2025

Year-End report

1 January – 31 December 2025

elicera
THERAPEUTICS

Elicera Therapeutics AB. Org.nr 556966-4955

Elicera Therapeutics AB (publ) Year-End report

1 January – 31 December 2025

Fourth quarter (October-December 2025)

- Operating profit/loss amounted to SEK -7,129,626 (-2,911,958).
- Loss for the period amounted to SEK -6,965,286 (-2,603,242).
- Cash flow from operating activities totaled SEK -10,076,983 (-4,232,090).
- Earnings per share before and after dilution totaled SEK -0.14 (-0.07).
- Proposed dividend of SEK 0.00 per share (0.00 for the preceding year).

Period (January-December 2025)

- Operating profit/loss amounted to SEK -17,944,682 (-16,884,056).
- Loss for the period amounted to SEK -17,406,665 (-16,110,327).
- Cash flow from operating activities totaled SEK -21,551,216 (-23,463,165).
- Earnings per share before and after dilution totaled SEK -0.38 (-0.51).

Key events during the fourth quarter

- Elicera decide to change Certified Adviser December 1, 2025 and Liquidity provider on January 22, 2026 to DNB Carnegie.

Key events during the period

- Elicera Continues Phase I/IIa CARMA Study with CAR T-Cell Therapy as Planned Following Safety Committee's Assessment of safety data in Cohort 2.
- Elicera reports: Active lymphoma eliminated in four out of six patients in the first two cohorts with the lowest doses in the CARMA study with iTANK-armed CAR T-cell therapy.
- Elicera clarifies that the ongoing patent application for the company's iTANK platform in the USA is still under review by the United States Patent and Trademark Office (USPTO).
- Elicera's drug candidate ELC-100 receives Orphan Drug Designation in the U.S. for the treatment of pancreatic neuroendocrine tumors.
- In March 2025 subscription of new shares with TO2 were exercised at high 96.3 %. A directed issue was made to guarantors at 3.7 %. Elicera receives 22.0 MSEK before costs.
- Elicera continues the Phase I/IIa CARMA study with its CAR T-cell therapy as planned following the safety committee's assessment of cohort 1.
- Elicera enters a Material Transfer Agreement with University Hospital Tübingen for testing of the company's oncolytic virus candidates, ELC-100 and ELC-201.
- Elicera postpones final reporting of ELC-100 study due to database transition.

Key events after the end of the period

- Elicera announces final data from its Phase I/IIa trial demonstrating a favorable safety profile and promising efficacy signals of oncolytic virus ELC-100 in neuroendocrine tumors
- No other events that impact earnings or the financial position occurred after the end of the period.

Contents

Cell and gene therapies for immune-based cancer treatments

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Condensed earnings and cash flow plus key performance indicators

(AMOUNTS IN SEK UNLESS OTHERWISE INDICATED)	2025 3 MOS OCT-DEC	2024 3 MOS OCT-DEC	2025 9 MOS JAN-JUN	2024 12 MOS JAN-DEC
Other operating income	29,997	2,870,916	10,855,180	7,128,288
Operating expenses	-7,159,623	-5,782,874	-28,799,862	-24,012,344
Operating result	-7,129,626	-2,911,958	-17,944,682	-16,884,056
Result for the period after net financial items	-6,965,286	-2,603,242	-17,406,665	-16,110,327
Cash flow from operating activities	-10,076,983	-4,232,090	-21,551,216	-23,463,165
KEY PERFORMANCE INDICATORS				
Working capital	23,361,361	20,769,437	23,361,361	20,769,437
Quick asset ratio, %	675	406	675	406
Equity/asset ratio, %	85	76	85	76
Earnings per share before dilution	-0.14	-0.07	-0.38	-0.51
Earnings per share after dilution	-0.14	-0.07	-0.38	-0.51
Average number of shares	48,535,544	35,093,286	45,584,106	31,569,593
Average number of warrants	-	11,908,764	2,610,140	9,168,117
Average no. of shares after dilution	48,535,544	47,002,050	48,194,246	40,737,710

Definitions of key performance indicators

Working capital
Sum total of current assets (including cash in hand) minus current liabilities.

Quick asset ratio
Sum total of current assets (including cash in hand) as a percentage of current liabilities.

Equity/asset ratio
Equity in relation to the balance sheet total.

Earnings per share before dilution
Earnings after tax divided by the average number of shares.

Average number of shares
The number of shares, on average, counted from the registration date of the issuance.

Average number of shares after dilution
The number of shares, on average, counted from the registration date.

CEO's Statement

A Year Marked by Clinical Progress

CEO and co-founder, Jamal El-Mosleh

The past year has primarily been characterized by advances in our clinical studies, particularly in the CARMA study. The most recent data report from the CARMA study, presented at the inauguration of the Karolinska ATMP Center in Flemingsberg on August 25, 2025, showed very promising preliminary results. Of the six patients treated with the two lowest dose levels, four exhibited a complete metabolic response – meaning no active cancer could be detected on PET/CT scanning. Among these is a patient who had previously stopped responding to CD19-targeted CAR T-cell therapy, which supports that the iTANK arming gives ELC-301 its unique potential, especially for this hard-to-treat patient group with metastatic or treatment-resistant B-cell lymphoma. The study has so far demonstrated a good safety profile with no reported serious adverse events. Two of the six planned patients have been treated in the third and highest dose group to date. Next data update, including from the third dose group, is planned to be presented during the second quarter of 2026.

> "The study has so far demonstrated a good safety profile with no reported serious adverse events"

Chief Scientific Officer Invited to Leading Scientific Conferences in Europe

Our Chief Scientific Officer, Professor Magnus Essand, has during the year been invited to speak at several prestigious scientific conferences across Europe. This is a clear recognition that both our research work and the results generated are of high international standard and are generating interest in the global immunotherapy field.

