Interim Report January–March 2026

Cinclus Pharma Holding AB (publ)

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Quarter 1 2026

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Finanancial summary January - March 2026

- Net sales amounted to TSEK 10,036 (0).
- Operating profit (EBIT) amounted to TSEK -90,854 (-47,519).
- The result for the period was TSEK -88,768 (-33,672) and earnings (loss) per share before and after dilution were SEK -1.91 (-0.72) SEK.
- Total cash flow for the period amounted to TSEK -12,712 (-42,133).
- Cash and cash equivalents at the end of the period amounted to TSEK 475,849 (523,899).

22
FTE[1)]

476 MSEK
CASH AND
CASH EQUIVALENTS

91% R&D
OF
OPERATING EXPENSES

[1)] Refers to both employees and consultants. Totally, 17 employees and 27 consultants are affiliated with the company at the end of the period.

General informatin about the report

The information in this report refers to the Group unless otherwise stated. Comparative figures in brackets refer to the corresponding period of the previous year. Comparative figures in brackets for balance sheet items refer to the end of the previous financial year. This report has not been subject to the auditors review. The report has been prepared in a Swedish and an English version. In the event of any discrepancies between the versions, the Swedish version will take precedence.

Upcoming information events

May 21 2026	Annual General Meeting 2026
August 19 2026	Interim Report Q2 2026
November 11 2026	Interim Report Q3 2026

For further information please contact

Christer Ahlberg
Chief Executive Officer
and President
+46 70 675 33 30
ir@cinclupharma.com

Magnus Christensen
Chief Financial Officer
+46 70 675 49 55

The webcast will be held on May 13 2026 at 10:00 via Inderes. Link to the event:
https://cinclus-pharma.events.inderes.com/q1-report-2026
The report is available on the company's website:
https://cinclupharma.com/sv/investerare/finansiella-rapporter/

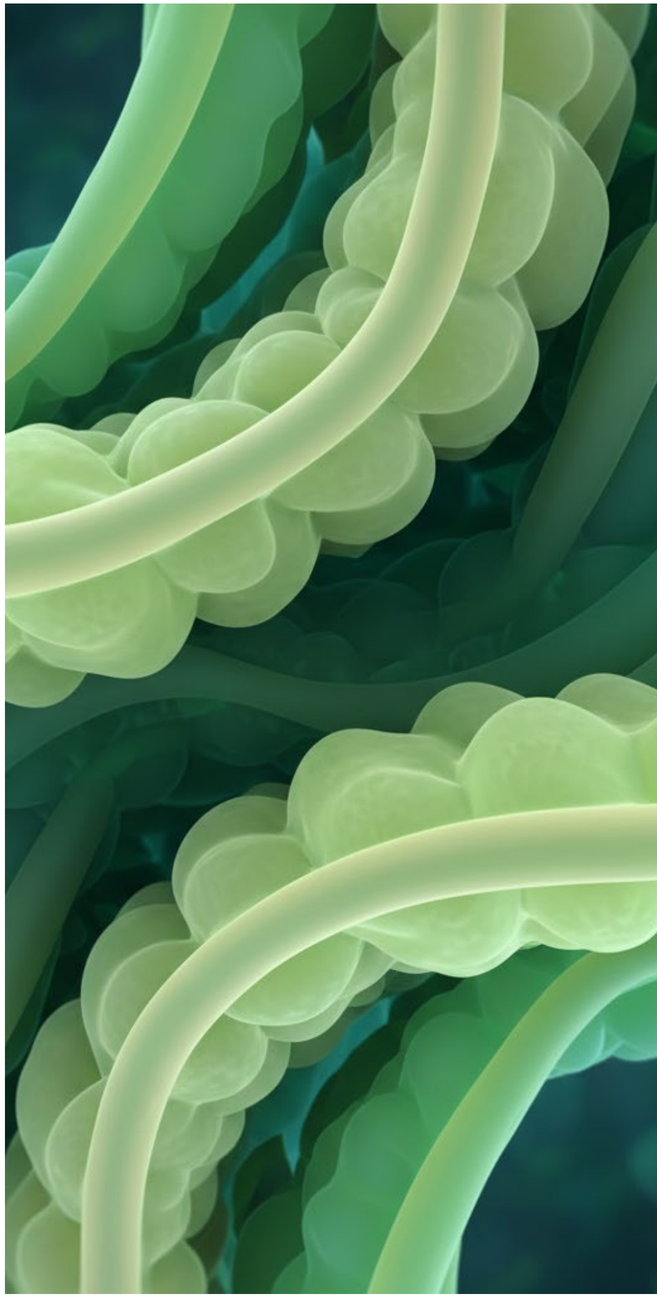
Significant events

Significant events during the period January-March 2026

- At an Extraordinary General Meeting on January 19, Kjell Andersson was elected as a new board member after Peter Unge had announced his wish to step down from his position on the company's board.
- In February, Cinclus Pharma announced that the European Medicines Agency (EMA) had issued a positive opinion following completed scientific advice regarding CMC (Chemistry, Manufacturing and Controls) for linaprazan glurate.
- In February, the FDA issued a positive assessment of the company's preclinical development plan for linaprazan glurate following completed scientific advice.
- In February, Cinclus Pharma announced that the FDA had determined that a single pivotal study will become the new standard for market approval of drugs in the US.
- In March, Cinclus Pharma announced that a financing agreement of EUR 28 million had been entered into with Claret Capital Partners, enabling an earlier start of the final stage of the Phase III program. The financing is structured as a secured loan facility with warrants and convertible instruments.

Significant events after the end of the period

- On April 16, Cinclus Pharma's annual report for 2025 was published, along with the notice convening the Annual General Meeting to be held on May 21, 2026.



Phase III study approaching full enrollment

Patient enrollment in our first Phase III study has continued to progress well and we have good visibility on the timeline with expected topline results in the fourth quarter of this year. The strengthened financial position provides increased flexibility and enables an efficient start to the HEEALING 2 study, the final part of the Phase III program for erosive GERD. During the quarter, Cinclus Pharma also made further regulatory progress toward a future marketing authorization for linaprazan glurate, a drug with the potential to improve quality of life for patients suffering from severe acid related diseases who are not helped by existing treatment options.

HEEALING 1 entering its final phase

Our first Phase III study, HEEALING 1, has progressed according to plan since its launch in October. At present, around 85 percent of the patients in the study have been screened, and more than two thirds have been randomized. We expect the last patient to be screened within the coming month, which means that we have good visibility regarding the timeline for HEALING 1, with topline results expected to be presented during the fourth quarter of this year.

The objective of the study, conducted across eight European countries, is to demonstrate superior healing compared with the proton pump inhibitor lansoprazole in patients with moderate to severe erosive GERD after four weeks of treatment, as well as healing and symptom relief for up to eight weeks.

New financing enables an efficient start to the final step of the program

In March, we entered into a long term structured financing agreement with Claret Capital Partners, Europe's largest independent growth debt fund manager. The financing is structured as a secured loan facility with warrants and convertibles, with conversion price that is more than 80 percent above the current share price,totaling up to EUR 28 million, of which EUR 8 million was drawn during the quarter.

The capital injection enables us to initiate a number of important preparatory processes ahead of the start of the second part of the Phase III program, thereby ensuring a smooth start of our second healing study, HEEALING 2, following the presentation of the results from HEEALING 1.

This includes, among other things, preparatory work related to regulatory applications and study protocols as well as the procurement of a CRO (contract research organization) for the study.

The agreement with Claret also contributes to increased financial flexibility for Cinclus Pharma, both in the short and long term. Under the current business plan, Cinclus Pharma is funded through the third quarter of 2027, while we expect potentially value-driving topline results as early as the fourth quarter of 2026. In addition, the financing will support the remainder of the development program as well as pre-market activities, and is tied to clearly defined milestones that reflect our long-term planning.

Important regulatrory progress

In February, we received positive feedback from the European Medicines Agency (EMA) regarding our CMC (Chemistry, Manufacturing and Controls) strategy. The agency confirmed that the planned CMC approach is well aligned with the requirements for the upcoming marketing authorization application.

