicating financial targets, pending a more stable and predictable development in sales.

### Biovica in brief – Treatment decisions with greater certainty

- Biovica develops and commercializes the blood-based biomarker assay, DiviTum® TKa, which enables early-stage evaluation of treatment effectiveness. The initial focus is on breast cancer.
- DiviTum TKa has obtained FDA 510(k) clearance in the USA and has CE marking in the EU.
- Biovica's shares are traded on the Nasdag First North Premier Growth Market (BIOVIC B).

### Webcast:

When: 18 December 2025, 3 PM to 3.30 PM CET Where: registration via: <u>Biovica Q2 Earnings Call</u>

Broadcast language: in English

This is a translation of the Swedish original Annual Report. In the event of any discrepancies, the Swedish version shall prevail.

### CEO's comments

We made significant progress during the quarter and continued to build a strong platform for future growth. Net sales during the quarter increased by 16% (27% in local currency) compared to the same quarter previous year and by 31% (45% in local currency) compared to the same period previous year. Thanks to good cost control, we also improved our cash flow significantly.

There has been a delay in the study to be carried out in collaboration with the large integrated healthcare network in the USA, which is something we flagged already in the previous quarterly report. Because of that, we did not meet our sales target for the period.

It means that we will neither achieve our target of SEK 50 million for the current financial year, nor our target of SEK 150 million for the coming year. The Board has decided that Biovica will refrain from publishing these types of targets until the sales pipeline has matured. Nevertheless, we have the ambition of achieving positive cash flow during 2027. The plan is based on continued positive development within Pharma Services and the progress achieved in collaboration with the US healthcare network toward the initiation of the planned breast cancer clinical trial.

For Pharma Services, we previously reported that there are agreements in place with a contract value of approximately SEK 25 million. The revenue from those will be recognized during the next two to three years and we have planned deliveries that will boost revenue in this segment already during Q3 and Q4.

We are seeing a clear trend towards both larger projects and larger pharmaceutical companies in our customer base. Five of our current customers each report annual sales in excess of SEK 100 billion. There is enormous potential here to expand on the collaborations and establish DiviTum TKa as a standard tool in the development of new oncology drugs.

Our collaboration with the large integrated healthcare network in the USA to launch a largescale clinical study is progressing. In the previous quarterly report, we reported that the original study protocol needed to be revised, resulting in a delay. With the protocol now revised the goal is for the study to get underway during the first half of 2026, providing an increase in revenue as well as important clinical results.

Following nine quarters of growth, sales within clinical use in the USA have temporarily stalled. The break in the trend is explained, among other things, by the fact that an NCI/NCCN-designated cancer center, where a study was recently completed, has postponed clinical implementation pending a commercial agreement. After the end of the reporting period, such an agreement has now been signed, creating favorable conditions for an accelerated adoption of DiviTum TKa. It is part our strategy of focusing more sharply on major NCI/NCCN centers, along with signing commercial agreements to boost sales. We have also noticed higher interest for use in early breast cancer and have developed an LDT version of the assay that can be offered to the market. In light of these developments, we are confident in our ability to reignite strong growth in the coming quarters.

Our strategic collaboration with Tempus – a leader in data-driven precision medicine – continues. The partnership expands our reach to a broader network of oncologists in the USA and is a key element of our long-term plan to establish DiviTum TKa as a standard tool for monitoring treatment efficacy. Efforts to develop a joint value proposition have taken much longer than initially planned, but we now have common ambition to be ready for launch during the second half of 2026.

Our good standing in the scientific community also improved during the period. In December of 2025, two new studies were presented at SABCS confirming the role of DiviTum TKa in metastatic breast cancer. The findings demonstrate the value of DiviTum TKa for dose optimization and early identification of treatment resistance, factors that are critical to treatment outcomes and cost efficiency.

The studies were conducted in the USA at: Yale Cancer Center, Connecticut and Mass General

Brigham (MGB) Cancer Institute, Massachusetts. Both institutions are among the leading and most prestigious cancer research centers in the United States.

The Yale study concluded that DiviTum TKa is a promising tool for monitoring patient adherence to CDK4/6 inhibitors and for detecting potential drugdrug interactions in the clinical setting.

The study at MGB Cancer Institute confirms TKa as a promising biomarker and shows how TKa patterns, combined with ctDNA profiles, can be linked to genomic alterations and treatment response in HR+/HER2- metastatic breast cancer.

With the aim of strengthening customer relationships and accelerating sales growth, I plan to spend more time in the United States in 2026. It is a strategic investment aimed at enhancing our relationships with key partners and achieving commercial success in the near term. In summary, we have a strong position in a growing market, a product that makes a meaningful

difference for both patients and caregivers, and an organization that is ready to deliver. As we move into the next quarter, our focus remains on driving sales growth and creating value for patients, our partners, and shareholders.



Anders Rylander, CEO

Significant events during the first quarter (May-July 2025).

### Biovica is now collaborating with Tempus to expand the commercial reach of DiviTum TKa

Biovica signed an agreement with Tempus, a leader in AI and data-driven precision medicine.

### Biovica announced that it has signed a Master Service Agreement (MSA) with a US-based pharma/biotech company.

An initial work order valued at SEK 4 million has also been signed.