Business Development and Strategic Collaborations

During the past year, with the preliminary CARMA data from six patients available, we have initiated targeted updates to potential licensees for ELC-301 and/or the iTANK platform. The patient dataset is still relatively limited, and we are aware that additional data – particularly from the highest dose group – will likely be required to realize a licensing agreement. However, the signals so far have been positive.

In parallel, three academic collaborations are ongoing around the iTANK platform and our oncolytic virus programs. Elicera provides the platform and drug candidates for preclinical studies, while the academic partners conduct preclinical studies in various areas of application. This type of collaborative project is an efficient way for Elicera to validate the broad potential of iTANK, identify new application areas, and lay the groundwork for future drug candidates and possibly new patents.

Final Reporting of the Phase I/IIa Study with ELC-100 Shows Good Safety and Early Efficacy Signals

In the recently completed clinical Phase I/IIa study with ELC-100 (oncolytic virus) in 12 patients with advanced, metastatic neuroendocrine tumors, a favorable safety

profile was observed with no dose-limiting toxicity. The side effects were manageable and in line with the virus's mechanism of action. Among the eight patients evaluable for efficacy signals, partial response was noted in two, providing meaningful early evidence of anti-tumor activity in a disease with a significant unmet medical need. We are grateful to the Victory NET Foundation for their generous support, which enabled the study's completion. With these positive signals, we are now evaluating various strategic options for the continued development of the ELC-100 program.

Looking Ahead: Preclinical Programs and Financing

We are actively advancing efforts to secure financing for our preclinical programs, with particular focus on ELC-401 for the treatment of glioblastoma – one of the most aggressive brain tumor forms with a very short expected median survival. ELC-401, which also builds on our iTANK platform (like ELC-301), has shown promising preclinical results by effectively activating the immune system against this difficult cancer form. The program is now in late preclinical phase, and we are actively planning for clinical studies with a possible start in 2027. During the year, we have conducted extensive work searching for various grants, and we are well aware and prepared that, similar to the grant we previously received from the European Innovation Council (EIC) Accelerator Fund for the CARMA study (ELC-301), it may require repeated applications to obtain funding.

In parallel with the financing efforts, we intend to hold an advisory meeting with the Swedish Medical Products Agency in the spring of 2026 to discuss the study's design and planning.

Regarding the ELC-201 program (our next-generation oncolytic virus candidate), it is also in the preclinical phase with strong preclinical proof-of-concept data, but without as advanced clinical planning as for ELC-401 at present. We continue to evaluate opportunities for financing and strategic partnerships to advance these programs and address the large unmet medical need in patients with hard-to-treat solid tumors.

Summary and Thanks

2025 has been a year of significant progress for Elicera: strong early clinical data from CARMA, successful completion of the ELC-100 study, increased international visibility, and forward-looking collaborations. We are entering 2026 continuing to focus on maintaining our momentum and look forward to an eventful year where we position ourselves as a leading player in cell and gene therapies for difficult cancer forms.

A warm thank you to our dedicated team, the board, our academic and industrial collaboration partners, and to you shareholders for your continued trust and support. Together, we work toward the goal of developing new, effective treatment options for cancer patients with significant needs.

Jamal El-Mosleh
CEO Elicera Therapeutics

Introduction to Elicera Therapeutics

Elicera Therapeutics AB is a clinical stage cell and gene therapy company developing the next generation of armed cancer treatments. The company has developed a portfolio consisting of the patented iTANK gene technology platform and four drug candidates in clinical and preclinical development phase.

iTANK permits strengthening of the efficacy of CAR T-cell therapies and oncolytic viruses – what we call "arming" them – against aggressive and recurrent solid cancers. In preclinical studies, this method has demonstrated potent efficacy against solid tumors, which are known for being extremely difficult to treat with current approved CAR T-cell therapies. The method is being applied in three of the company's drug candidates under development (ELC-301, ELC-401 and ELC-201) and the technology is being offered on a license basis to other pharmaceutical companies that are active in the field of CAR T-cell therapies. This platform thus opens the door to new possibilities for treating solid tumors where current CAR T-cell therapies have not yet been successful.

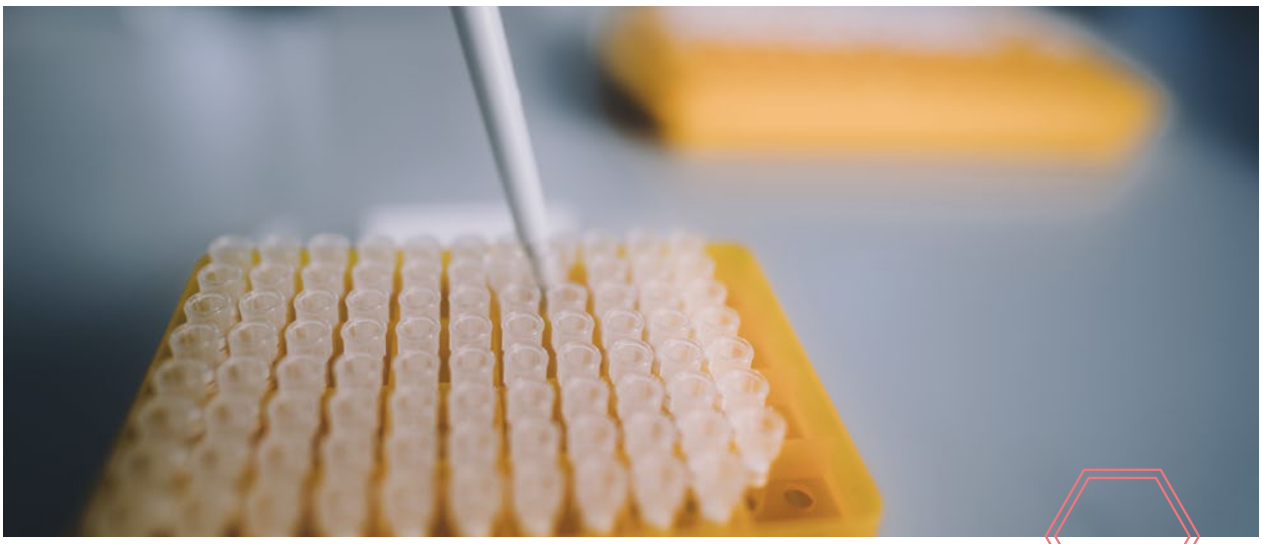
Elicera's drug candidates comprise two CAR T-cell therapies, ELC-301 and ELC-401, and two oncolytic viruses, ELC-201 and ELC-100. ELC-100 recently completed a clinical Phase I/II trial showing a good safety profile and promising signs of clinical efficacy, while ELC-301 initiated a clinical phase I/IIa study (CARMA) in November 2024 showing 4 out 6 patients in the first two dose groups with complete metabolic responses.. ELC-201 and ELC-401 are in the preclinical development phase.