Last autumn, Cinclus Pharma received similar feedback from the FDA. Having both the FDA and EMA provide consistent support for our regulatory strategy reinforces the view that Cinclus Pharma is well positioned for the steps ahead. This type of constructive dialogue with regulatory authorities is essential to ensuring quality, predictability, and efficiency in the continued development.

Shortly thereafter, we received positive feedback from the FDA following scientific advice on the preclinical development plan for linaprazan glurate. The FDA confirmed that, based on the available data, no additional toxicology studies are required ahead of the marketing authorization application. This is an important conclusion that reduces risk and contributes to a more time efficient development process.

A clear goal - addressing the medical needs of the patients who suffer the most

Severe erosive reflux disease occurs when stomach acid repeatedly flows back into the esophagus, leading to inflammation, erosions, and persistent symptoms that significantly impact patients' health and quality of life. Existing treatment options on the market do not provide sufficient relief for these chronic conditions, and a more effective therapy would offer patients a meaningful improvement in daily life. The total market is estimated at 19 million patients globally, including 10 million in Europe and the United States.

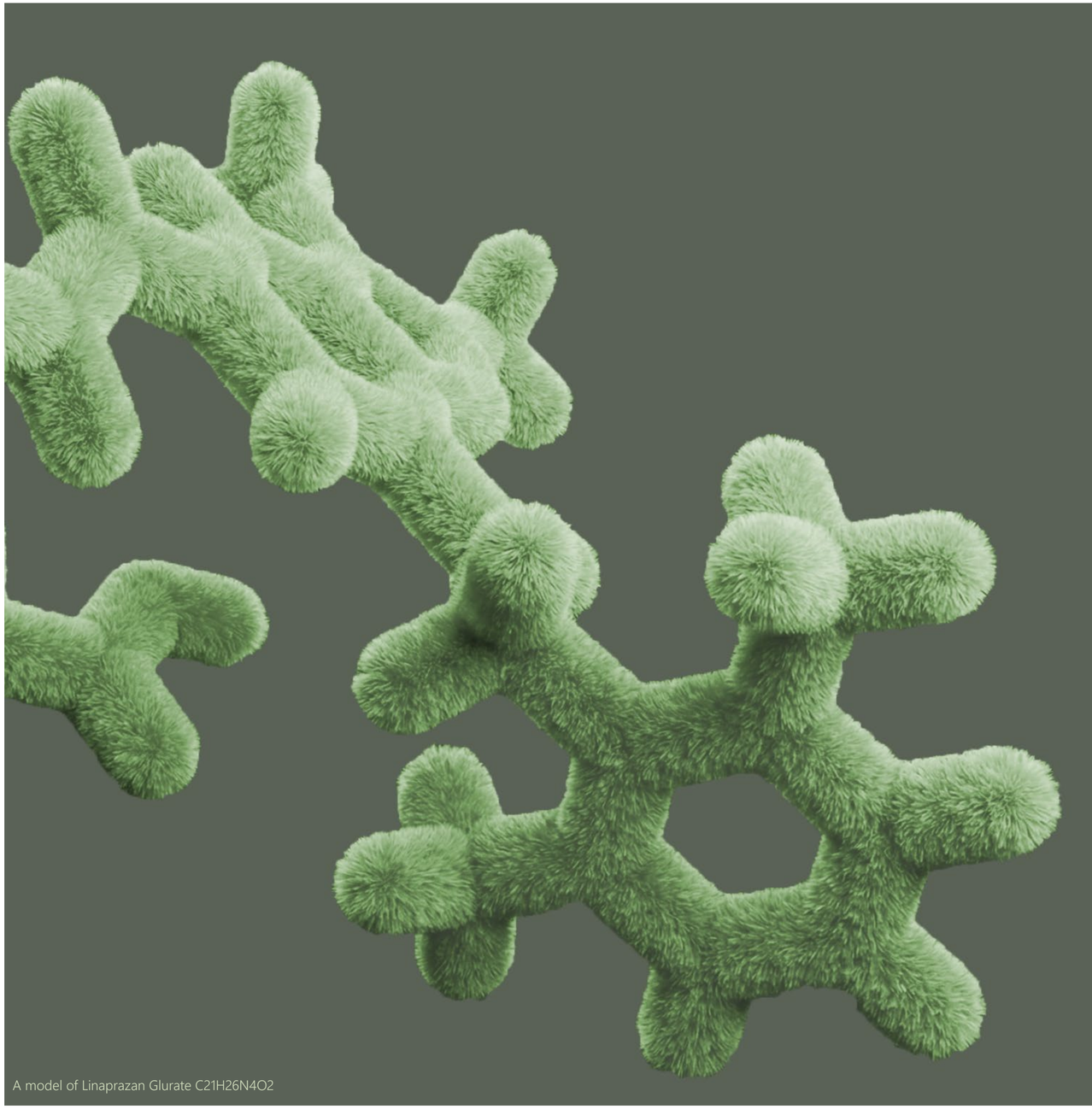
In our study, linaprazan glurate is compared with a proton pump inhibitor, the current standard of care for acid related diseases. In recent years, PCABs (potassium competitive acid blockers) have rapidly gained market share in regions where they have been launched, as they offer improved acid control, which is essential for healing.

Linaprazan glurate is the next generation of PCABs and has demonstrated improved acid control compared with both proton pump inhibitors and other PCABs. In our studies, Linaprazan glurate has shown the ability to maintain gastric pH above 4 for 23[1)] out of 24 hours, providing a unique capacity to deliver effective treatment for patients with severe erosive GERD. With the potential to address a major unmet medical need in a large patient population suffering from severe symptoms and insufficient treatment options, we see significant market potential for linaprazan glurate.

Overall, the first quarter of the year has enabled us to strengthen Cinclus Pharma in several key areas. With a dedicated team and clear focus, we are now moving forward into 2026 with strong determination. Our goal remains unchanged: to offer patients with severe GERD a new and long-awaited treatment option as quickly and responsibly as possible.

Christer Ahlberg,
President and CEO

[1)] Source: Miner P et al, Am J Gastro 2003, Phathom Pharmaceuticals Voquezna (Vonoprazan) FDA labeling 2023, Cinclus Pharma Study CX842A2107 (data on file) Phase I pH control 1,5-24 hours Day 1 and 0-24 hours Day 14 (data on file), Publicly available company reports from Phathom Pharmaceuticals.



A model of Linaprazan Glurate C21H26N4O2

19 million patients worldwide seek medical care for severe erosive reflux disease

Reflux disease affects people worldwide and is a growing health problem. The most common treatment today is proton pump inhibitors (PPIs), but their effect is often insufficient – especially in patients with more severe symptoms and erosive damage. An alternative treatment option, potassium-competitive acid blockers (PCABs), is now gaining ground in several countries.

The key is to control stomach acid

In reflux disease, stomach acid repeatedly flows back into the esophagus, causing irritation, inflammation, and in severe cases erosive damage to the lining. The primary goal of drug treatment is therefore to control stomach acid – that is, to keep gastric pH high for most of the day so the acid does not damage the tissue.

Today's standard treatment is not sufficient

Proton pump inhibitors (PPIs) have long been the most common medical treatment for reflux disease. These drugs block the activity of the proton pump in the acid-producing cells of the stomach, thereby reducing acid production.

PPI treatment often provides symptom relief and improvement for many patients. However, in some cases acid control is not sufficient throughout the entire day, which can result in incomplete healing of the esophageal lining and a slower therapeutic effect.

Linaprazan shows strong and sustained acid control

In recent years, a new class of drugs – potassium-competitive acid blockers (PCABs) – has been introduced in several markets. PCABs have a different mechanism of action than PPIs and can provide faster and more stable suppression of acid production. First-generation PCABs are now beginning to replace standard treatment in these markets.

Cinclus Pharmas Cinclus Pharma's drug candidate linaprazan glurate is in late-stage clinical development and represents the next generation of PCAB therapy. Previous clinical studies show that linaprazan glurate is more effective than first-generation PCABs, providing strong and long-lasting acid control. This level of control is critical for effective treatment of reflux disease and also supports significant commercial potential.