# New data on DiviTum TKa use in three areas of cancer

#### presented at ASCO

Biovica presented three abstracts based on studies with DiviTum TKa at ASCO. The new data further reinforces DiviTum TKa's role as a predictive biomarker across three different cancer indications:

- Hormone receptor—positive (HR+)
  metastatic breast cancer (MBC) in patients
  treated with CDK4/6 inhibitors, as studied
  in the high-profile PEARL trial
- BRAF V600–mutated metastatic melanoma treated with immune checkpoint inhibitors (ICIs)
- Ovarian cancer treated with platinumbased chemotherapy

### Biovica signed an agreement with its fifth Tier 1 biopharma company in the USA

Biovica has signed a new MSA with a company that has also placed an initial work order of approximately KSEK 800 thousand. It is the fifth Tier 1 biopharma company in the USA to join the Biovica customer base in Pharma Services.

### Biovica announced financial targets subsequent to important partnership and commercial success

Biovica announced financial targets for the next two fiscal years.

Biovica's financial targets are:

- Fiscal year 2024/25 (ending 30 April 2025): Net sales of SEK 8.5 million
- Fiscal year 2025/26: Net sales of SEK 50 million
- Fiscal year 2026/27: Net sales of SEK 150 million

With expected operating expenses at the current level of approximately SEK 90 million per year, Biovica anticipates that it will become cash flow positive during the third quarter of the 2026/27 financial year.

Significant events during the second quarter (Aug-Oct 2025).

# A rights issue generated approximately SEK 80.1 million, prior to issue costs.

# A directed share issue generated approximately SEK 42.2 million, prior to issue costs.

The Board of Directors resolved to exercise the option of oversubscription in the rights issue through a directed share issue to the investors who have entered into guarantee undertakings as top-down guarantors (the "Anchor Investors"). The directed share issue corresponds to additional liquidity of approximately MSEK 42.2 prior to issue costs and the set-off of a bridge loan of approximately MSEK 10.1, which means that the company will receive a total of approximately MSEK 122.3 prior to issue costs from the rights issue and the directed share issue.

# Significant events after the end of the period

### Two new studies with DiviTum TKa presented at SABCS 2025

Both studies highlight the clinical relevance of DiviTum TKa as a dynamic biomarker for monitoring treatment response.

Researchers from Yale Cancer Center evaluated whether early TKa measurements could identify suboptimal CDK4/6 inhibitor activity caused by medication non-adherence or drug interactions. A multi-center case series by the Mass General Brigham Cancer Institute and Washington University in St. Louis examined the associations between baseline circulating tumor DNA (ctDNA) profiles and early TKa response patterns in 22 patients starting CDK4/6 inhibitor therapy. The case studies demonstrated how ctDNA and TKa provide complementary insights — ctDNA reveals resistance biology, while TKa reflects treatment response in real time.

### Biovica signed a commercial agreement with a leading US cancer center that will increase availability of DiviTum TKa

The agreement strengthens Biovica's presence among leading US cancer institutions. It also supports introduction at a leading academic cancer center and the company's commercialization in the USA. The introduction of DiviTum TKa underscores the biomarker's growing role in treatment monitoring and establishes a reference institution that increases confidence among clinicians and supports dialog with payers — paving the way for broader integration into clinical practice.

#### Other

#### 2025 AGM

The Annual General Meeting was held on 23 September 2025. Summary of events at the AGM:

- The financial statements were adopted and the Board of Directors and CEO were discharged from liability for the financial year.
- The AGM resolved that no dividends would be distributed to shareholders.
- Fredrik Alpsten and Niels Bogerd were newly elected as Board members.
- The following Board members were reelected: Annika Carlsson Berg, Marie-Louise Fjällskog, Maria Holmlund, Anders Rylander and Jesper Söderqvist.
   Lars Holmqvist declined re-election.
- Fredrik Alpsten was elected as Chairman of the Board.
- Grant Thornton Sweden AB was re-elected as the company's auditor, with Stéphanie Ljungberg as head auditor.
- The Board was granted the authority to issue new shares equal to 20% of the current number of shares.
- Implementation of Performance Share Program 2025/2028:1-3 for the company's employees and the Board of Directors.
- Approval of directed share issue.

### Comments on the financial performance of the Group

### Q2 - Sales and earnings

The quarter covers the period 1 August 2025 through 31 October 2025. The comparison figures are for the period 1 August 2024 through 31 October 2024.

Net sales for the period amounted to SEK 2,675 (2,312) thousand. Sales in the second quarter are derived from three different product groups. These are: Tests (IVD) for the US market, Tests (RUO) and DiviTum Kits (RUO), which are primarily sold to the pharmaceutical industry and used for research purposes.

Net sales increased by 16% (27% excluding currency effects) compared to the same period previous year, which is primarily attributable to strong growth in Pharma Services for Tests (RUO) by 86% (122% excluding currency effects) along with Tests (IVD) for the US market by 47% (62% excluding currency effects). Net sales of Divitum Kits (RUO) declined compared to the same period previous year, corresponding to a trend we have noticed, where more pharmaceutical companies are choosing to send samples to Biovica for analysis, rather than purchasing the kit and doing the analysis themselves. There were no sales to the European market during the quarter. More information is provided in Note 1.

The operating loss for the quarter was SEK -14,964 (-19,753) thousand.

The earnings improvement compared to last year is attributable to a reduction in expenses after the Group restructuring that was implemented in April 2024, along with higher sales.

Net financial items amounted to SEK -18 (216) thousand. Loss after financial items was SEK - 14,982 (-19,537) thousand. Loss for the period was SEK -15,111 (-21,152) thousand.

The average number of employees for the quarter was 24 (27) employees, of which 14 (14) are women.

### Q1 and Q2 - Combined sales and earnings

The quarter covers the period 1 May 2025 through 31 October 2025. The comparison figures are for the period 1 May 2024 through 31 October 2024.

Net sales for the period amounted to SEK 5,271 (4,027) thousand. Sales in the first half of the year are derived from three different product groups. These are: Tests (IVD) for the US market, Tests (RUO) and DiviTum Kits (RUO), which are primarily sold to the pharmaceutical industry and used for research purposes.