Elicera's operations and product portfolio are based on years of research conducted by Professor Magnus Essand, who has a sterling reputation in the field at Uppsala University. Elicera's strengths are based on a profound understanding of how cells and viruses can be genetically modified to trigger a robust immune response to cancer.

CAR T-cell therapies in brief

CAR T-cells are a form of cell therapy that are produced by using gene modification to place a synthetic receptor (chimeric antigen receptor, or CAR) in the patient's T-cells. This receptor has been customized for a high degree of affinity against a specific tumor antigen – a molecule that is visible on the surface of the cancer cell – and helps the T-cell locate, bind to and kill the cancer cell.

CAR T-cell therapies have made it possible to cure forms of cancer that were previously incurable, but the seven treatments that have been approved to date are only effective against various forms of hematological cancers, meaning ones found in the blood, lymph system or bone marrow. Despite the major advances that have occurred in this field of treatment, around 50 percent of the patients who suffer from these hematological cancer forms still succumb to these diseases.



Oncolytic viruses in brief

Oncolytic viruses are genetically modified viruses that are designed to selectively infect and destroy cancer cells without harming normal cells. When the tumor cell "bursts" and dies through this process, known as oncolysis, an immune response against tumor cells is initiated through tumor neoantigens (mutated antigens, the antigens that provoke the strongest immune response) being released and captured by the patient's dendritic cells, which then teach the T-cells to attack cancer cells wherever they are found in the body.

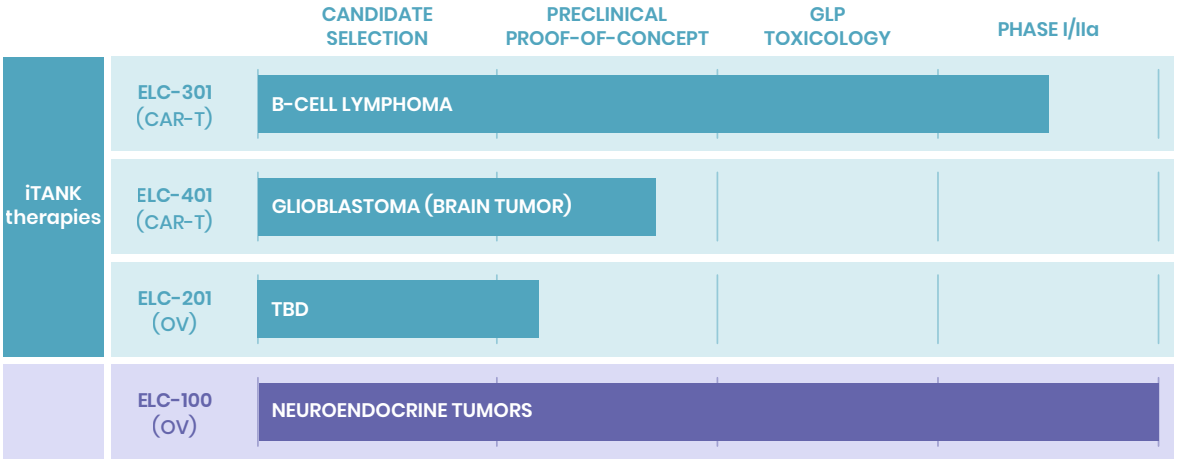
Business model and strategy

Elicera's business model is to develop and, over the long term, outlicense its in-house and patented arming technology iTANK and treatment methods for cancers. The iTANK platform is ready for commercialization via non-exclusive licenses to various CAR T-cell therapy developers, while Elicera's four internal development programs in immunotherapy are intended to be licensed exclusively at various stages of development. All outlicensing is expected to generate significant revenue in the form of technology upfront payments, milestone payments and royalties. The strategy for generating revenue from commercial partnerships is built on:

- Conducting preclinical and clinical trials that demonstrate the mechanism of action and efficacy of the programs.
- Benefiting from the company's competence in cell and tumor immunology in order to develop drugs that address major medical needs that are not being met.
- Continuing to build on its strong patent portfolio and accumulate valuable know-how.

Product portfolio

The company's product portfolio consists of the iTANK platform technology and four drug candidates – two are in the field of oncolytic viruses (ELC-100 and ELC-201) and two in the field of CAR T-cell therapies (ELC-301 and ELC-401).



PoC: Proof-of-Concept GLP: Good Laboratory Practice

Figure 1: Elicera's product portfolio.

