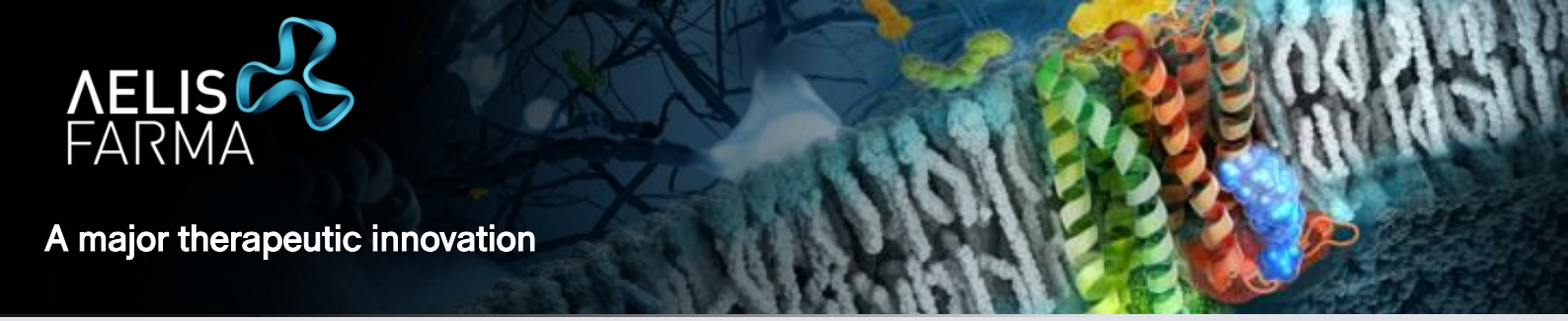

Press release

Availability of the 2025 Universal Registration Document

Bordeaux, April 24, 2026 – 6:00 p.m. CEST – Aelis Farma (ISIN: FR0014007ZB4 – Ticker: AELIS), a clinical-stage biopharmaceutical company specializing in the development of treatments for brain and peripheral diseases involving the CB_1 receptor, today announces the publication of its 2025 Universal Registration Document (URD) as filed with the French Financial Markets Authority (*Autorité des Marchés Financiers* (AMF)) on April 23, 2026, under number D.26-0284.

The Universal Registration Document 2025 includes in particular:

- The 2025 annual financial report;
- The statutory auditor's reports;
- Information on the statutory auditor's fees;
- Information relating to corporate social responsibility.

The URD can be consulted and downloaded on the Company's website under the heading Investors / Documentation, as well as on the AMF website (https://www.amf-france.org).

Financial Agenda:

- Annual General Meeting, June 3, 2026
- 2026 Half-Year Results, September 15, 2026

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About AELIS FARMA

Founded in Bordeaux in 2013, Aelis Farma is a biopharmaceutical company that is developing a new class of drugs, the Signaling Specific inhibitors of the CB_1 receptor of the endocannabinoid system (CB_1-SSi). CB_1-SSi have been developed by Aelis Farma based on the discovery of a natural regulatory mechanism of CB_1 hyperactivity made by the team led by Dr Pier Vincenzo Piazza, the Company's CEO, when he was the director of the Neurocentre Magendie of INSERM in Bordeaux. By mimicking this natural mechanism, CB_1-SSi appear to selectively inhibit the disease-related activity of the CB_1 receptor without disrupting its normal physiological activity. CB_1-SSi have consequently the potential to provide new safe treatments for several brain and peripheral organ diseases.

Aelis Farma currently has two first-in-class clinical-stage drug candidates. AEF0117 for the treatment of cannabis use disorders (CUD), that has shown to be able to decrease cannabis use across two studies. AEF0217 for cognitive disorders, which has shown in a Phase 1/2 to be safe and able to improve adaptive behavior in young adults with Down syndrome (Trisomy 21) and has started a Phase 2b in Europe aiming to confirm its efficacy and safety in people with Down syndrome. The clinical results obtained with these 2 compounds have confirmed the safety and therapeutic activity of CB_1-SSi in humans. The Company also develops a portfolio of new innovative CB_1-SSi for the treatment of

other disorders associated with a dysregulation of the activity of the CB_1 receptor, including diseases involving peripheral organs, such as obesity and related metabolic conditions. The new drugs developed by the Company belong to the same general pharmacological class, the CB_1-SSi, but have distinct functional effects allowing to target different types of dysregulations of the CB_1 receptor and guaranteeing that the different compounds are not substitutable one with the others.

For more information, visit www.aelisfarma.com and follow us on LinkedIn and Twitter.

ISIN: FR0014007ZB4
Ticker: AELIS
C Compartment of Euronext Paris

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Disclaimer

Forward-looking statements

Some information contained in this press release is forward-looking statements, not historical data. These forward-looking statements are based on current beliefs, expectations, and assumptions, including, but not limited to, assumptions about Aelis Farma's current and future strategy and the environment in which Aelis Farma operates. They involve known and unknown risks, uncertainties, and other factors, which may cause actual results, performance, achievements, or industry results or other events, to differ materially from those described or implied by such forward-looking statements. These risks and uncertainties include those set out and described in detail in Chapter 3 "Risk Factors" of Aelis Farma's Universal Registration Document filed with the *Autorité des Marchés Financiers* on April 23, 2026, under number D.26-0284.

These forward-looking statements are made only as of the date of this press release and Aelis Farma expressly disclaims any obligation or undertaking to release any updates or corrections to the forward-looking statements included in this press release to reflect any change in expectations or events, conditions, or circumstances on which any such forward-looking statement is based. Forward-looking information and statements are not guarantees of future performance and are subject to various risks and uncertainties, many of which are difficult to predict and generally beyond Aelis Farma's control. Actual results could differ materially from those described in, or implied or projected by, forward-looking information and statements.