130 million

130 million people worldwide live with some form of reflux disease

19 million patients worldwide seek medical care for severe erosive reflux disease

19 million

10 million

The number of patients is estimated at 10 million in the United States and Europe

The commercial strategy combines partnerships, patent protection, and growth

Cinclus Pharma's commercial strategy is built on a clear unmet medical need and a large global market. Through clinical development, strong intellectual property protection, and strategic partnerships, the company is creating the foundation for long-term value and new treatment options for patients with severe erosive reflux disease.

A partner-based path to market

In the second quarter of 2025, Cinclus Pharma entered into a strategic collaboration and licensing agreement with Zentiva, a leading European pharmaceutical company. The agreement covers the commercialization of linaprazan glurate across the EEA, including the United Kingdom and Switzerland. The total deal value amounts to up to EUR 220 million and includes upfront payments, regulatory and commercial milestone payments, as well as well as royalties of approximately 20 percent on future sales.

In Asia, Cinclus Pharma has previously licensed the rights to linaprazan glurate to Jiangsu Sinorda Biomedicine Co. Ltd for development and commercialization in China and other selected regions. In December 2025, linaprazan glurate was included on China's National Reimbursement Drug List for the treatment of gastroesophageal reflux disease, enabling commercial launch in 2026.

For other markets, including the United States, Cinclus Pharma continues to evaluate the most value-creating route to market.

Strong patent protection and data exclusivity

Strong intellectual property protection is a core element of Cinclus Pharma's commercial strategy. Linaprazan glurate is protected by multiple patents, including a polymorph patent in the United States valid until 2042 and a formulation patent in Europe valid until 2040.

In addition to patent protection, Cinclus Pharma also relies on regulatory data exclusivity, which provides effective protection against generic competition following potential market approval. In Europe, this provides up to 10–11 years of data exclusivity from the date of approval, while the corresponding protection in the United States is five years. In addition, the company may be eligible for a further five years of exclusivity in the United States if linaprazan glurate is approved for the treatment of *H. pylori* as the first indication.

Focus on long-term value creation

The combination of strong intellectual property protection, established partnerships, and a flexible market strategy provides Cinclus Pharma with a solid foundation for the next steps. With a clear focus on quality, long-term value, and collaboration, the company continues to advance linaprazan glurate toward market and toward patients in need of improved treatment options.

H. pylori indication expands the potential use of linaprazan glurate

Cinclus Pharma is primarily developing linaprazan glurate for the treatment of severe erosive reflux disease. In parallel, the company is also evaluating the potential use of linaprazan glurate as part of the treatment of *Helicobacter pylori* infection.

*H. pylori i*s a bacterium that infects the stomach and is a common cause of peptic ulcers and chronic gastritis. The World Health Organization (WHO) classifies *H. pylori* as a cancer-causing bacterium, and infection should therefore be treated. Current treatment typically consists of a combination of acid-suppressing medication and antibiotics.

Linaprazan glurate has the potential to provide stable and long-lasting acid control, which may improve treatment outcomes for *H. pylori* infection. Cinclus Pharma has also received regulatory waivers from pediatric studies for this indication, simplifying future development.

Approval for an *H. pylori* indication could, over time, expand the clinical use of linaprazan glurate and support additional regulatory data exclusivity.

Linaprazan glurate is in late-stage clinical development

In 2025, linaprazan glurate entered Phase III, marking an important milestone in its clinical development. The program is based on a comprehensive clinical data package from earlier studies demonstrating both effective acid control and high healing rates.

Phase I: Effective acid control

Across multiple Phase I studies, linaprazan glurate has demonstrated dose-dependent and sustained acid control. Gastric pH, used as a biomarker, provides an early and reliable indication of future clinical efficacy. For patients with severe erosive reflux disease, where esophageal damage is caused by prolonged exposure to gastric acid, effective acid suppression is critical to enable healing.

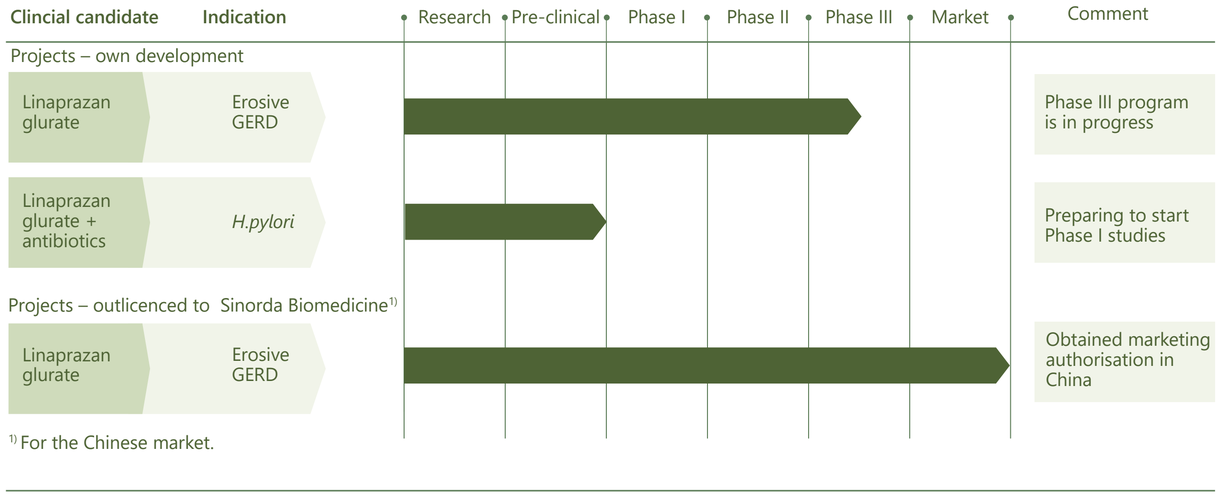
The relationship between gastric pH and healing is well established. When gastric pH is maintained above 4 for most of the day, conditions for esophageal healing improve significantly. In one of Cinclus Pharma's Phase I studies, linaprazan glurate maintained gastric pH above 4 for 96% of the day at the dose now used in the Phase III program. This level of acid control is unusually high and unique compared to currently approved therapies worldwide, making linaprazan glurate particularly promising for patients with erosive reflux disease.

Phase 2: Rapid healing

In the company's Phase II study, a healing rate of 93% was achieved in patients with severe erosive reflux disease in the optimal dose group. In comparison, the healing rate for the reference proton pump inhibitor (PPI) was 38%. These results demonstrate a rapid and robust treatment effect, even in the most difficult-to-treat patient populations, and provide strong support for continued clinical development.

Phase 3: Confirm efficacy at scale

In the Phase III program, linaprazan glurate is being evaluated in large, comparative studies against the current standard of care for erosive reflux disease. The objective is to confirm that the strong acid control and rapid, effective healing observed in earlier studies can also be demonstrated in a broader patient population and in a manner that meets regulatory requirements.



The share

Cinclus Pharma's share (CINPHA) has been listed on Nasdaq Stockholm since June 20, 2024.

The opening price January 2 2026 was SEK 19.26 per share. The closing price on the last trading day in March was SEK 16.88 per share.

The average volume-weighted share price during the first quarter was SEK 17.51 per share. The market capitalization on the last trading day in March was MSEK 800.

The company had 47,392,219 outstanding shares of which 46,537,789 are ordinary shares and 854,430 are C shares which give the right to 1/10 vote of an ordinary share. The C-shares are held by Cinclus Pharma Holding (publ).

At the end of the first quarter, the company had approximately 4,000 shareholders.

Trading	**Nasdaq Stockholm**
Ticker	**CINPHA**
ISIN	**SE0020388577**
LEI-code	**549300TJBPSNZ3DO6B42**
Share price at 2026-03-31	**16.88 SEK**
Market cap. 2026-03-31	**800 MSEK**

Owner information at year-end

Shareholding in the company at the end of the period	Number of shares	Share (%)
Trill Impact Ventures	3,721,221	7.9%
Fjärde AP-fonden	3,700,000	7.8%
Movestic Livförsäkring AB	2,342,397	4.9%
Linc AB	2,318,322	4.9%
Peter Unge via company	2,061,075	4.3%
Kjell Andersson via company	1,908,000	4.0%
Futur Pension Försäkringsaktiebolag	1,666,056	3.5%
Mikael Dahlström estate	1,569,613	3.3%
Nordnet Pensionsförsäkring	1,367,436	2.9%
Nylof Holding AB	1,164,575	2.5%
Lennart Hansson via company	1,084,771	2.3%
Eir Ventures I AB	898,750	1.9%
Cinclus Pharma *	854,430	1.8%
Avanza Pension	850,829	1.8%
Postamentet Holding AB	636,512	1.3%
Fifteen largest shareholders	***26,143,987***	***55.2%***
Others	*21,248,232*	*44.8%*
Total	***47,392,219***	***100.0%***

* Refers to C shares which give the right to 1/10 vote.