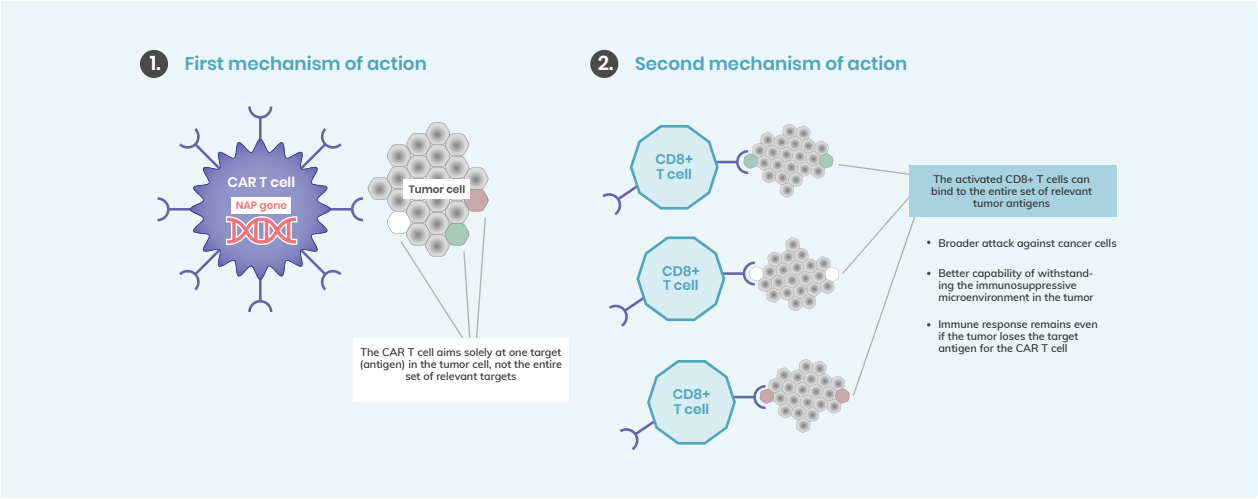


Figure 2: The iTANK platform results in a second parallel mechanism of action and a broad attack on tumor cells via CD8+ T-cells. The CD8+ T-cells are activated against the entire set of relevant targets on the tumor cell.

Product portfolio

iTANK

Elicera has developed iTANK, a patented and commercially available platform technology for expanding the areas of application for CAR T-cell therapy. iTANK makes it possible to impact the microenvironment in solid tumors, activate a robust immune response against cancer and develop a long-term immunological memory related to several different tumor targets, which aims to counteract recurrences of cancer.

The technology arms CAR T-cells with the bacteria protein NAP (neutrophil-activating protein from Helicobacter pylori). When the CAR T-cells are introduced into the body, NAP is set free around the cancer cells, which initiates an inflammatory process that involves the body's immune system signaling other immune cells to accumulate in the tumor. The process leads to the immune cells being triggered to kill those cancer cells that the CAR T-cells are incapable of attacking. The aim is to create an immunological memory via the lymphatic system in pace with the destruction of the tumor, to drastically reduce the risk of relapse.

The capacity among CAR T-cells armed with iTANK to activate the body's immune system in a non-specific manner (as opposed to the specificity directed via the CAR) against several unique tumor targets yields completely new possibilities for developing better CAR T-cell treatments against blood cancers and new treatments against solid cancers.

Preclinical studies with iTANK were able to confirm that CAR T-cells armed with NAP generate robust immunological activity in the tumor tissue by attracting other immune cells. This is believed to be able to meet the challenge with a hostile tumor microenvironment.

All together, the results from the preclinical studies support the possibilities of using Elicera's unique method to create CAR T-cell therapies against a range of solid forms of cancer – something that at present is very difficult.

The results from the preclinical studies were published in 2022 in the high-impact scientific journal Nature Biomedical Engineering[1], and constitutes a fundamental pillar for the validity of the scientific concept.

Figure 2 above illustrates the advantages of the iTANK platform and shows how CAR T-cells armed with NAP generate another mechanism of action through killer T-cells that focus broadly on the entire set of relevant tumor antigens in cancer cells – not only a single target, as often is the case for conventional CAR T-cells.

1 https://www.nature.com/articles/s41551-022-00875-5

Elicera's four drug candidates

ELC-301: B-cell lymphoma

The ELC-301 program is being developed to treat B-cell lymphoma. Diffuse large B-cell lymphoma (DLBCL), the most common non-Hodgkin lymphoma, is an aggressive form of cancer that starts out from the immune system's B-cells. DLBCL is one of the most common forms of B-cell cancer and the disease progresses rapidly, which requires treatment to be administered as soon as possible after a diagnosis has been established.

The specific target group that ELC-301 is being developed for is patients who are suffering from a particularly difficult form of DLBCL or who have relapsed after several rounds of standard treatment. The current standard treatment comprises a combination of chemotherapy and antibodies, and 60 to 70 percent of patients can be cured this way. Among the patients who suffer a relapse, CAR T-cell therapy comprises the next step in the treatment hierarchy. Despite the initial disappearance of the disease among many after CAR T-cell treatment, the frequency of recurrence in the patients remains high – between 40 and 50% – and the treatment alternatives, in the form of more advanced therapies following current CAR T-cell therapy, are limited.[2]

All of the currently approved CAR T-cell therapies in B-cell lymphoma target the tumor antigen CD19 – a common B-cell protein that is overproduced on the surface of cancer cells in DLBCL. Among many of the individuals who suffer relapses, this tumor antigen disappears and further treatments with the same CAR T-cell therapy therefore become ineffectual. ELC-301 targets CD20 instead, which is also overrepresented in B-cell lymphoma. By switching the target protein to CD20 and arming the CAR T-cells with the iTANK platform, ELC-301 facilitates treatment of relapse patients who are in need of a new efficacious alternative.