Net sales increased by 31% (45% excluding currency effects) compared to the same period previous year, which is primarily attributable to strong growth with Tests (IVD) for the US market by 59% (76% excluding currency effects) and in Pharma Services with Tests (RUO) by 66% (83% excluding currency effects). Sales of Divitum Kits (RUO) were essentially unchanged compared to the same period previous year. There were no sales to the European market during the first half of the year. More information is provided in Note 1.

The operating loss for the period was SEK 34,298 (43,316) thousand.

The earnings improvement compared to last year is attributable to higher sales along with a reduction in expenses after the Group restructuring that was implemented in April 2024.

Net financial items amounted to SEK 179 (-407) thousand. Loss after financial items was SEK -34,119 (-43,722) thousand. Loss for the period was SEK -32,886 (-44,041) thousand.

The average number of employees for the quarter was 24 (27) employees, of which 14 (14) are women.

Financial position, cash flow and investments

The closing amount for cash & cash equivalents on 31 October 2025 was SEK 100,664 (61,883) thousand.

Net investments in property, plant and equipment in the form of equipment for the year amounted to SEK -2 (0) thousand.

Intangible assets consist of capitalized development expenditure for the various versions of DiviTum TKa along with patents. There has not been any new capitalization during the year. More information is available under the heading, Intangible assets, on page 10.

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### **Funding**

During the period, a rights issue and directed issue to anchor investors of the rights issue were carried out in order to generate the capital required for the continued launch of DiviTum TKa. The proceeds from this were approximately SEK 122.3 million prior to issue costs and the set-off of a bridge loan.

With cash and cash equivalents of SEK 101 million at the end of the period, along with the current sales and cost forecast, the Board assesses that the Company has sufficient resources to support continued operations as of the date of this report.

### Related party transactions

During the period, the company, represented by parties related to the CEO and Board member, Anders Rylander, leased office facilities to the Parent Company. The total fee for rent paid was SEK 68 (68) thousand. Transactions were in accordance with market-based terms and conditions.

### Incentive programs

Program	То	Country	Options / Saving Shares	Subscription price	Subscription period	Equity Increase	Number of class B shares	Dilution
23/26:1*	Employees	US	240,000	10.13	1 June – 30 September 2026	16,000	240,000	0.08%
23/26:2*	Employees	US	56,000	10.12	11 July 2023 – 15 September 2026	3,733	56,000	0.02%
23/26:3*	Employees	SE	358,000	8.24	1 October- 1 November 2026	23,867	358,000	0.12%
23/26:4*	Board of Directors	SE	195,000	8.24	1 October- 1 November 2026	13,000	195,000	0.07%
23/26:5*	Employees	US	155,250	12.66	1 October- 1 November 2026	10,350	155,250	0.05%
23/26:6*	Employees	US	51,750	11.10	15 September - 1 November 2026	3,450	51,750	0.02%
SSP 24/27:1**	Employees	SE	621,600	2.90	1 October 2027- 1 November 2027	41,440	621,600	0.21%
SSP 24/27:2**	Board of Directors	SE	420,000	2.90	1 October 2027- 1 November 2027	28,000	420,000	0.14%
ESOP 24/27:3*	Employees	US	176,400	3.65	1 October 2027- 1 November 2027	11,760	176,400	0.06%
PRSU 24/27:4**	Employees	US	176,400	3.91	1 October 2027- 1 November 2027	11,760	176,400	0.06%
PRSU 2025/2028:1**	Board of Directors	SE	1,853,100	0.7416-1.0506	23 September 2028 - 23 October 2028	123,540	1,853,100	0.63%
PRSU 2025/2028:2**	Employees	SE	1,980,900	0.7416-1.0506	23 September 2028 - 23 October 2028	132,060	1,980,900	0.68%
PRSU 2025/2028:3**	Employees	US	1,022,400	0.7416-1.0506	23 September 2028 - 23 October 2028	68,160	1,022,400	0.35%
	•		7,306,800	_		487,120	7,306,800	2.50%

### Incentive programs

Valuation is as per the Black & Scholes pricing model for Warrants/ Options\* and as per Monte Carlo simulation for Share Savings Program/Performance Share Program\*\*. The programs 23/26:3-6 were never implemented due to the unfavorable stock price trend during fall 2023.

Resolutions were passed at the EGM on 23 September 2025 on 3 performance share programs 25/28: 1-4, which were distributed during fall of 2025. The incentive programs distributed free-of-charge have been calculated and reported in accordance with IFRS 2. Accordingly, the increase in both personnel expenses (debit) and equity (credit), amounted to SEK 57 (107) in the second quarter. The corresponding figure for May-Oct is SEK 210 (200) thousand. Additional information is available in the Annual Report for 2024/2025.

As of the closing date, the company had 7,306,800 (2,574,854) subscription rights, options and performance shares outstanding from the employee long-term incentive program. A total of 67,496 (67,496) of the stock options have been earned, a total of 5,577,410 (910,610) unearned but still possible to earn and the remainder expired since the person they had been allocated to had left the company.

#### Shares

As of 31 October 2025, the number of outstanding shares in Biovica was 291,911,199, of which 14,423,973 shares are Class A and 277,487,223 shares are Class B. The total number of votes amounted to 320,759,145. The rights issue was implemented in July 2025 and the directed issue to anchor investors of the rights issue was implemented in August 2025. The proceeds from this were approximately SEK 122.3 million prior to issue costs and the set-off of a bridge loan, which were received and registered with the Swedish Companies Registration Office during the period.