Key Share Data

	Quarter 1		Year
Share data	**2026**	**2025**	**2025**
Net income / loss, TSEK	–88,768	–33,672	–183,972
Cash flow for the period, TSEK	–12,712	–42,133	–77,836
Number of shares at the beginning of the period	46,537,789	46,537,789	46,537,789
Number of shares at the end of the period	46,537,789	46,537,789	46,537,789
Average number of shares	46,537,789	46,537,789	46,537,789
Number of warrants at the beginning of the period*	697,737	1,051,897	1,051,897
Number of warrants at the end of the period*	682,955	769,737	697,737
Average number of warrants*	687,772	942,168	793,909
Share capital at the end of the period, TSEK	920	920	920
Equity at the end of the period, TSEK	281,052	511,625	369,391
Earnings per share before dilution, SEK	–1.91	–0.72	–3.95
Earnings per share after dilution, SEK	–1.91	–0.72	–3.95
Equity per share, SEK	6.04	10.99	7.94
Cash flow for the period per share, SEK	–0.27	–0.91	–1.67

* Number of warrants is recalculated so that all programs must meet the 1:1 conversion condition

Financial summary, January – March 2026

	Quarter 1		Year
Financial summary for the group	**2026**	**2025**	**2025**
Net sales, TSEK	10,036	–	57,470
Operating income / loss (EBIT), TSEK	−90,854	−47,519	−199,558
Net income / loss, TSEK	−88,768	−33,672	−183,972
Operating expenses, TSEK	−100,705	−46,249	−255,650
R&D expenses vs. operating expenses, %	91%	83%	84%
Cash flow from operating activities, TSEK	−95,641	−41,793	−74,647
Cash and cash equivalents at the end of the period, TSEK	475,849	523,899	487,254
Quick ratio, %	289%	1271%	392%
Equity, TSEK	281,052	511,625	369,391
Equity ratio, %	55%	92%	68%
Average number of employees during the period	12	18	19
Average number of shares, before dilution	46,537,789	46,537,789	46,537,789
Average number of shares, diluted	46,564,654	46,561,439	46,564,368
Number of shares at the end of the period, before dilution	46,537,789	46,537,789	46,537,789
Number of shares at the end of the period, diluted	46,564,654	46,561,439	46,564,368
Earnings per share, before dilution 1), SEK	−1.91	−0.72	−3.95
Earnings per share, diluted 1), SEK	−1.91	−0.72	−3.95

1) The period's earnings per share before and after dilution are defined in IFRS. Other key figures in the above table are alternative key figures and thus not defined in IFRS, see further section for definitions and reconciliation of key figures and alternative key figures later in this report.

Net sales

Net sales amounted to TSEK 10,036 (0) during the quarter. The revenue consists partly of licensing revenue from the partnership with Zentiva for commercialization of linaprazan glurante in Europe, and partly from Sinorda in China.

Operating expenses

Research and development expenses

Research and development (R&D) expenses during the quarter amounted to TSEK -91,273 (-38,436), which correspond to an increase of TSEK 52,837 or 137 %. The research and development expenses related mainly to the ongoing Phase III study, where patients are recruited and screened while the corresponding period last year consisted of expenses for the start up of the Phase III program.

Administrative expenses

The administrative expenses amounted to TSEK -9,432 (-7,813) for the quarter, which correspond to an increase of TSEK 1,620 or 21 %. The increased expenses are largely due to concultancy fees and increased rental costs due to the move to new premises.

Other operating income and expenseses

Other operating income and expenses amounted to net TSEK -185 (-1,270) during the quarter, corresponding to a change of TSEK 1,085. Other operating income and expenses consist of realized and unrealized exchange rate effects on operating receivables and liabilities.

Operating income/loss (EBIT)

The Group's operating income/loss for the quarter amounted to TSEK -90,854 (-47,519), corresponding to a change of TSE K -43,335.

Financial items

Financial income and expenses (net finacial income/expense) amounted to TSEK 2,088 (13,920) during the quarter, corresponding to a change of TSEK -11,833. The financial income for the interim period attributed to interest income on bank funds and the exchange rate development of the Swedish krona while the financial expenses are attributed to the convertible loan communicated in March.

Income tax

The Group recognized a tax expense of TSEK -1 (-73) during the quarter. The tax consisted of Swiss federal and cantonal tax.

Net income/loss

The Group reported net income/loss after tax of TSEK -88,768 (-33,672) for the quarter. This corresponded to a change of TSEK -55,096.

Equity and indebtedness

Equity in the Group as of March 31, 2026 amounted to TSEK 281,052 compared to TSEK 369,391 at the end of previous year 2025, a decrease of TSEK 88,339.

Non-current liabilities at the end of the peroid amounted to TSEK 56,629 (40,373). The non-current liabilities consisted mainly of non-current contractual liabilities, attributed to the out-licensing of the commercial rights to Zentiva in the second quarter of 2025.

Current liabilities in the Group at the end of the period amounted to TSEK 173,958 (136,792), an increase of TSEK 37,166.
The increase attributed mainly to current interest bearing liabilities of TSEK 45,788 (0) as well as convertibles and warrants of TSEK 32,945 (0) arising from the first tranche of the convertible loan arrangement. Furthermore, current liabilities consisted of trade payables of TSEK 29,522 (48,464), lease liabilities of TSEK 3,174 (3,111), tax liabilities of TSEK 1 (207), other liabilities of TSEK 3,615 (9,656), current contract liabilities of TSES 35,708 (54,524) and accrued expenses of 23,205 (20,827).

Liquid funds and cash flow

Cash and cash equivalents at the end of the period amounted to TSEK 475,849 (487,254), a decrease of TSEK 11,405 compared to the end of 2025. During the period, the company received the first drawn down of the convertible loan of approximately MSEK 86 before transaction costs.

Cash flow from operating activities before change in working capital was TSEK -88,953 (-46,077) for the quarter.

Cash flow from operating activities including change in working capital amounted to TSEK -95,641 (-41,793) for the quarter.
Cash flow from investing activities amounted to TSEK -24 (0) for the quarter attributed to equipment.

Cash flow from financing activities amounted to TSEK 82,953 (-340) for the quarter attributed to amortization of lease liabilities and convertible loan received.

The total cash flow for the quarter amounted to TSEK -12,712 (-42,133).

Financing

Following the settlement of the convertible loan, the Company estimates as of March 31, 2026 that the current working capital is sufficient to the the third quarter of 2027, after the read out of the first Phase III program. Read out is assessed per the date of this report, following the current development plan to take place during the fourth quarter of 2026.

Parent company

Cinclus Pharma Holding AB (publ), reg.no. 559136–8765, is the parent company of the Group. The business consists of work with pre-clinical and clinical development, marketing, administrative and corporate management functions. The parent company has two wholly owned subsidiaries, one in Switzerland and one in Sweden, which together form the Group.

The total revenues of the parent company amounted to TSEK 10,192 (44) for the quarter and consist of revenues from Zentiva and Sinorda.

Operating income/loss for the quarter amounted to TSEK -90,794 (-46,271)

Net financial income/expense for the quarter amounted to TSEK 2,640 (12,727). The positive net financial income for the interim period attributed to interest income on bank funds and the currency exchange development of the swedish krona.

Income/loss for the quarter amounted to TSEK -88,154 (-33,544).