In November 2024, Elicera started a clinical phase I/IIa trial, called the CARMA-study (NCT06002659), with ELC-301 in patients with severe or recurring DLBCL. CARMA is conducted in two parts: a dose-escalation phase (phase I) and a dose-expansion phase (phase IIa). The initial part is planned to include three cohorts (dosing groups) with three patients in the first and second dosing groups and six patients in the third dosing group, who are expected to receive the maximum dose. The objective is to study safety and to identify the optimal dose for treatment with the CAR T-cell therapy ELC-301, which will then be tested in an additional six patients in the phase IIa part of the study.Elicera's Chief Scientific Officer, Professor Magnus Essand, presented preliminary efficacy data from the first two cohorts with the lowest dose level at the inauguration of the Karolinska ATMP Center in Flemingsberg, Sweden, on August 25. The results showed that 4 out of 6 treated patients exhibited complete metabolic response, meaning no active disease was detected. Following the safety committee's positive assessment of safety data in cohort 2, recruitment is now proceeding for patients in the third and final cohort with the maximum planned dose. The CARMA-study is being financed in part with EUR 2.5 million in grants from the EIC Accelerator Fund. The agreement between Elicera and Uppsala University regulates the partnership and Elicera's ownership rights to the data.

2 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC9561408/

ELC-401: Glioblastoma

The ELC-401 program is being developed to treat glioblastoma (GBM), a solid tumor. Glioblastoma is an aggressive form of brain cancer with an extremely high mortality rate, and the expected median survival rate among persons with the diagnosis is approximately 15 months.

At present, glioblastoma is treated primarily with surgery and radiation therapy since it is a challenge to develop drugs that can pass through the blood-brain barrier and be efficacious in the central nervous system. Elicera's drug candidate ELC-401 targets the IL13Ra2 tumor antigen, which is a receptor protein that is overrepresented in GBM. In a preclinical study, the company was able to demonstrate that IL13Ra2 is an effective tumor target for CAR T-cells strengthened with iTANK. Owing to iTANK, ELC-401 is expected to also be able to counteract the robust immunosuppressive micro-environment in glioblastoma and mobilize an immune response against other targets in this heterogeneous form of cancer as well.

A study published in Nature Communications in 2023[3] evaluated the synthetic receptor that forms the basis of ELC-401. The results included the finding that the CAR T-cell had a potent cell-killing efficacy and prolonged survival in the disease model. ELC-401 is currently in a preclinical evaluation phase.

ELC-201: Solid tumors

Alongside its CAR T-cell program and ELC-100, Elicera is developing ELC-201, a program to develop oncolytic virus treatment with the potential to treat several different forms of solid cancer.

It is expected that ELC-201 will form a double attack on cancer tumors, both through the oncolytic virus and via a parallel T-cell response against cancer owing to the reinforcement with iTANK and an additional T-cell stimulating factor.

The company has extensively surveyed potential cancer indications for ELC-201 based on both scientific and commercial considerations, and is now evaluating alternatives for financing the program of clinical trials, with a focus on commercial partnership and various types of soft financing.

3 https://www.nature.com/articles/s41467-023-40303-z

Phase I/II trial on neuroendocrine tumors

Dose escalation in 12 patients – completed

In partnership with Uppsala University, which is acting as sponsor for the study

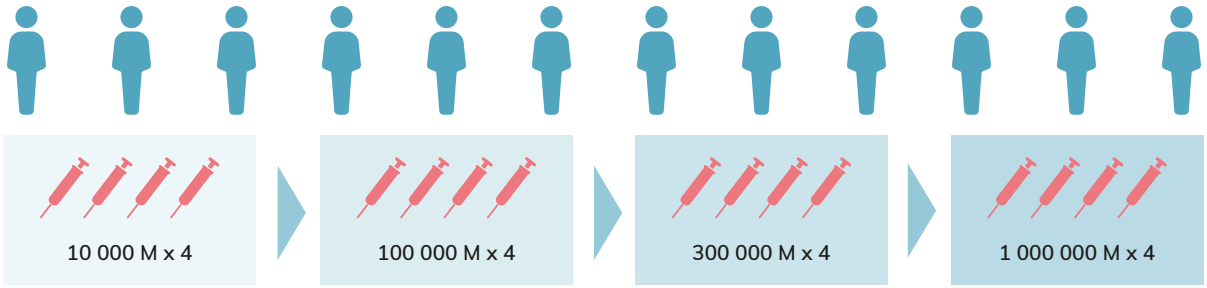


Figure 3: Ongoing Phase I/II trial on neuroendocrine tumors has recently been completed showing a good safety profile and promising signs of clinical efficacy.

ELC-100 (AdVince): Neuroendocrine tumors

ELC-100, also known as AdVince, is a program for developing and treating neuroendocrine tumors (NETs), which arise from cells in the neuroendocrine system. The tumors can be found anywhere in the body, but occur primarily in the stomach and intestines (43%) as well as in the lungs (30%) and in the pancreas (7%)[4].

In preclinical studies on mice, ELC-100 demonstrated extended survival compared with different types of standard treatments such as tyrosine kinase inhibitors and radioactive medicines.

ELC-100 is based on a genetically modified adenovirus, Ad5PTD, and has been optimized with regard to its ability to enter specifically neuroendocrine cancer cells and not healthy cells, where they propagate until the tumor cell bursts and dies in a process known as oncolysis.

In addition to the selective propagation in NET cells, ELC-100 has also been specifically modified to prevent propagation in liver cells in order to reduce the risk of damage to liver cells since the oncolytic virus is administered via the hepatic artery.

ELC-100 recently completed a clinical Phase I/II trial (ClinicalTrials.gov identifier: NCT02749331) with Uppsala University as sponsor (agreements between Elicera and Uppsala University regulate the partnership and Elicera's ownership rights to the data). The main purpose of the trial is to study the safety of the treatment and determine the maximum tolerated dose. In early January 2026, Elicera reported that ELC-100, was generally well-tolerated with no dose-limiting toxicities observed. Importantly, the trial also revealed promising efficacy signals, including partial tumor responses in two out of eight patients evaluable for efficacy, providing early evidence of anti-tumor activity in this highly treatment-resistant population.