### Subscription rights TO4 B

Based on the outcome of the Rights Issue that was published on 5 August 2025, the Board decided on 13 August to utilize the Oversubscription Option and carry out a directed share issue for a total of 67,002,517 Class B shares in the Company to anchor investors, aimed at ensuring their full allocation in the Rights Issue.

As compensation to the anchor investors, they obtain, free-of-charge, the same number of Warrants from TO4 B (67,002,517) as the number of guaranteed shares, meaning that the company's share capital could increase by SEK 4,466,857 with full subscription of TO4 B.

One warrant of series TO4 B entitles the holder to subscription of one new Class B share in the

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company during the period ranging from registration of the warrants with the Swedish Companies Registration Office through 30 June 2030, at a subscription price of SEK 0.95 if the warrant is exercised by 30 June 2028, and at a subscription price of SEK 1.25 if the warrant is exercised during the period from 1 July 2028 through 30 June 2030. For subscription of the Class B shares, the portion of the subscription price that exceeds the quotient value of the previous shares will be added to the share premium reserve. Full subscription at a price of SEK 0.95, would generate approximately SEK 63,652,391 for the company, prior to issue costs. Full subscription at a price of SEK 1.25, would generate approximately SEK 83,753,146 for the company, prior to issue costs.

#### Reclassification of shares

At the end of each calendar quarter, class A shareholders are offered the opportunity of reclassifying their shares to B shares.

Reclassification from Class A to Class B shares lowers the voting power, in that Class A shares carry three votes each and Class B shares carry one vote each. The Class A shares are unlisted, while Biovica's Class B shares are traded on Nasdaq First North Premier Growth Market, Stockholm. No reclassification occurred on 30 September 2025.

# Policies for preparing the interim report *Accounting policies*

This interim report was prepared in accordance with IAS 34, Interim Financial Reporting. The Group applies the Annual Accounts Act, International Financial Reporting Standards (IFRS) that have been adopted by the EU and RFR 1 Additional Accounting Regulations for Groups when preparing the financial statements. The Parent Company applies RFR 2 Accounting for Legal Entities when preparing the financial statements. The applied accounting policies otherwise correspond with those described in the Annual Report for 2024/2025.

### New standards and interpretations that enter into force in 2025 and later

As of the date when these financial statements were approved for release, no new standards, revisions or interpretations of existing standards that have not yet entered into force or been published by the International Accounting Standards Board (IASB) have been early-adopted by the Group.

### Significant risks and uncertainties

There are a number of risks and uncertainties associated with the company's operations, including market, regulatory and financial risks. For a more detailed description of the risks (in Swedish), please see the Annual Report for 2024/2025.

### Liquidity risk

Conservatism in managing liquidity risk involves holding sufficient liquid funds or agreed credit facilities in order to be able to run the business.

With cash and cash equivalents of SEK 101 million at the end of the period, along with the current sales and cost forecast, the Board assesses that the Company has sufficient resources to support continued operations as of the date of this report.

### Uncertainties in the global situation

The Board and management continuously monitor the global situation and increased risks arising from, among other things, Russia's invasion of Ukraine and the war in Gaza. An increased risk of trade wars and the introduction of high tariffs — particularly between Europe and the United States — could negatively impact the company's earning capacity.

#### Financial risk management

The Group's business activities are associated with a variety of financial risks such as currency risk and interest rate risk on cash flows, credit risk and liquidity risk. The Group's overall risk management policy, which has been established by the Board, is to strive for minimal adverse effects on financial results and financial position.

### Currency risks

The Group has operations both domestically (in Sweden) and internationally, which means that there is exposure to fluctuations in different currencies, particularly USD and EUR. Currency risk arises through future business transactions and reported assets and liabilities. The increased scope of the company's operations has increased its net exposure to foreign currencies compared to prior years.

#### Interest rate risk on cash flows

Interest rate risk is the risk that the value of financial instruments will fluctuate due to changes in market interest rates. The Group currently only has

interest-bearing financial assets in the form of bank balances, which is why this risk is assessed as low.

#### Credit risk

Credit risk is the risk that a party to a transaction involving a financial instrument is unable to fulfill its obligation. Exposure to credit risks is marginal for both the Group and Parent Company.

### Significant assessments

### Assessments and estimates in the financial statements

In preparing the financial statements, the executive management team must make assessments and estimates that affect both the application of accounting policies and the reported amounts of assets, liabilities, income and expenses. The actual outcome may deviate from these estimates and assessments. A selection of these assessments is presented below.

#### Revenue from contracts with customers

Revenue from contracts with customers is reported at net realizable value and recognized when the performance obligation has been fulfilled and control over the goods or services has been transferred to the customer, in accordance with IFRS 15. This assessment shall occur from the customer's perspective, taking into consideration such things as transfer of ownership and risks, the customer's acceptance, physical access and the right to invoice. An assessment must also be made of whether control has been transferred at a specific point in time, or over time. All net sales are sales at a particular point in time. No sales are reported as sales over time. The timing of revenue recognition for services coincides with the reporting of test results to the customer. For goods, revenue is recognized when the risks and rewards associated with the item are transferred to the customer. Revenue is recognized at net realizable value at a specific point in time, which is when control of the goods or services has been transferred to the customer. The contract terms and conditions may vary but typically, transfer of control and thus revenue recognition occurs at the time of delivery.

### Segment reporting

The Group's operations consist of development, manufacturing and sales of blood analysis products. The Group's organizational structure is by function as follows: production, sales & marketing, administration and R&D. The Group is considered to be a single unit, where all of its sub-components are integrated and dependent upon each other.