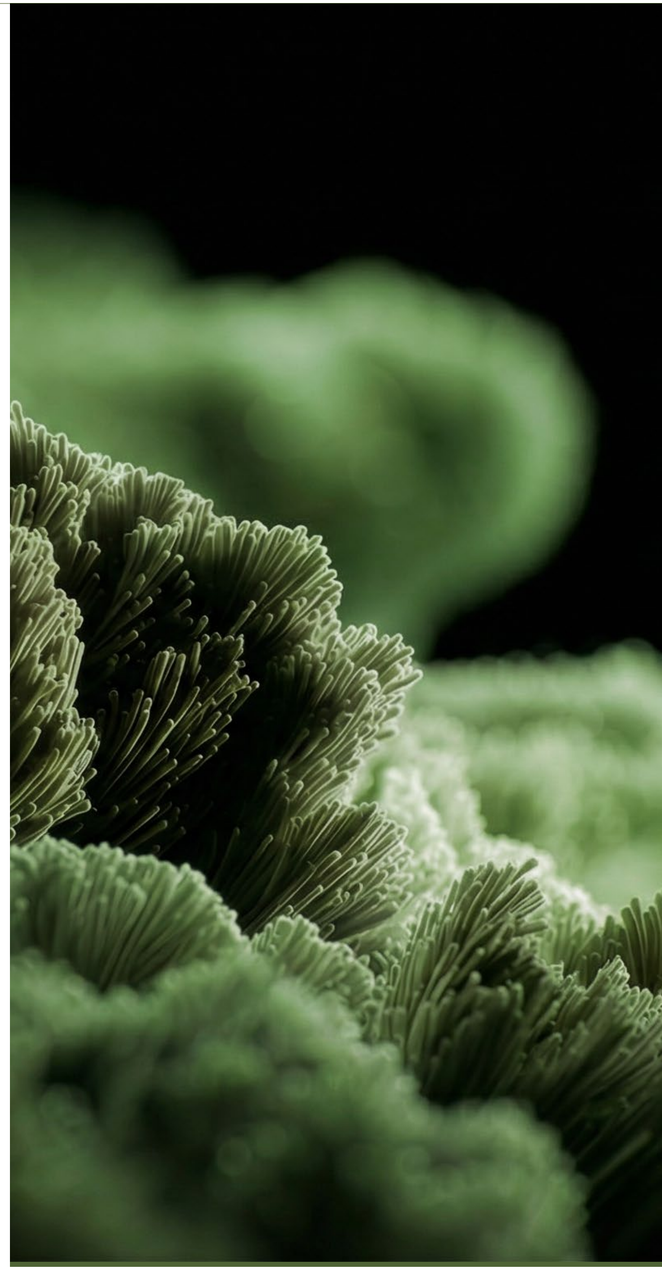
With the transfer of patents and IP rights to the parent company from the swiss subsidiary as of January 1, 2022, the parent company recognizes an intangible asset of TSEK 320,463 (320,463).

Cash and cash equivalents at the end of the period amounted to TSEK 459,156 compared to TSEK 480,535 at the end of previous year, a decrease of TSEK 21,379.

Total Equity in the parent company as of March 31, 2026 amounted to TSEK 593,954 compared to TSEK 681,859 at the end of previous year, corresponding to a decrease of TSEK 87,905. Share capital amounted to TSEK 920 (920). The company had on the balance sheet day, March 31, 2026, 46,537,789 ordinary shares and 854,430 C-shares.

Non-Current liabilities in the parent company amounted to TSEK 52,666 (35,587) in the end of the period, attributed to contract liabilities deriving from the Zentiva deal in May 2025.

Current liabilities in the parent company amounted to TSEK 172,709 (125,373) in the end of the period, an increase of TSEK 47,336 mainly due to the convertible loan.



Other information

Personnel

At the end of the quarter, the number of employees was 17, compared to 18 in the same period previous year. The average number of employees during the quarter was 12, compared to 18 in the same period last year. All employees are employed by the parent company. In addition, the company had 27 consultants attached to the company, at the end of the quarter.

Risks

As the company is dependent on additional financing to continue the development of linaprazan glurate in the long term, the refinancing risk is described below. For other risks, reference is made to the description of the Group's significant financial and business risks in the Directors' Report and Note 19 in the Annual report for 2025.

Refinancing risk

Refinancing risk refers to the risk that cash and cash equivalents are not available and that financing can only be obtained partially or not at all, or at an increased cost. The Group is currently financed with equity and loan and the refinancing risk has been significantly reduced in view of the new share issue that took place in connection with the listing of the company's share on Nasdaq Stockholm on June 20, 2024, the out-licensing of the European commercial rights to Zentiva and the settlement of the convertible loan this quarter. In the longer term, the Group is in need of more extensive financing to be able to conduct and implement a second Phase III study and registration of the erosive GERD indication.

Consolidated income statement in summary

(TSEK)	Note	Quarter 1 2026	Quarter 1 2025	Year 2025
Revenues				
Net sales	4	10,036	–	57,470
Operating expenses				
Administrative expenses		–9,432	–7,813	–57,248
Research and development expenses		–91,273	–38,436	–198,402
Other operating income and expenses		–185	–1,270	–1,379
Operating income / loss		**–90,854**	**–47,519**	**–199,558**
Net financial income/expense	8	2,088	13,920	15,847
Income / loss before tax		**–88,767**	**–33,599**	**–183,710**
Income tax		–1	–73	–262
Net income/ loss for the period attibutable to parent company shareholders		**–88,768**	**–33,672**	**–183,972**
Earnings per share, calculated on earnings attributable to the parent company ordinary shareholders (SEK):				
Before dilution		–1.91	–0.72	–3.95
Diluted		–1.91	–0.72	–3.95

Consolidated statement of comprehensive income in summary

(TSEK)	Note	Quarter 1 2026	Quarter 1 2025	Year 2025
Net income / loss for the period		**–88,768**	**–33,672**	**–183,972**
Other comprehensive income /loss				
Items that can later be reclassified to the income stament:				
Translation differences from operations abroad		179	–10,828	–5,136
Other comprehensiveincome / loss, net after tax		**179**	**–10,828**	**–5,136**
Comprehensive income / loss for the period		**–88,588**	**–44,499**	**–189,108**
Comprehensive income/ loss for the period as a whole attributable to the parent company shareholders		**–88,588**	**–44,499**	**–189,108**

Consolidated statement of financial position in summary

(TSEK)	Note	Mar 31, 2026	Mar 31, 2025	Dec 31, 2025
ASSETS				
Property, plant and equipment				
Inventories		990	38	1,046
Right-of-use assets		7,566	720	8,360
Financial assets				
Other non-current assets		296	1	296
Total fixed assets		**8,851**	**759**	**9,703**
Current assets				
Trade receivables		1,418	–	16,062
Current tax assets		4	–	–
Other receivables		4,054	1,814	4,992
Prepaid expenses and accrued income		21,463	28,962	28,546
Cash and cash equivalents		475,849	523,899	487,254
Total current assets		**502,788**	**554,675**	**536,854**
TOTAL ASSETS		**511,639**	**555,434**	**546,556**

(TSEK)	Note	Mar 31, 2026	Mar 31, 2025	Dec 31, 2025
EQUITY ALD LIABILITIES				
Equity				
Share capital		920	920	920
Other contributed capital		1,297,740	1,297,740	1,297,740
Translation difference		23,710	17,839	23,530
Retained earnings including profit for the period		–1,041,318	–804,874	–952,800
Equity attributable to the parent company shareholders		**281,052**	**511,625**	**369,391**
Non-current liabilities				
Non-current lease liabilities		3,963	161	4,787
Non-current contract liabilities		52,666	–	35,587
Total non-current liabilities		**56,629**	**161**	**40,373**
Current liabilities				
Current interest bearing liabilities	8	45,788	–	–
Derivative convertible and warrants	8	32,945	–	–
Trade payables		29,522	14,389	48,464
Lease liabilities, short-term		3,174	222	3,111
Current tax liabilities		1	6,829	207
Other liabilities		3,615	3,736	9,656
Current contract liabilities		35,708	–	54,527
Accrued expenses		23,205	18,473	20,827
Total current liabilities		**173,958**	**43,648**	**136,792**
TOTAL EQUITY AND LIABILITIES		**511,639**	**555,434**	**546,556**