In January, ELC-100 was granted Orphan Drug Designation by the U.S. Food and Drug Administration (FDA) for the treatment of pancreatic neuroendocrine tumors. Orphan Drug Designation (ODD) is intended to promote the development of drugs that address rare diseases. In the United States, the Food and Drug Administration (FDA) grants this status to drugs or biological products designed to treat diseases affecting fewer than 200,000 people in the country. During the development of the drug candidate, this designation provides certain advantages, such as tax credits for clinical trials conducted in the U.S. At a later stage, ODD offers the opportunity to waive fees associated with applications for marketing approval, as well as up to seven years of market exclusivity.

4 https://www.cancer.net/cancer-types/neuroendocrine-tumors/introduction

Financial information

Financial performance during the fourth quarter, October 1 - December 31, 2025

Operating loss

Operating result for the quarter totaled SEK -7,129,626 (SEK -2,911,958), which is a change of SEK -4,217,668 compared to the year-earlier period. The change is due primarily to a decrease grants booked SEK -2,840,919 and SEK -1,376,749 increase in costs.

Loss for the quarter

Result for the quarter amounted to SEK -6,965,286 (–2,603,242). Earnings per share totaled SEK -0.14 (-0.07).

Liquidity and cash flow

- Cash flow from operating activities totaled SEK -10,076,983 (-4,232,090).
- Cash flow from investing activities totaled SEK 0 (0) SEK.
- Cash flow from financing activities totaled SEK 0 (0).
- Cash flow for the quarter amounted to SEK -10,076,983 (-4,232,090).
- At the end of the period, the company's cash and cash equivalents totaled SEK 24,846,480 (26,399,108).

Financial performance during the period, January 1–December 31, 2025

Operating loss

Operating loss for the period totaled SEK -17,944,682 (-16,884,056), which is a change of SEK -1,060,626 compared to the year-earlier period. The change is due primarily to an increase grants booked SEK +3,726,892 and SEK -4,787,518 increase in costs.

Loss for the period

Loss for the period amounted to SEK -17,406,665 (SEK -16,110,327). Earnings per share totaled SEK -0.38 (-0.51).

Liquidity and cash flow

- Cash flow from operating activities totaled SEK -21,551,216 (-23,463,165).
- Cash flow from investing activities totaled SEK 1 000 (0) SEK.
- Cash flow from financing activities totaled SEK 19,997,589 (20,479,306).
- Cash flow for the period amounted to SEK -1,552,628 (-2,983,859).
- At the end of the period, the company's cash and cash equivalents totaled SEK 24,846,480 (26,399,108).

EU accelerator program

In June 2022 Elicera is selected, in very hard competition, for a grant from EU accelerator program amounting to SEK 2.5 m (about SEK 27 m). EU has paid the first part amounting at SEK 17.7 m previously. In February 2025 SEK 5.6 m was paid. The remaining part approximately SEK 1,5 m is expected to be paid during H1 2026.

The amount is booked as prepaid income. The income has been booked as the costs occur in the project and the prepaid income have been reduced.

During the period SEK 10,8 m has been booked as income.

Investments

Elicera's investments for the period totaled SEK 1 000 (0).

Personnel and organization

The number of employees at the end of the period was 2. Elicera's organization comprises all the competence and experience that is necessary to run the company. Close collaboration has been established with a number of key consultants in patents, preclinical, clinical trials, development of pharmaceuticals, regulatory expertise for manufacture and documentation, quality assurance, finance, and law.

Warrants serie TO2

The subscription of new shares supported by TO2 took place February 26 to March 11, 2025.

For securing the subscription an agreement was signed Mangold. Bottom guarantor signed up for up to 70 % and top guarantors for the remaining 30 %. The agreement secured that the full amount should be subscribed.

In total 96.3 % of warrants were used for new shares. Guarantors received 3.7 %. This is a strong outcome. Through the usage of TO2 for signing new shares and the directed issue the number of shares increase with 11,908,764 from 35,093,268 shares to 47,002,032 shares. The share capital increase with 500,168.09 SEK from 1,473,917.26 to 1,974,085.35.

Top guarantors were compensated with new shares and bottom guarantors could receive cash or shares (higher compensation). A directed issue was made to guarantors of 1,533,512 shares that increase the number of shares to 48,535,544. The share capital increase with 64,407.50 SEK to 2,038,492.85 SEK.

The compensation new issue had no cash impact.

Annual general meeting 2025

The Annual General Meeting was held on May 15, 2025 in Stockholm. The AGM followed the proposals from the nomination committee.

The AGM resolved to re-elect its Board of Directors: Agneta Edberg (chair), Magnus Essand, Christina Herder, Margareth Jorvid and Sharon Longhurst. Di Yu was re-elected deputy. Board fees was fixed SEK 360,000 for Chairman of the Board Agneta Edberg and SEK 150,000 for the other members. Cedra Väst KB, with signatory auditor Kristoffer Håkansson, was re-elected as auditor. The Board of Directors was authorized to conduct a private placement with a maximum dilution of 20 %.

Nomination committee

On the Annual General Meeting confirmed rules to guide the work of the Nomination Committee. The largest owners at September 30, 2025 were Di Yu, Magnus Essand and

Jamal El-Mosleh, who control 20 % of the votes, and have therefore been appointed to the Nomination Committee with Magnus Essand as chair.

Shareholders with viewpoints and proposals are asked to contact the chairman of the Nomination Committee, Magnus Essand, via email at info@elicera.com.

Annual general meeting 2026

The AGM will be held on May 5, 2025 at 1:00 p.m. CEST, at the offices of Advokatfirman Delphi, Mäster Samuelsgatan 17 in Stockholm.