Biovica's highest decision-making body monitors the consolidated income statement and statement of comprehensive income.

### Intangible assets

Expenditure for research that is for the purpose of gaining new scientific or technical knowledge is expensed as incurred. Expenditure for development (where the research results or other knowledge is used to achieve new or improved products or processes) is recognized as an intangible fixed asset in the statement of financial position if the product or process is technically or commercially usable and the company has adequate resources for monitoring the development and thereafter using or selling the intangible asset. Decisions on whether or not to capitalize expenditure on development projects are made by the company's Board of Directors based on documentation and support provided by the Audit Committee. The decision is based on whether it is possible to implement the project using existing or future resources and on whether conclusion of the project and launch is expected to occur in the foreseeable future. Directly attributable expenditure that is capitalized as part of the cost of the asset includes expenditure for employees and materials. With capitalization, consideration is given to the portion of expenditure recognized as revenue against received/expected grants. Capitalized development expenditure is reported as intangible assets and amortized as of the date when the asset is ready for use. The estimated useful life for capitalized development expenditures is 10 years. Other development expenses are recognized in the income statement as incurred. Patents are recognized at the cost of acquisition and they are amortized on a straightline basis over their estimated useful lives. The estimated useful life is assessed based on the legal life of the patent.

For a detailed description of these assessments, please see the Annual Report for 2024/2025.

### Note 1. Sales per product group

Net sales for the quarter are derived from the following product groups:

	Aug-Oct	Aug-Oct
SEK 000s	2025	2024
Tests (IVD) - USA	1,011	687
DiviTum Kits (IVD) - EU	0	280
Tests (RUO)	1,383	744
DiviTum Kits (RUO)	280	601
Total net sales	2,675	2,312

Corresponding figures for the period May - Oct

	May-Oct	May-Oct
SEK 000s	2025	2024
Tests (IVD) - USA	2,089	1,314
DiviTum Kits (IVD) - EU	0	280
Tests (RUO)	2,067	1,194
DiviTum Kits (RUO)	1,113	1,239
Total net sales	5,271	4,027

### Note 2 Reclassification of materials costs and inventory changes

As of Q2 2025–26, the Group has changed the presentation of certain production-related costs in the income statement. Direct labor costs and manufacturing overhead allocations incurred in production are no longer recognized as Materials cost and are instead included in Changes in inventory.

The reclassification was made to better reflect the link between production costs and changes in inventory, resulting in a more accurate presentation of the cost structure.

The change affects neither operating income nor net income for the year and relates solely to a reclassification between line items in the income statement.

Comparative figures have not been restated, as the change relates solely to classification and has no impact on earnings.

### KPIs for the Group

	Aug-Oct	Aug-Oct	May-Oct	May-Oct	Full year
SEK 000s	2025	2024	2025	2024	24/25
Net sales	2,675	2,312	5,271	4,027	8,619
Operating profit (loss)	-14,964	-19,753	-34,298	-43,316	-85,839
Profit (loss) for the period	-15,111	-21,152	-32,886	-44,041	-87,624
Earnings per share, before dilution	-0.06	-0.23	-0.18	-0.50	-0.95
Earnings per share, after dilution	-0.06	-0.23	-0.18	-0.50	-0.95
Cash and cash equivalents at the end of the period	100,664	61,883	100,664	61,883	24,415
Cash flow from operating activities	-18,111	-21,053	-35,306	-50,096	-85,367
Cash flow for the period	84,429	-3,231	76,359	-17,466	-54,730
Equity	123,804	86,879	123,804	86,879	43,206
Equity per share, SEK	0.48	0.96	0.69	0.99	0.44
Equity ratio (%)	89%	80%	89%	80%	67%
Average number of employees	24	27	24	26	26

Definitions are the same as those presented in the Annual Report for 2024/2025.

### Alternative key performance indicators

Of the KPIs presented above, the only one that is obligatory to report, and which is defined in accordance with IFRS is: Earnings per share, before and after dilution. For the other KPIs, the following are in accordance with IFRS presentation requirements: Profit (loss) for the year, Cash & cash equivalents at the end of the period, Cash flow for the period and Equity.

mow for the period and Ex		Reason for using alternative KPIs, which are
KPIs	Definition	not defined in accordance with IFRS.
Net sales	Income from goods sold	Shows the demand for the product.
Operating profit (loss)	Profit (loss) before financial items and tax.	Operating profit (loss) is an indication of the company's earnings generated from ordinary operations.
Earnings per share, before and after dilution	Profit (loss) divided by the weighted average number of shares during the period, before and after dilution.	
Cash & cash equivalents and short-term investments	Bank balances and short-term investments	
Cash flow from operating activities	Cash flow before the cash flow from investing activities and financing activities	
Cash flow for the period	Change in cash & cash equivalents for the period not including the effect from unrealized exchange gains and losses.	
Equity per share	Equity divided by the number of shares at the end of the period.	Management uses this KPI to monitor the value of equity per share.
Equity ratio	Equity as a percentage of total assets.	Management uses this KPI because it provides an indication of the company's financial stability.
Average number of employees	The average number of employees is calculated as the average during the period of the number of employees per month.	<i>,</i>

