

SEQUANA MEDICAL ANNOUNCES NEW SHARE CAPITAL AMOUNT AND NEW NUMBER OF SHARES

Ghent, Belgium, 26 September 2025 – Sequana Medical NV (Euronext Brussels: SEQUA) (the "Company" or "**Sequana Medical**"), a pioneer in the treatment of drug-resistant fluid overload in liver disease, heart failure and cancer, announces today a capital increase in the framework of (i) a contribution in kind of certain convertible receivables (for an aggregate amount of EUR 214,000.00) due under the convertible loan agreement entered into on 17 March 2025 between the Company and certain lenders (as amended), (ii) a contribution in kind of certain convertible receivables (for an aggregate amount of EUR 1,829,628.00) due under the loan agreement entered into on 19 July 2022 between the Company and Kreos Capital VII (UK) Limited (as amended), and (iii) contributions in cash in the framework of the outstanding "restricted share unit" or "RSU" plan for non-executive independent directors of the Company. As a result of aforementioned transactions, the Company's share capital has increased on 26 September 2025 from EUR 6,258,283.87 to EUR 6,542,176.98 and the number of issued and outstanding shares has further increased from 60,404,799 to 63,145,080 ordinary shares, through the issuance of a total of 2,740,281 new shares. The applicable issue prices of the relevant new shares were determined in accordance with the terms of the relevant convertible loan agreements and RSU plan.

The total current number of outstanding subscription rights amounts to 7,489,576, which entitles their holders (if exercised) to subscribe to 8,656,304 new shares with voting rights in total, namely:

- up to 261,895 new shares can be issued upon the exercise of 90,780 share options that are still outstanding under the 'Executive Share Options' plan for staff members and consultants of the Company, entitling the holder thereof to acquire ca. 2.88 new shares when exercising one of his or her share options (the "**Executive Share Options**");
- up to 687,784 new shares can be issued upon the exercise of 687,784 share options (each share option having the form of a subscription right) that are still outstanding under the '2018 Share Options' plan for directors, employees and other staff members of the Company and its subsidiaries, entitling the holder thereof to acquire one new share when exercising one of his or her share options (the "**2018 Share Options**");
- up to 188,370 new shares can be issued upon the exercise of 188,370 share options (each share option having the form of a subscription right) that are still outstanding under the '2021 Share Options' plan for directors, employees and other staff members of the Company and its subsidiaries, entitling the holder thereof to acquire one new share when exercising one of his or her share options (the "**2021 Share Options**");
- up to 1,000,000 new shares can be issued upon the exercise of 1,000,000 share options (each share option having the form of a subscription right) that are still outstanding under the '2023 Share Options' plan for directors, employees and other staff members of the Company and its subsidiaries, entitling the holder thereof to acquire one new share when exercising one of his or

her share options (the "**2023 Share Options**");

- up to 1,000,000 new shares can be issued upon the exercise of 1,000,000 share options (each share option having the form of a subscription right) that are still outstanding under the '2025 Share Options' plan for directors, employees and other staff members of the Company and its subsidiaries, entitling the holder thereof to acquire one new share when exercising one of his or her share options (the "**2025 Share Options**");
- up to 302,804 new shares can be issued to Bootstrap Europe S.C.SP. upon the exercise of 10 warrants (each warrant having the form of a subscription right) that are still outstanding that have been issued by the extraordinary shareholders meeting of 27 May 2022 (the "**Bootstrap Warrants**");
- up to 1,567,819 new shares can be issued to Kreos Capital VII Aggregator SCSp. upon the exercise of 875,000 warrants (each warrant having the form of a subscription right) that are still outstanding that have been issued by the extraordinary shareholders meeting of 20 December 2024 (the "**Kreos Warrants**")¹;
- up to 1,057,632 new shares can be issued upon exercise of 1,057,632 subscription rights that are still outstanding that have been issued by the board of directors (within the framework of the authorized capital) on 27 April 2023 and 10 May 2023 in the framework of the private placement of new shares and new subscription rights (the "**2023 Investor Warrants**"); and
- up to 2,590,000 new shares can be issued to GEM Global Yield LLC SCS ("**GEM**") upon the exercise of 2,590,000 warrants (each warrant having the form of a subscription right) that are still outstanding that have been issued by the extraordinary shareholders meeting of 22 May 2025, entitling GEM to acquire one new share when exercising one of its warrants (the "**GEM Warrants**").