Shareholders will be notified that the meeting has been called through an announcement in Post- och Inrikes Tidningar and on the company's web site, as well as through an announcement in Svenska Dagbladet, at the earliest six weeks and at the latest four weeks prior to the meeting.

Shareholders wishing to have a matter addressed at the AGM can submit a written request to Elicera Therapeutics AB, Attn: Board of Directors, World Trade Center Göteborg, Mässans gata 10, 7th floor, SE-412 51 Gothenburg, Sweden. The request must be received by the Board at the latest seven weeks prior to the AGM, or enough in advance so that the matter, if required, can be included in the notification to attend.

The Annual Report will be published on March 27, 2026.

Risks and uncertainties

In addition to the general uncertainty related to research and development operations and delays in the start of clinical trials, there are no known tendencies, uncertainties, potential receivables or other demands, commitments or events that could be expected to have a material impact on the company's future prospects in addition to the risks presented in the annual report (pages 27-30).

Equity

Equity was impacted by the new share issue and earnings during the period. At the end of the period, equity totaled SEK 23,361,361 (20,770,437).

Proposal for appropriation of profits

The Board of Directors and the CEO propose that no dividend (SEK 0.00 per share, same as the previous year) be paid for the fiscal year January 1 – December 31, 2025.

The share

The Elicera share was listed on Nasdaq First North Growth Market on June 11, 2021. The share register is managed by Euroclear.

Loss after tax divided by the average number of shares for the period totaled SEK -0.38 (-0.51) for the reporting period. At the end of the period Elicera had approximately 5,500 shareholders, an increase with around 3,000 since end of 2024. The number of shares at the end of the period was 48,535,544.

NAME	NUMBER OF SHARES	SHARE OF VOTES/ CAPITAL (%)
Avanza	3,862,262	8.0
Di Yu	3,463,715	7.2
Magnus Essand	3,397,059	7.1
Jamal El-Mosleh	2,712,200	5,6
SC Holding	1,392,000	2.9
Other owners	33,679,778	69.4
Total number of shares	**48,535,544**	**100.0**

Change of certified adviser and liquidity provider to dnb carnegie

Elicera has entered into an agreement with DNB Carnegie Investment Bank AB regarding the roles as Certified Adviser and Liquidity provider. DNB Carnegie will take over as Certified Adviser on December 1, 2025 and Liquidity provider on January 22, 2026.

Transactions with affiliated parties

Board member Magnus Essand is part-time employee as CSO and received a salary of 470,000 SEK (371,610).

Board deputy Di Yu is part-time employee as Head of translational research and received a salary of 378,000 SEK (411,480) and payment to pension plan at 192,566 SEK (103,684).

Advanced Biologics, the company that employs Sharon Longhurst has invoiced 0 SEK (110,797).

The pricing took place under market conditions.

Accounting principles – change to IFRS (RFR2)

Elicera changes from K3 to RFR2 (IFRS for companies without subsidiaries). The change is for the planned move to First North Premier. No adjustments have been identified and as consequence the profit & loss, balance sheet and cash flow are unchanged also for previous periods.

Audit

This year-end report has not been audited.

Events after the end of the period

No key events that impact the financial statements occurred after the end of the period.

Assurance of the Board

The Board of Directors and CEO give their assurance that this Year-end report provides a true and fair overview of the company's operations, financial position, and earnings, and that it describes the material risks and uncertainties faced by the company.

Gothenburg, February 13, 2026.

The Board of Directors of Elicera Therapeutics (publ)

Agneta Edberg, Chairman

Magnus Essand

Christina Herder

Margareth Jorvid

Sharon Longhurst

Jamal El-Mosleh, CEO

Statement of income and other comprehensive

(AMOUNTS IN SEK)	2025 3 MOS OCT-DEC	2024 3 MOS OCT-DEC	2025 12 MOS JAN-DEC	2024 12 MOS JAN-DEC
Other income	29,997	2,870,916	10,855,180	7,128,288
Operating expenses				
Other external expenses	-5,467,796	-4,457,093	-21,999,543	-18,941,803
Personnel expenses	-1,691,827	-1,322,643	-6,800,319	-5,058,765
Depreciation of property, plant and equipment	-	-2,938	-	-11,776
Total operating costs	**-7,159,623**	**-5,782,874**	**-28,799,862**	**-24,012,344**
Operating result	**-7,129,626**	**-2,911,958**	**-17,944,682**	**-16,884,056**
Interest income and similar profit/loss items	169,408	318,086	545,894	826,579
Interest expenses and similar profit/loss items	-5,068	-9,370	-7,877	-52,850
Result before taxes	**-6,965,286**	**-2,603,242**	**-17,406,665**	**-16,110,327**
Tax	-	-	-	-
Result for the period	**-6,965,286**	**-2,603,242**	**-17,406,665**	**-16,110,327**

Balance sheet

(AMOUNTS IN SEK)	DEC 31 2025	DEC 31 2024
ASSETS		
Financial assets		
Securities	-	1,000
Total financial assets	**-**	**1,000**
Total non-current assets	**-**	**1,000**
Other receivables	545,898	881,867
Other interim receivables	2,029,078	284,770
Cash and bank	24,846,480	26,399,108
Total current assets	**27,412,456**	**27,565,745**
TOTAL ASSETS	**27,421,456**	**27,566,745**
EQUITY		
Restricted equity		
Share capital	2,038,493	1,473,917
Total restricted equity	**2,038,493**	**1,473,917**
Non restricted equity		
Share premium reserve	106,055,937	86,622,924
Profit or loss carried forward	-67,326.404	-51,216,077
Loss of the year	-17,406,665	-16,110,327
Total non-restricted equity	**21,322,865**	**19,296,520**
Total equity	**23,361,361**	**20,770,437**
Current liabilities		
Account payables	2,568,602	2.043,872
Other current liabilities	248,961	354,399
Accrued expenses and prepaid income	1,242,532	4,398,037
Total current liabilities	**4,060,095**	**6,796,308**
TOTAL EQUITY AND LIABILITIES	**27,421,456**	**27,566,745**