# Consolidated income statement and summary statement of comprehensive income

	Aug-Oct 2025	Aug-Oct 2024	May-Oct 2025	May-Oct 2024	Full year 2024/2025
Amount in SEK thousands					
Net sales (Note 1)	2,675	2,312	5,271	4,027	8,619
Other income	96	584	308	1,325	2,341
Operating income	2,772	2,896	5,578	5,351	10,961
Change in WIP inventory (Note 2)	714	0	714	0	0
Materials cost (Note 2)	-283	-214	-454	-370	-535
Other external costs	-5,795	-6,368	-12,249	-13,702	-28,332
Employee benefit expenses	-10,206	-13,095	-23,120	-29,215	-57,299
Depreciation/amortization	-2,133	-2,124	-4,272	-4,503	-8,843
Other operating expenses	-32	-848	-494	-877	-1,791
Operating expenses	-17,735	-22,649	-39,876	-48,667	-96,800
Operating profit (loss)	-14,964	-19,753	-34,298	-43,316	-85,839
Financial income	56	146	489	341	996
Financial expenses	-74	70	-310	-748	-1,139
Profit (loss) before tax	-14,982	-19,537	-34,119	-43,722	-85,983
Tax	-128	-1,615	1,233	-319	-1,641
Profit (loss) for the period	-15,111	-21,152	-32,886	-44,041	-87,624
Consolidated statement of comprehensive income					
Profit (loss) for the period	-15,111	-21,152	-32,886	-44,041	-87,624
Exchange differences when translating foreign operations	-185	-40	-111	-134	-632
Other comprehensive income for the period	0	0	0	0	0
Comprehensive income for the period	-15,295	-21,192	-32,997	-44,175	-88,256
Earnings per share					
Earnings per share, before dilution (SEK)	-0.06	-0.23	-0.18	-0.50	-0.95
Average number of shares, before dilution	259,439,634	90,818,790	178,613,009	87,437,175	92,569,248
Earnings per share, after dilution (SEK)	-0.06	-0.23	-0.18	-0.50	-0.95
Average number of shares, after dilution	259,439,634	90,818,790	178,613,009	87,437,175	92,569,248

Biovica International AB (publ)

### Consolidated statement of financial position, in summary

Amount in SEK thousands	2025-10-31	2024-10-31	2025-04-30
ASSETS			
Intangible assets	24,128	28,944	26,536
Machinery, equipment, tools, fixtures and fittings	1,020	984	1,049
Right-of-use assets	1,986	5,909	3,719
Other non-current receivables	387	438	396
Deferred tax asset	2,123	2,928	2,455
Total fixed assets	29,643	39,203	34,154
Inventories	2,139	2,077	1,930
Accounts receivable	3,051	1,372	1,815
Current receivables	3,073	3,888	2,634
Cash and Bank	100,664	61,883	24,415
Total current assets	108,928	69,220	30,794
TOTAL ASSETS	138,571	108,423	64,949
EQUITY			
Share capital	19,461	6,519	6,519
Other contributed capital	678,478	577,416	577,824
Reserves	-333	276	-222
Retained earnings (losses), including loss for the period	-573,801	-497,332	-540,915
Total equity	123,804	86,879	43,206
LIABILITIES			
Right-of-use liabilities	899	3,296	1,736
Deferred tax liability	706	1,609	1,849
Total non-current liabilities	1,605	4,905	3,585
Right-of-use liabilities	2,138	3,559	2,915
Advance payments from customers	1,783	0	0
Accounts payable	1,228	3,474	3,544
Current tax liabilities	0	106	14
Other liabilities	702	874	912
Accrued expenses and deferred income	7,311	8,627	10,774
Current liabilities	13,162	16,639	18,158
TOTAL EQUITY AND LIABILITIES	138,571	108,423	64,949

Biovica International AB (publ)

### Consolidated statement of changes in equity, in summary

	CI.	Other		<b>5</b>	<b>-</b>
Amount in SEK thousands	Share capital	contributed capital	Reserves	Retained earnings	Total equity
Opening balance, 1 May 2024	5,604	543,918	410	-453,291	96,640
Appropriation in accordance with AGM decision					0
New share issue	915	34,922			35,837
Issue costs		-1,604			-1,604
Share-based payments, employees		588			588
Transaction with owners	6,519	577,824	410	-453,291	131,461
Profit (loss) for the year				-87,624	-87,624
Other comprehensive income			-632		-632
Comprehensive income for the year		_			
(loss)	0	0	-632	-87,624	-88,257
Closing balance, 30 April 2025	6,519	577,824	-222	-540,915	43,206
Opening balance, 1 May 2024 Appropriation in accordance with AGM	5,604	543,918	410	-453,291	96,640
decision					0
New share issue	915	34,922			35,837
Issue fees		-1,604			-1,604
Share-based payments, employees		180			180
Transaction with owners	6,519	577,416	410	-453,291	131,054
Profit (loss) for the period				-44,041	-44,041
Other comprehensive income			-134		-134
Comprehensive income for the period	0	0	-134	-44,041	-44,175
Closing balance, 31 October 2024	6,519	577,416	276	-497,332	86,879
Opening balance, 1 May 2025 Appropriation in accordance with AGM	6,519	577,824	-222	-540,915	43,206
decision	12.010	100.057			0
New share issue	12,942	109,357			122,299
Issue fees		-8,926			-8,926
Share-based payments, employees		223			223
Transaction with owners	19,461	678,479	-222	-540,915	156,802
Profit (loss) for the period				-32,886	-32,886
Other comprehensive income			-111		-111
Comprehensive income for the period	0	0	-111	-32,886	-32,997
Closing balance, 31 October 2025	19,461	678 <i>,</i> 478	-333	-573,801	123,804

