

Cessatech A/S Announces Financial Results for the Full Year 2025

Copenhagen, Denmark - 27 February 2026 - Cessatech A/S (“Cessatech” or the “Company”), a late-stage paediatric specialty company developing and commercializing non-invasive hospital medicines for children, today announced financial results for the year ended December 31, 2025, which also includes financial results for the fourth quarter of 2025.

Full year 2025 for the period 1 January - 31 December (Q4'2025 results in brackets):

- Net revenue: KDKK 5,718 (1,989)
- Operating result: KDKK -14,967 (-1,836)
- Net result: KDKK -10,904 (-773)
- Cash at bank end of the period: KDKK 8,591 (8,591)
- Earnings per share*: KDKK -0.60 (-0.04)
- Solidity**: 91% (91%)

**Earnings per share (DKK per share): Operating result divided by the average number of shares during the period. The total number of shares as of 31 December 2025 amounted to 18,576,437 shares, the average number of shares during the full year was 18,090,665 **Solidity: Total equity divided by total capital and liability.*

The Annual Report is presented for approval at the Annual General Meeting, to be held on 28 March 2026. The Board of Directors and the CEO propose that no dividend be paid for the financial year from 1 January 2025 to 31 December 2025.

The Annual Report 2025 is available on Cessatech's website.

About Cessatech A/S

Cessatech is a pivotal-stage paediatric biotech company developing first-in-class specialty hospital medicines for children, focused on high-impact unmet needs in acute and emergency care. The company's lead program, CT001, is advancing toward near-term commercialization. Cessatech operates a capital-efficient, partnership-driven model, combining a lean, experienced team with best-in-class partners for development, manufacturing and commercial execution across key markets. Headquartered in Hellerup, Denmark, Cessatech is led by a seasoned leadership team with a strong track record in drug development and product launches in Europe, the US and Asia.

Highlights during the full year 2025

- Announced positive top-line results from Study 0202, the final required paediatric clinical study for CT001, confirming its safety and efficacy profile and enabling initiation of the European regulatory submission process.
- Received a positive Notified Body opinion under the EU Medical Device Regulation (MDR) for CT001, confirming that the device component meets applicable EU safety and performance requirements.
- Submitted a Paediatric Use Marketing Authorisation (PUMA) application for CT001 to the European Medicines Agency (EMA), which was subsequently validated.
- Advanced U.S. activities through an extended collaboration with STAQ Pharma and completed technology transfer activities supporting near-term hospital supply.
- Strengthened the company's financial position through a directed share issue raising approximately DKK 15 million.
- Enhanced Executive Management with the appointment of Martin Juhl as Chief Scientific Officer (CSO).

For more information about Cessatech, please contact:

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