This announcement is made in accordance with Article 15 of the Belgian Act of 2 May 2007 on the disclosure of major participations in issuers of which shares are admitted to trading on a regulated market and regarding miscellaneous provisions.

¹ The exercise price of the Kreos Warrants is equal to the lowest subscription price paid or agreed to be paid for a share in the share capital of the Company pursuant to any round of equity financing (or other financing convertible or exchangeable into equity) by the Company (taking into account any discounts including those arising on conversion or cancellation or indebtedness and/or interest thereon, but not taking into account any further anti-dilution adjustment mechanisms included in such rights or securities) prior to the exercise of the Kreos Warrants, and subject to certain exempted events that shall not be taken into account when determining the applicable exercise price per underlying new share. The number of new shares issuable upon exercise of the Kreos Warrants has been calculated on the basis of an exercise price that is equal to the lowest applicable issue price of the new shares issued on 24 January 2025 in the framework of contributions in kind of certain receivables (*i.e.*, EUR 0.5581 per share).

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About Sequana Medical

Sequana Medical NV is a pioneer in treating fluid overload, a serious and frequent clinical complication in patients with liver disease, heart failure and cancer. This causes major medical issues including increased mortality, repeated hospitalizations, severe pain, difficulty breathing and restricted mobility. Although diuretics are standard of care, they become ineffective, intolerable or exacerbate the problem in many patients. There are limited effective treatment options, resulting in poor clinical outcomes, high costs and a major impact on their quality of life. Sequana Medical is seeking to provide innovative treatment options for this large and growing "diuretic resistant" patient population. **alfapump**[®] and DSR[®] are Sequana Medical's proprietary platforms that work with the body to treat diuretic-resistant fluid overload, and are intended to deliver major clinical and quality of life benefits for patients, while reducing costs for healthcare systems.

The Company received US FDA approval for the **alfapump** System for the treatment of recurrent or refractory ascites due to liver cirrhosis in December 2024, following the grant of FDA Breakthrough Device Designation in 2019. Sequana Medical has commenced US commercialisation through its specialty commercial team that will target the 90 US liver transplant centers that perform more than 90% of liver transplants. In August 2025, CMS announced that it approved the New Technology Add-on Payment for the **alfapump** when performed in the hospital inpatient setting as of October 1, 2025.

Results of the Company's RED DESERT and SAHARA proof-of-concept studies in heart failure published in European Journal of Heart Failure in April 2024 support DSR's mechanism of action as breaking the vicious cycle of cardiorenal syndrome. All three patients from the non-randomized cohort of MOJAVE, a US randomized controlled multi-center Phase 1/2a clinical study, have been successfully treated with DSR, resulting in a dramatic improvement in diuretic response and virtual elimination of loop diuretic requirements². The independent Data Safety Monitoring Board approved the start of the randomized MOJAVE cohort of up to a further 30 patients, which is dependent on securing additional financing.

Sequana Medical is listed on the regulated market of Euronext Brussels (Ticker: SEQUA.BR) and headquartered in Ghent, Belgium. For further information, please visit www.sequanamedical.com.

Important Safety Information: For important safety information regarding the **alfapump**[®] system, see <https://www.sequanamedical.com/wp-content/uploads/ISI.pdf>.

The **alfapump**[®] System is currently not approved in Canada.

² Data reported in press release of [March 25, 2024](#); mean increase of 326% in six-hour urinary sodium excretion at 3 months follow up vs baseline, and 95% reduction of loop diuretics over same period.

DSR® therapy is still in development and is currently not approved in any country. The safety and effectiveness of DSR® therapy has not been established.

Note: **alfapump**® and DSR® are registered trademarks.

Forward-looking statements

This press release may contain predictions, estimates or other information that might be considered forward-looking statements. Such forward-looking statements are not guarantees of future performance. These forward-looking statements represent the current judgment of Sequana Medical on what the future holds, and are subject to risks and uncertainties that could cause actual results to differ materially. Sequana Medical expressly disclaims any obligation or undertaking to release any updates or revisions to any forward-looking statements in this press release, except if specifically required to do so by law or regulation. You should not place undue reliance on forward-looking statements, which reflect the opinions of Sequana Medical only as of the date of this press release.