Consolidated statement of changes in equity in summary

(TSEK)	Mar 31, 2026	Mar 31, 2025	Dec 31, 2025
Opening balance	**369,391**	**555,330**	**555,330**
Comprehensive income / loss for the period	−88,588	−44,499	−189,108
Share-related remuneration, staff vested value	249	794	3,170
Closing balance	**281,052**	**511,625**	**369,391**

Consolidated statment of cash flow in summary

		Quarter 1		Year
(TSEK)	Note	**2026**	**2025**	**2025**
Operating activities				
Operating income / loss		−90,854	−47,519	−199,558
Adjustments for items not included in the cash flow				
Depreciations		875	448	2,379
Exchange rate differences		0	10	−68
Share-based remuneration		249	794	3,170
Interest received		2,078	460	12,216
Interest paid		−1,090	−55	−449
Taxes paid		−211	−215	−7,279
Cash flow from operating activities before change in working capital		**−88,953**	**−46,077**	**−189,591**
Cash flow from change in working capital				
Increase(-)/Decrease (+) of operating receivables		22,045	5,644	−16,047
Increase(+)/Decrease (-) of account payables		−20,459	−4,538	29,536
Increase (+) /Decrease (-) of contract liabilities		−1,740	–	90,114
Increase(+)/Decrease (-) of other operating liabilities		−6,534	3,178	11,340
Cash flow from operating activities		**−95,641**	**−41,793**	**−74,647**
Investing activities				
Investments in tangible assets		−24	–	−1,134
Investments in financial assets		–	–	−295
Cash flow from investing activities		**−24**	**–**	**−1,429**
Financing activities				
Amortisation of lease liabilities		95	−340	−1,760
Received convertible loan	8	82,858	–	–
Cash flow from financing activities		**82,953**	**−340**	**−1,760**
Cash flow for the period		**−12,712**	**−42,133**	**−77,836**
Cash and cash equivalents at the beginning of the period		**487,254**	**566,716**	**566,716**
Exchange rate differences in cash and cash equivalents		1,307	−684	−1,627
Cash and cash equivalents at the end of the period		**475,849**	**523,899**	**487,254**

Parent company income statement in summary

(TSEK)	Note	Quarter 1 2026	Quarter 1 2025	Year 2025
Revenues				
Net sales	4	10,192	44	62,894
Operating expenses				
Administrative expenses		−9,132	−7,734	−62,124
Research and development expenses		−91,348	−37,760	−198,220
Other operating income and expenses		−506	−822	−1,155
Operating income / loss		**−90,794**	**−46,271**	**−198,604**
Results from shares in group companies		–	–	68,139
Net financial income/expense	8	2,640	12,727	13,436
Income / loss after financial items		**−88,154**	**−33,544**	**−117,029**
Group contribution		–	–	–
Income / loss before tax		**−88,154**	**−33,544**	**−117,029**
Corporate tax		–	–	–
Net income / loss for the period		**−88,154**	**−33,544**	**−117,029**

In the parent company, there are no items that are reported as other comprehensive income, which is why the total comprehensive income for the period corresponds to the period's result.

Parent company balance sheet in summary

(TSEK)	Note	Mar 31, 2026	Mar 31, 2025	Dec 31, 2025
ASSETS				
Intangible assets				
Concessions, patents, licenses, etc.		320,463	320,463	320,463
Property, plant and equipment				
Inventories		990	38	1,046
Financial assets				
Shares in group companies		2,025	88,543	2,025
Other non-current assets		296	–	296
Total fixed assets		**323,774**	**409,044**	**323,831**
Current assets				
Trade receivables		1,418	–	399
Receivables in group companies		10,644	3,590	7,911
Prepaid expenses and accrued income		2,262	1,808	969
Other current receivables		22,075	24,857	29,174
Cash and cash equivalents		459,156	517,023	480,535
Total current assets		**495,555**	**547,278**	**518,988**
TOTAL ASSETS		**819,329**	**956,322**	**842,818**

(TSEK)	Note	Mar 31, 2026	Mar 31, 2025	Dec 31, 2025
EQUITY AND LIABILITIES				
Equity				
Restricted equity				
Share capital		920	920	920
Non restricted equity				
Share premium fund		1,297,509	1,297,509	1,297,509
Retained earnings		−616,320	−501,917	−499,541
Profit or loss for the period		−88,154	−33,544	−117,029
Equity attributable to the parent company's shareholders		**593,954**	**762,968**	**681,859**
Non-current liabilities				
Non-current contract liabilities		52,666	–	35,587
Total non-current liabilities		**52,666**	**–**	**35,587**
Current liabilites				
Current interest bearing liabilities	8	45,788	–	–
Derivative convertible and warrants	8	32,945	–	–
Trade payables		29,522	14,347	41,728
Current tax liabilities		1	–	–
Liabilities to group companies		1,912	156,497	1,937
Other liabilities		3,615	3,736	6,352
Current contract liabilities		35,708	–	54,527
Accrued expenses		**23,219**	**18,775**	**20,828**
Total current liabilities		**172,709**	**193,354**	**125,373**
TOTAL EQUITY AND LIABILITIES		**819,329**	**956,322**	**842,818**

Notes to the financial information

Note 1 General inforamtion

Cinclus Pharma Holding AB (publ), (hereafter Cinclus Pharma) corporate registration number 559136–8765 is a limited company registrered in Sweden with its registered office in Stockholm. The address of the head office is Kungsbron 1, 111 22 Stockholm, Sweden. The company is listed on Nasdaq Stockholm since June 20, 2024 and the object of the company's operations is to develop and commercialize pharmaceuticals. Cinclus Pharma Holding AB (publ) is the parent company in the Group Cinclus Pharma, which consists of the parent company and its two subsidiaries (hereafter the Group). Unless otherwise specifically stated, all amounts are reported in thousands of kronor (TSEK). All amounts are, unless otherwise stated, rounded to the nearest thousand. Figures in parentheses refer to the comparison period.

Except for Debt to credit institutions, the Group's financial assets and liabilities that are recognized at amortized cost are deemed to be a reasonable estimate of the fair value. They refer essentially to short-term receivables and liabilities, whereby the discounting effect is immaterial.

Note 2 Accounting principles

The most important accounting principles applied when these consolidated accounts have been prepared are stated below. These principles have been applied consistently for all periods presented, unless otherwise stated. The consolidated financial statements have been prepared in accordance with the Annual Accounts Act (1995:1554), RFR 1 Supplementary accounting rules for groups, and the International Financial Reporting Standards (IFRS) and interpretations from IFRS Interpretations Committee (IFRS IC) as established by the European Union. This interimreport has been prepared in accordance with IAS 34 Interim Financial Reporting and the Annual Accounts Act.

The parent company interim report has been prepared in accordance with the Annual Accounts Act and Swedish Corporate Reporting Board recommendation RFR 2.

Applied accounting principles and explanations for these can be found and are consistent with those described in the 2025 annual report. The accounting principles have been extended to also include the accounting and valuation of the Group's convertible loan.

Classification and valuation of financial liabilities
The Group's financial liabilities are classified as amortised cost with the exception of derivative instruments, see below. Financial liabilities recognized at amortized cost are initially measured at fair value, net of transaction costs. After the first accounting session, they are valued at amortised cost using the effective interest method.

Convertible Debentures
The convertible debentures issued can be converted into company shares by the counterparty exercising the option to convert the debt into shares. The conversion right is regulated by an exchange of a fixed number of shares for a fixed amount in EUR, i.e. a currency other than the functional currency. The conversion right therefore does not meet the criteria for classification as equity but is reported as a derivative instrument. The conversion option have equity-like characteristics and is thus not closely related to the debt.

The convertible debentures are reported as a composite financial instrument, divided into a debt component and a derivative component (embedded derivative). At the time of initial recognition, the segregated derivative is measured at fair value, and the remaining amount received is allocated to the liability component. In the subsequent accounting, the debt component is valued at amortised cost using the effective interest method. At the time of initial accounting and at subsequent accounting, the derivative component is measured at fair value through profit or loss through the application of generally accepted valuation methods. The liabilities component and the derivatives component are recognised on the balance sheet as current liabilities, as the counterparty has the right to convert at any time during the maturity. Changes in the fair value of the derivative component are recognised in the income statement in net financial items.