### Consolidated statement of cash flows, in summary

	Aug-Oct	Aug-Oct	May-Oct	May-Oct	May-April
Amount in SEK thousands	2025	2024	2025	2024	24/25
Cash flow from operating activities					
before changes in working capital	-12,665	-17,328	-29,481	-39,236	-77,113
Change in current receivables	331	180	-1,481	-4	-216
Change in current liabilities	-5,183	-3,670	-4,106	-10,937	-7,953
Change in inventories	-594	-235	-238	80	-85
Changes in working capital	-5,446	-3,725	-5,825	-10,861	-8,254
Cash flow from operating activities	-18,111	-21,053	-35,306	-50,096	-85,367
Investing activities					
Investments in PPE	-2	0	-147	0	-287
Coch flow from investing activities	2	0	1.47	0	207
Cash flow from investing activities	-2	0	-147	0	-287
Financing activities					
New share issue	122,299	19,422	122,299	35,837	35,837
Issue fees	-8,926	-804	-8,926	-1,604	-1,604
Borrowings	0	0	10,000	0	0
Amortization of loans	-10,831	-796	-11,561	-1,603	-3,309
	,		,	,	,
Cash flow from financing activities	102,542	17,822	111,812	32,630	30,925
Cash flow for the period	84,429	-3,231	76,359	-17 <i>,</i> 466	-54,730
Cash and cash equivalents at the	4 2 2 2 =	65.00-		70.00	70
beginning of the period	16,305	65,209	24,415	79,407	79,407
Translation difference, cash and cash	70	-96	110	-58	-262
equivalents  Cash and cash equivalents at the end of	-70	-96	-110	-58	-262
the period	100,664	61,883	100,664	61,883	24,415
the period	100,004	01,003	100,004	01,000	27,713

### Parent Company income statement, in summary

	Aug-Oct 2025	Aug-Oct 2024	May-Oct 2025	May-Oct 2024	May-April 24/25
Amount in SEK thousands					
Net sales	1,755	1,629	3,454	17,751	28,385
Other operating income	96	584	308	1,325	2,341
Total revenue	1,850	2,213	3,762	19,076	30,726
Change in WIP inventory (Note 2)	714	0	714	0	0
Materials cost (Note 2)	-278	23	-588	-573	-640
Other external costs	-10,601	-18,315	-24,174	-40,240	-78,062
Employee benefit expenses	-5,802	-7,211	-13,833	-16,415	-33,024
Depreciation/amortization	-1,242	-1,242	-2,483	-2,734	-5,217
Other expenses	-32	-848	-494	-877	-1,791
Operating expenses	-17,241	-27,593	-40,859	-60,839	-118,734
Operating profit (loss)	-15,391	-25,380	-37,097	-41,764	-88,008
Other interest income and similar items Other interest expenses and similar	221	145	548	340	994
items	-33	-66	-234	-254	-975
Profit (loss) before tax	-15,203	-25,301	-36,783	-41,678	-87,990
Tax on profit for the year	0	0	0	0	0
Profit (loss) for the period	-15,203	-25,301	-36,783	-41,678	-87,990

Comprehensive income (loss) equals the loss for the period.

### Parent Company balance sheet, in summary

Amount in SEK thousands	2025-10-31	2024-10-31	2025-04-30
ASSETS			
Intangible assets	24,128	28,944	26,536
Machinery, equipment, tools, fixtures and fittings	707	424	636
Financial assets	693	4,551	4,082
Total fixed assets	25,528	33,919	31,254
Inventories	2,062	1,938	1,866
Accounts receivable	1,750	1,000	1,120
Current receivables	1,984	2,749	1,796
Cash and Bank	97,663	60,213	22,722
Total current assets	103,459	65,901	27,504
TOTAL ASSETS	128,988	99,820	58,758
EQUITY			
Restricted equity	40,509	30,904	27,567
Non-restricted equity	77,362	56,059	13,491
Total EQUITY	117,871	86,963	41,059
LIABILITIES			
Current liabilities	11,117	12,857	17,699
Total LIABILITIES	11,117	12,857	17,699
TOTAL EQUITY AND LIABILITIES	128,988	99,820	58,758

### Glossary

**Abstract** - A short summary of a longer document, such as a dissertation or research article. It briefly states the purposes and results of the research. Abstracts are submitted to scientific conferences in order to spread knowledge of new research.

ASCO American Society of Clinical Oncology The world's leading professional organization for physicians and oncology professionals caring for people with cancer. Together with the Association for Clinical Oncology, ASCO represents nearly 45,000 oncologists.

Imaging These are methods that currently serve as the cornerstones for diagnostics and treatment planning for essentially all types of solid tumors. It includes computer tomography (CT) scans and other X-ray methods, magnetic resonance tomography (MRT) scans, positron emission tomography (PET) scans and ultrasound.

**CDK4/6 inhibitors** A new type of targeted, selective drugs that have been shown to be effective against several forms of cancers, including hormone receptor-positive breast cancer.

**CLIA laboratory** (The Clinical Laboratory Improvement Amendments): a clinical laboratory that has been certified to accept human samples from people in the USA for diagnostic testing. The Center for Medicare and Medicaid Services (CMS) is the regulatory body that grants certification.

CDx - Companion Diagnostics. These are diagnostic tests used to identify patients that would likely respond to a specific treatment, as well as to monitor the treatment effect on individual patients. They thus facilitate personalization of treatment.

ctDNA Circulating tumor DNA is found in the bloodstream and it is DNA that comes from cancerous cells and tumors. Most DNA is found inside the nucleus of a cell. As a tumor grows, cells die and are replaced by new ones. The dead cells are broken down and their contents, including DNA, are released into the bloodstream. ctDNA is small pieces of DNA, usually comprising less than 200 building blocks (nucleotides) in length.