Statement of changes in equity

(AMOUNTS IN SEK)	SHARE CAPITAL	SHARE PREMIUM RESERVE	RETAINED EARNINGS	LOSS FOR THE YEAR	TOTAL EQUITY
Opening balance at January 1, 2024	**830,844**	**66,786,691**	**-34,818,100**	**-16,397,977**	**16,401,458**
Proposed appropriation of earnings to AGM			-16,397,977	16,397,977	-
New issue	643,073	26,917,209	-	-	27,506,282
Capitalization costs		-7,080,976			-7,080,976
Loss for the period	-	-	-	-13,507,085	-13,507,085
Closing balance at September, 2024	**1,473,917**	**86,622,824**	**-51,216,077**	**-13,507,085**	**23,373,679**

(AMOUNTS IN SEK)	SHARE CAPITAL	SHARE PREMIUM RESERVE	RETAINED EARNINGS	LOSS FOR THE YEAR	TOTAL EQUITY
Opening balance at October 1, 2024	**1,473,917**	**86,622,924**	**-51,216,077**	**-13,507,085**	**23,373,679**
Loss for the period	-	-	-	-2,603,242	-2,603,242
Closing balance at December 31, 2024	**1,473,917**	**86,622,924**	**-51,216,077**	**-16,110,327**	**20,770,437**

(AMOUNTS IN SEK)	SHARE CAPITAL	SHARE PREMIUM RESERVE	RETAINED EARNINGS	LOSS FOR THE YEAR	TOTAL EQUITY
Opening balance at January 1, 2025	**1,473,917**	**86,622,924**	**-51,216,077**	**-16,110,327**	**20,770,437**
Proposed appropriation of earnings to AGM			-16,110,327	16,110,327	-
New issue	500,169	21,531,046	-	-	22,031,215
Capitalization costs		-2,033,626			-2,033,626
Compensation issue	64,407	2,772,590			2,836,997
Compensation costs		-2,836,997			-2,836,997
Loss for the period	-	-	-	-10,441,379	-10,441,379
Closing balance at September 30, 2025	**2,038,493**	**106,055,937**	**-67,326,404**	**-10,441,379**	**30,326,646**

(AMOUNTS IN SEK)	SHARE CAPITAL	SHARE PREMIUM RESERVE	RETAINED EARNINGS	LOSS FOR THE YEAR	TOTAL EQUITY
Opening balance at October 1, 2025	**2,038,493**	**106,055,937**	**-67,326,404**	**-10,441,379**	**30,326,646**
Loss for the period	-	-	-	-6,965,286	-6,965,286
Closing balance at December 31, 2025	**2,038,493**	**106,055,937**	**-67,326,404**	**-17,406,665**	**23,361,360**

DISCLOSURE OF SHARES AND WARRANTS	NUMBER OF SHARES
Number of shares at the beginning of the year	35,093,268
Number of shares at 2025-12-31	48,535,544
Number of warrants at the beginning of the year	11,908,764
Number of warrants at 2025-12-31	0

Cash flow statement

(AMOUNTS IN SEK)	2025 3 MOS OCT-DEC	2024 3 MOS OCT-DEC	2025 12 MOS JAN-DEC	2024 12 MOS JAN-DEC
OPERATING ACTIVITIES				
Operating loss before financial items	-7,129,626	-2,911,958	-17,944,682	-16,884,056
Reversal of depreciation	–	2,938	–	11,776
Interest received	169,408	318,086	545,730	826,526
Interest paid	-5,068	-9,370	-7,877	-52,850
Taxes paid	–	–	–	–
Cash flow from operating activities before changes in working capital	**-6,965,286**	**-2,600,304**	**-17,406,665**	**-16,098,551**
Increase/Decrease in short-term receivables	-202,472	-685,846	-1,408,339	-382,361
Increase/Decrease in account payable	526,036	1,502,666	524,730	1,160,857
Increase/Decrease in other current liabilities	-3,435,261	-2,448,606	-3,260,942	-8,143,110
Cash flow from operating activities	**-10,076,983**	**-4,232,090**	**-21,551,216**	**-23,463,165**
Investing activities				
Investments in intangible assets	–	–	1,000	–
Change in non-current financial assets	–	–	–	–
Cash flow from investing activities	**–**	**–**	**1 000**	**–**
Financing activities				
New share issue	–	–	22,031,214	27,560,282
Capital raising costs	–	–	-2,033,626	-7,080,976
Cash flow from financing activities	**–**	**–**	**19,997,588**	**20,479,306**
Cash flow for the period	-10,076,983	-4,232,090	-1,552,628	-2,983,859
Cash and cash equivalents at the beginning of the period	34,923,463	30,631,198	26,399,108	29,382,967
Cash and cash equivalents at the end of the period	**24,846,480**	**26,399,108**	**24,846,480**	**26,399,108**

Financial calendar

Annual report	March 27, 2026
Interim Report January–March 2026	May 5, 2026
Annual meeting	May 5, 2026
Interim Report January–June 2026	August 21, 2026
Interim Report January–September 2026	November 27, 2026
Year-end Report 2026	February 16, 2027

If you have questions, please contact:

Jamal El-Mosleh, CEO
+46 (0) 703 319 051
jamal.elmosleh@elicera.com

Adresses

Elicera Therapeutics AB
World Trade Centre Gothenburg
Mässans gata 10, vån 7, 412 51 Gothenburg
www.elicera.com

Certified Adviser
Certified Adviser is DNB Carnegie Investment Bank AB (publ)
certifiedadviser@dnbcarnegie.se

elicera
THERAPEUTICS

www.elicera.se