Derivatives
Derivatives are initially reported at fair value at the time of entry and are subsequently revalued at fair value at each reporting date. The resulting profit or loss is immediately recognized in the income statement.

A derivative with positive fair value is recognized as a financial asset, while a derivative with negative fair value is recognized as a financial liability. Derivatives are not net recognized in financial statements unless the Group has both a legally binding right and intention to net reduce. A derivative is presented as a long-term asset or a long-term liability if the remaining maturity of the instrument is longer than 12 months and it does not mature or be settled within 12 months. Other derivatives are presented as short-term.

Embedded derivatives
An embedded derivative is a component of a composite instrument that also includes a non-derivative hosting contract and has the effect of varying some of the cash flows of the composite instrument in a similar manner to that of a stand-alone derivative.

Derivatives embedded in composite instruments where the host contract is not a financial asset within the scope of IFRS 9 (e.g. financial liabilities) are separated from the host contract in cases where the derivative meets the definition of a derivative, its risks and nature are not closely related to the risks and nature of the host contract, and the composite instrument as a whole is not measured at fair value through profit or loss.

Warrants
The warrants issued in connection with the issuance of the convertible loan are settled through an exchange of a fixed number of shares for a fixed amount in EUR, i.e. a currency other than the functional currency. The warrants therefore do not meet the criteria for classification as equity but are reported as derivative instruments.

Judgements and estimates

To prepare reports in accordance with IFRS requires the use of some important estimates for accounting purposes. Furthermore, management is required to make certain judgments when applying the Group's accounting principles. The areas that include a high degree of assessment that are complex or such areas where assumptions and estimates are of significant importance for the consolidated accounts, have been reported in the Group's annual report for 2025.

Going concern principle

This interim report has been prepared with the assumption that the company has the ability to continue as a going concern for the next 12 months in line with the going concern principle.

See further sections on financing, risks and risk management and note 3.

Note 3 Risks and risk management

Cinclus Pharma's operations, results and position are affected by a number of risk factors that are described in detail in the company's prospectus prepared in connection with the listing of the company's share on Nasdaq Stockholm on June 20, 2024 but also in the annual report for 2025.

The risks and associated risk management considered in the preparation of this interim report apply to all periods and are consistent with what is presented in the risk factors section in the annual report for 2025.

With the new share issue in connection with the listing of the company's shares on Nasdaq Stockholm, the partnership with Zentiva and the convertible loan the refinancing risk has been reduced.

Refinancing risk refers to the risk that liquid funds are not available, and that financing can only be obtained partially or not at all, alternatively at an increased cost. The Group is currently financed with equity. In the longer term, the Group is in need of more extensive financing. It can not therefore be ruled out that the Group will be exposed to risks related to for example external loan financing.

Note 4 Net sales

Net sales are based on the license agreement with Zentiva k.s., an European pharmaceutical company, regarding the commercial rights of linaprazan glurate in Europe.

The agreement with Zentiva includes an upfront payment, regulatory and commercial milestone payments, sales milestones and royalties on Zentiva's future product sales revenue of linaprazan glurate.

For the quarter, the net sales relates to parts of the upfront payment Cinclus Pharma received upon signing of the agreement with Zentiva. The upfront payment has been allocated over the estimated period of time that the Phase III program runs.

Net sales are also based on the agreement between Cinclus Pharma and its Chinese partner Sinorda Biomedicine Co. Ltd. The income refers to license related revenues that Sinorda Biomedicine has received from out-licensing to its partner in China, HuaDong Medicine Co. Ltd.

Note 5 Incentive programs

The summary table below shows the current programs as of the balance sheet date:

Option programs

Program	Opening balance Jan 2026	Allocated options	Expired options	Closing balance Mar 2026	Terms	Corresponding number of shares	Exercise price/ option (SEK) *
QESO 2022	4,450	–	–	4,450	1:80	356,000	47.33
QESO 2024	51,737	–	–14,782	36,955	1:1	36,955	47.33
ESOP 2024/2027 series 1	290,000	–	–	290,000	1:1	290,000	54.60
Total						**682,955**	

* The exercise prise is recalculated in accordance with the split of the company's shares, which was resolved upon the extraordinary general meeting on 29 May 2023.

QESO = Qualified Employee Stock Options
ESOP = Emloyee Stock Option Program

Performance share program

		Employees per category and series		Investment in number of shares per category			Max. share rights at the end of the vesting period per category		
Category	Series	Max no. of employees	Actual no. of employees	Max. per employee	Max. total	Actual total	Per employee	Total	Vesting period
CEO (1 person)	1	1	1	11,600	11,600	11,600	104,400	104,400	2407-2708
Executve management (maximum 3 persons)	1	3	0	5,375	16,125	-	26,875	-	2407-2708
R&D-management (maximum 7 persons)	1	7	5	3,325	23,275	12,465	16,625	62,325	2407-2708
Employees level 2 (maximum 2 persons)	1	2	–	1,775	3,550	-	8,875	-	2407-2708
Employees level 1 (maximum 8 persons)	1	8	1	1,025	8,200	1,025	5,125	5,125	2407-2708
Total series 1		**21**	**7**		**62,750**	**25,090**		**171,850**	
Employees level 2 (maximum 2 persons)	2	2	1	1,775	3,550	1,775	8,875	8,875	2412-2712
Total series 2		**2**	**1**		**3,550**	**1,775**		**8,875**	
TOTAL series 1 and 2		**23**	**8**		**66,300**	**26,865**		**180,725**	

At an extraordinary general meeting on June 3, 2024, two new long-term incentive programs were adopted (one employee stock option program, PO 2024/2027 series 1, and a performance share program). New articles of associations were also adopted at the extraordinary general meeting, pursuant to which the company may issue class C shares in order to secure delivery of shares to the participants in the programs and to secure payment of future social security contributions. In December 2024, 854,430 C shares were issued, which the company repurchased and holds in its own custody.
The performance share program for employees and the employee stock option program for the CEO and a scientific advisor have been granted and started to be expensed in quarter three, 2024, see adjacent tables.

The performance share program runs for just over three years and participants must retain their employment and invested shares throughout the vesting period in order to receive an allocation of new shares. The number of shares allocated depends on the share price performance and the employment status at the end of the vesting period. As regards the development of the share price, a comparison is made at the end of the vesting period between the initial share price, i.e. the IPO price of SEK 42 per share, and the price at the end of the vesting period. A range between 20% and 60% in share price development results in linearly different allocations of shares. However, a maximum of 360,150 shares can be allocated to participants in the program.

The performance share program generates personnel costs, in accordance with IFRS2 and is initially estimated at approximately SEK 6.8 million and social costs estimated at SEK 6.2 million according to certain assumptions, see also the company's listing prospectus from June 2024.

The new employee stock option program, PO 2024/2027 series 1, also generates personnel costs in accordance with IFRS2 of approximately SEK 1.5 million and social costs of SEK 1.4 million. The dilution for all incentive programs in the company, at maximum allocation, including C shares, is 1.9%.

Note 6 Related party transactions

Transactions with related parties take place on market terms. The table below shows purchases in the Group's parent company and subsidiaries. For further information about transactions with related parties, see annual report 2025.

(TSEK)	Quarter 1		Year
Supplier / Related to	**2026**	**2025**	**2025**
PetoMaj Invest AB 6) Peter Unge, Board member	–	288	1,066
PCW Consultants AB Peter Wallich, Chief Commercial Officer	122	95	583
WBC Europe GmbH 1) Jesper Wiklund, Corporate & business development director	1,216	768	3,456
Arexela AB, 2) Margit Mahlapuu, Executive R&D director	512	475	1,742
Felicia Ahlberg 3) Project manager event	6	13	32
Two Tribes AB 4) Patrik Norgren, CFO	723	–	294
Invegio AB 5) Carina Palm Sundqvist, People & Culture	134	–	399

[1] Related party from quarter 3, 2024

[2] Related party from quarter 4, 2024

[3] Employee since September 2024. Related party to Christer Ahlberg, CEO,

[4] Related party to March 2026

[5] Related party from quarter 4, 2025

[6] Related party to January 2026

Note 7 Number of shares and share capital

Date	Transaction	Change no. of ordinary shares	Total no. of ordinary shares	Total no. of C-shares	Change share capital	Total share capital	Nominal value
2024–01–01	Opening balance 2024	-	26,227,040	-	-	509,153	0.019
2024–06–19	New share issue ordinary shares	17,023,810	43,250,850	-	330,488	839,641	0.019
2024–06–19	Conversion of bridge loan	3,286,939	46,537,789	-	63,810	903,451	0.019
2024–12–03	New share issue C-shares	-	-	854,430	16,587	920,039	0.019
2026–03–31	Closing balance	-	46,537,789	854,430	-	920,039	0.019

* C shares give the right to 1/10 vote.