**DiviTum Kits (IVD or RUO)** - This is the DiviTum TKa analysis kit. It can be sold as DiviTum Kits (IVD) to analyze samples taken from patients in a clinical setting, or DiviTum Kits (ROU), which are samples taken from patients for Research Use Only. Biovica's customers purchase DiviTum Kits and conduct the analyses in their own laboratories. DiviTum Kits (RUO) are primarily sold to

pharmaceutical companies or Clinical Research Organizations. Divitum Kits (IVD) are sold to European partners.

Fulvestrant (Faslodex) A drug that is used to treat hormone receptor (HR)-positive metastatic breast cancer in postmenopausal women with disease progression and HR-positive, HER2-negative advanced breast cancer in combination with palbociclib in women with disease progression after endocrine treatment. Fulvestrant is a Selective Estrogen Receptor Degrader (SERD). It works by binding to the estrogen receptor and destabilizing it, causing the cell's normal protein degradation processes to destroy it.

**IVD** In vitro diagnostics (IVD) are generally defined as a product which, regardless of whether they are used alone or in combination, are designed for performing in vitro tests on samples that have been taken from the human body. The main purpose is to obtain information for diagnostic, monitoring or compatibility purposes.

**Tests (IVD) USA -** This testing service is conducted at Biovica's CLIA laboratory in the USA, which receives patient samples from a caregiver, analyses them with DiviTum TKa and then sends a report with the results back to the caregiver.

**Palbociclib** A new type of targeted, selective drug that has been shown to be effective against several forms of cancers, including hormone receptorpositive breast cancer.

**Pemetrexed (Alimta)** is a type of chemotherapy for treating pleural mesothelioma (cancer of the outer covering of the lungs) and non-small cell lung cancer (NSCLC).

Poster session - These are sessions held at a congress or conference with an academic or professional focus to present research information in the form of a paper poster that conference participants may view. A poster session is an event at which many such posters are presented.

**Posters** - These are used to summarize information or research and present it in an attractive way as a means of generating interest in publishing it and sparking discussion at events such as scientific conferences.

**Predictive** Anticipation about what will happen in the future and used in the contexts like the predictive ability of a particular test.

PREDIX study A randomized trial of neoadjuvant

chemotherapy to treat HER2-positive breast cancer that was carried out during the period 2014–2019 at nine Swedish clinics under the supervision of Karolinska Institutet (KI).

Prospective studies Used to study the relationship between various risk factors and a particular disease. This type of study follows individuals both with risk factors and without (the control group), for a period of time into the future. At the end of the study, a comparison is made of the percentage that fell ill in each group.

**PYTHIA study** - A clinical study of patients with metastatic breast cancer. The primary aim of the PYTHIA study is to discover potentially innovative biomarkers for the selection of patients to Palbociclib/Fulvestrant treatment.

**Reimbursement -** Compensation for costs (in this context, it is payment from insurance companies to cover treatment costs)

**RUO Research Use Only -** An ROU product is an IVD (In Vitro Diagnostic) product that is in the

development stage and may only be used for laboratory research and clinical studies.

**Tests (ROU)** - These are tests that are performed on patient samples that have been taken for Research Use Only. It is a service sold primarily to pharmaceutical companies or universities that are conducting research in trials. Biovica receives the samples and sends back analysis reports. For the USA, the tests are conducted with DiviTum TKa at our CLIA laboratory in San Diego and for the EU, from our laboratory in Uppsala.

**Tymidine kinase** is an enzyme (kinase), subclass of phosphotransferase.

Estrogen receptor-positive - To determine whether a patient might benefit from hormone treatment, the tumor is studied to assess whether receptors for either estrogen or progesterone. If so, it is hormone-receptor positive, which is the case for around 70 percent of all breast cancer tumors. It is primarily estrogen that has a stimulating effect on tumor growth.

This report has been reviewed by the company's auditor.

#### Board of Directors' assurance

The Board of Directors and CEO hereby certify that this interim report provides a true and fair summary of the Parent Company's and the Group's operations, earnings and financial position as well as describing any significant risks or uncertainties faced by the Parent Company or any of the companies belonging to the Group.

Uppsala 18 December 2025

Fredrik Alpsten Chairman of the Board

Annika Carlsson Berg Board member

Maria Holmlund Board member

Anders Rylander President/CEO, Board member CP (Niels) Bogerd Board member

Marie-Louise Fjällskog Board member

Jesper Söderqvist Board member

#### Calendar

Interim Report for Q3: November-January 2025/2026 Interim report for Q4: February-April 2025/2026

**Annual Report** 

Interim Report for Q1: May-July 2026/2027

AGM 2026 16 Sept 2026

Interim Report Q2: August-October 2026/2027 Interim Report Q3: November-January 2026/2027 Interim report for Q4: February-April 2026/2027 18 March 2026 17 June 2026 29 June - 3 July, 2026 10 September 2026

17 December 2026 11 March 2027 17 June 2027

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### Biovica – Treatment decisions with greater certainty

Biovica develops and commercializes blood-based biomarker assays for evaluating the effect of cancer treatments. Biovica's assay DiviTum TKa measures cell proliferation by detecting a biomarker in the blood stream. The assay has successfully demonstrated its capabilities to early evaluate therapy effectiveness in several clinical trials. The first application area for DiviTum TKa is evaluation of the treatment effect on metastatic breast cancer. Biovica's vision is that all cancer patients will get an optimal treatment from day one. Biovica collaborates with world-leading cancer institutes and pharmaceutical companies. DiviTum TKa has CE marking and it is registered with the Swedish Medical Products Agency. Biovica's shares are traded on the Nasdaq First North Premier Growth Market (BIOVIC B). FNCA Sweden AB is the company's Certified Adviser, info@fnca.se, +46 (0)8-528 00 399. For more information, please visit <a href="https://www.biovica.com">www.biovica.com</a>.