

## **Sequana Medical announces Special General Meeting of Shareholders on 26 June 2023**

**Ghent, Belgium – 26 May 2023 – Sequana Medical NV (Euronext Brussels: SEQUA)** (the "Company" or "Sequana Medical"), a pioneer in the treatment of fluid overload in liver disease, heart failure and cancer, today invites the holders of securities issued by the Company to attend the Special General Meeting of Shareholders on Monday, 26 June 2023.

The items on the agendas of the meeting include the proposed approval of the appointment of Dr Kenneth Macleod as non-executive director of the Company and the approval in accordance with Article 7:151 of the Belgian Companies and Associations Code of the terms and conditions of the subscription rights issued in the framework of the Offering in the event of certain change of control events.

The Special General Meeting of Shareholders will take place at the Company's registered offices in Ghent and will start at 09:00 am CEST. The full convening notice with the agenda and proposed resolutions can be accessed on the Sequana Medical website: [www.sequanamedical.com/investors/shareholder-information](http://www.sequanamedical.com/investors/shareholder-information).

The Company recommends the holders of its securities to use e-mail for all communications with the Company regarding the Special General Meeting of Shareholders. The Company's e-mail address for such communications is: [IR@sequanamedical.com](mailto:IR@sequanamedical.com).

### **For more information, please contact:**

#### **Sequana Medical**

Lies Vanneste  
Director Investor Relations  
E: [IR@sequanamedical.com](mailto:IR@sequanamedical.com)  
T: +32 (0)498 053579

#### **Optimum Strategic Communications**

Nick Bastin, Jonathan Edwards, Vici Rabbetts  
E: [Sequana@optimumcomms.com](mailto:Sequana@optimumcomms.com)  
T: +44 (0) 208 078 4357

#### **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. These patients can have up to 15 liters of extra fluid in their bodies, causing major medical issues including increased mortality, repeated hospitalizations, severe pain, difficulty breathing and restricted mobility that severely impacts daily life. Although diuretics are standard of care, the problem is that in many patients they are no longer effective and / or tolerable. There are limited effective treatment options for these patients, 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**<sup>®</sup> and **DSR**<sup>®</sup> are Sequana Medical's proprietary platforms that work with the body to treat diuretic-resistant fluid overload, delivering major clinical and quality of life benefits for patients and reducing costs for healthcare systems. The Company has reported positive primary endpoint data from the North American pivotal POSEIDON trial of the **alfapump** in recurrent or refractory ascites due to liver cirrhosis, enabling the filing of a Pre-Market Approval (PMA) application with the FDA, planned for H2 2023. Having delivered clinical proof-of-concept for **DSR** as a disease-modifying drug program for the treatment of heart failure, the Company will commence **MOJAVE**, a US randomized controlled multi-center Phase 1/2a clinical trial of **DSR 2.0**, with initial data expected in Q4 2023.

Sequana Medical is listed on Euronext Brussels (Ticker: SEQUA.BR) and headquartered in Ghent, Belgium. For further information, please visit [www.sequanamedical.com](http://www.sequanamedical.com).

### **Important Regulatory Disclaimers**

*The **alfapump**<sup>®</sup> system is currently not approved in the United States or Canada. In the United States and Canada, the **alfapump** system is currently under clinical investigation (POSEIDON Trial) and is being studied in adult patients with refractory or recurrent ascites due to cirrhosis. For more information regarding the POSEIDON clinical trial see [www.poseidonstudy.com](http://www.poseidonstudy.com). **DSR**<sup>®</sup> therapy is still in development and it should be noted that any statements regarding safety and efficacy arise from ongoing pre-clinical and clinical investigations which have yet to be completed. There is no link between **DSR** therapy and ongoing investigations with the **alfapump** system in Europe, the United States or Canada.*

*Note: **alfapump**<sup>®</sup> and **DSR**<sup>®</sup> 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.*