Note 8 Convertible loan

Convertible loan

During Q1, 2026, the company entered into a financing agreement and under the financing agreement issued a convertible loan with associated standalone warrants. The loan is EUR 8 million and can be converted at any time during the term. If the loan is not converted, it falls due for payment by amortization in 2029. The contractual coupon rate of 12% shall be paid on an ongoing basis. At the same time, 636,341 warrants were issued.

The loan is accounted for in accordance with its substance, divided into a debt and a conversion option. In addition, the warrants are reported as an integral part of the loan in that the market value of the warrants at the time of issue is considered to be part of the interest rate. The effective interest rate on the loan is therefore 37% compared to the coupon rate of 12% and is presented in net financial items.

The loan's conversion option and the warrants are both classified as derivatives, issued call options, and are measured at fair value via the income statement with changes in value presented in net financial items.

The valuation is attributed to level 2 in IFRS 13 and is made in Black Scholes with rolling 12-month historical volatility, risk-free interest on each remaining maturity and an exercise price of EUR 1.59 for the conversion option and EUR 1.67 for the warrant. For the conversion options, an increase in the share price of 10% would lead to a valuation loss of SEK 4.5 million and an increase in volatility of ten percentage points would lead to a valuation loss of SEK 4.2 million. For the warrants, an effect of SEK 0.9 million and SEK 0.7 million respectively would occur in the same scenario.

The convertible loan can be converted into shares or repaid to the counterparty. Any repayment will not be made until 2029. When classified in the balance sheet, the loan needs to be seen together with the option to convert, and since adjustment (conversion) can take place at any time during the term, convertible loans as well as derivatives are classified as current liabilities. As of March 31, 2026, the convertible loan amounts to SEK 45,788 thousand and derivatives regarding conversion options and warrants respectively to SEK 32,945 thousand.

Cinclus Pharma Holding AB has provided collateral for the convertible loan in the form of a primary pledge in all material assets of the Company and its subsidiaries as well as a corporate mortgage of EUR 8 million. There are no financial covenants.

Definitions of key figures and definitions and reconciliation of alternative performance measures

In the report, the company presents key figures in accordance with the IFRS regulations. The company also presents socalled alternative key figures, i.e. measures that are not defined according to IFRS. The alternative key figures found in the report are, among other things, costs related to researchand development as a percentage of total operating costs, equity ratio % and quick ratio %. The company considers the former to be an important complement because it enables a better evaluation of the company's financial trends. This financial performance measure should not be viewed in isolation or considered to replace the performance indicators that have been prepared in accordance with IFRS.

Furthermore, the alternative perform ance measure the company has defined should not be compared with other performance measures with a similar name used by other companies. This because the above-mentioned performancemeasure is not always defined similarly and other companies may calculate it differently than Cinclus Pharma,see adjacent table for further definitions and reconciliation of KPIs and alternative KPIs.

Alternative performance measures

	Quarter 1		Year
Reconciliation of alternative performance measures	**2026**	**2025**	**2025**
Administrative expenses, TSEK	−9,432	−7,813	−57,248
Research and development expenses, TSEK	−91,273	−38,436	−198,402
Operating expenses, TSEK	**−100,705**	**−46,249**	**−255,650**
Research and development expenses / Operating expenses, %	**91%**	**83%**	**84%**
Cash flow for the period, TSEK	−12,712	−42,133	−77,836
Average number of ordinary shares	46,537,789	46,537,789	46,537,789
Cash flow for the period per ordinary share, SEK	**−0.27**	**−0.91**	**−1.67**

	Mar 31, 2026	Mar 31,2025	Dec 31, 2025
Equity, TSEK	281,052	511,625	369,391
Total assets, TSEK	511,639	555,434	546,556
Equity ratio %	**55%**	**92%**	**68%**
Trade receivables, TSEK	1,418	–	16,062
Current tax debt, TSEK	4	–	–
Other receivables, TSEK	4,054	1,814	4,992
Prepaid expenses and accrued income, TSEK	21,463	28,962	28,546
Cash and cash equivalents, TSEK	475,849	523,899	487,254
Total current receivables, TSEK	**502,788**	**554,675**	**536,854**
Current interest bearing liabilities, TSEK	45,788	–	–
Derivative convertible and warrants, TSEK	32,945	–	–
Trade payables, TSEK	29,522	14,389	48,464
Leasing liabilities, TSEK	3,174	222	3,111
Current tax liabilities, TSEK	1	6,829	207
Other current liabilities, TSEK	3,615	3,736	9,656
Contract liabilities, TSEK	35,708	–	54,527
Accrued expenses and deferred income, TSEK	23,205	18,473	20,827
Total current liabilites, TSEK	**173,958**	**43,648**	**136,792**
Quick ratio %	**289%**	**1271%**	**392%**
Equity, TSEK	281,052	511,625	369,391
Number of ordinary shares at the end of the period	46,537,789	46,537,789	46,537,789
Equity per ordinary share, SEK	**6.04**	**10.99**	**7.94**

Definitions of key figures and alternative key figures

Key figures according to IFRS	Definitions
Earnings per share for the period before and after dilution	Profit for the period divided by the average number of shares during the period before and after dilution. Earnings per share after dilution is calculated by adjusting the weighted average number of ordinary shares outstanding for an estimated conversion of all potential ordinary shares giving rise to a dilutive effect, which is in accordance with IAS 33 Earnings per share.

Alternative key figures	Definitions	Reasons for using the key figures
Operating profit / loss (EBIT)	Profit before financial items and tax. The information is taken from the Statement of income.	The key figure helps the reader understand the profitability of the operating business.
Operating expenses	The sum of research and development expenses and administration expenses for the period. The information is taken from the Statement of income.	The key figure helps the reader understand the costs of the operational business.
Research and development expenses / Operating expenses %	Research and development expenses, divided by operating expenses, which consists of research and development expenses and administrative expenses.	The key figure helps the reader understand the proportion of costs attributable to the group's core operations, research and development.
Equity ratio, % *	The equity ratio at the end of each period is calculated by dividing total equity attributable to the parent company's shareholders by total assets.	The equity ratio measures the proportion of the total assets that is financed by the shareholders.
Quick ration, % *	Current assets in relation to current liabilities.	The key figure shows the group's short-term ability to pay.
Number of shares on the balance sheet date	Number of shares in the company at the end of the period.	The key figure gives the reader an understanding of the number of shares at the end of the period.
Equity per share	Equity divided by number of shares at the end of the period	The key figure gives the reader a possibility to compare book value with market value
Cash flow for the period per share	Cash flow for the period divided by average number of shares	The key figure shows the net cash generated or used on a per-share basis

* Reconciliation of these key figures can be found on the previous page.

Certification by the board of directors and the CEO

The board of directors certifies that this interim report gives a true and fair view of the group's operations, financial position and results. For a description of the risks faced by the Cinclus Pharma Group, which are deemed to be unchanged, please refer to the Group's latest annual report.

Stockholm May 13 2026

WENCHE ROLFSEN
Board member

KJELL ANDERSSON
Board member

TORBJÖRN KOIVISTO
Board member

ANDERS ÖHBERG
Board member

HELENA LEVANDER
Board member

NINA RAWAL
Board member

LENNART HANSSON
Chairman of the Board

CHRISTER AHLBERG
CEO and President

Cinclus Pharma

Cinclus Pharma Holding AB (publ

Kungsbron 1

111 22 Stockholm, SVERIGE

Org. nr.559136–8765

Phone: +46 8 13 33 10

info@cinclus pharma.com

cinclus pharma.com