Sequana Medical to present at Needham's Healthcare Conference and attend Kempen's Life Sciences Conference in April 2022

Ghent, Belgium – 8 April 2022 – Sequana Medical NV (Euronext Brussels: SEQUA) (the "**Company**" or "**Sequana Medical**"), an innovator in the treatment of diuretic-resistant fluid overload in liver disease, malignant ascites and heart failure, today announces that it will attend the following upcoming investor conferences in April 2022:

- **21st Annual Needham Healthcare Conference, Virtual, 11-14 April 2022**
  Presentation by Ian Crosbie, CEO, on Thursday 14 April at 08:45am ET / 14:45 CET
- **14th Kempen Life Sciences Conference, Amsterdam, 20-21 April 2022**
  Participation by Sequana Medical on Wednesday 20 April

Sequana Medical will be meeting with international investors in 1-to-1 and small group meetings.
Presentation slides will be available on Sequana Medical's Investors website shortly after the events.

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About Sequana Medical

Sequana Medical is a commercial stage medical device company utilizing its proprietary **alfa**pump® and DSR® (Direct Sodium Removal) technologies to develop innovative treatments for fluid overload in liver disease, malignant ascites and heart failure where diuretics are no longer effective. Fluid overload is a frequent complication of many large diseases – including advanced liver disease driven by NASH (non-alcoholic steatohepatitis)-related cirrhosis and heart failure – with diuretic resistance being widespread. The U.S. market for the **alfa**pump resulting from NASH-related cirrhosis is forecast to exceed €3 billion annually within the next 10-20 years. The heart failure market for DSR and the **alfa**pump DSR® is estimated to be over €5 billion annually in the U.S. and EU5 by 2026.

The **alfa**pump is Sequana Medical's unique, fully implanted wireless device that automatically pumps fluid from the abdominal cavity into the bladder, where it is naturally eliminated through urination. DSR is Sequana Medical's proprietary approach to managing sodium and fluid overload (congestion) through use of a sodium-free infusate administered into the abdominal cavity.

In the U.S., the Company's key growth market, the **alfa**pump has been granted breakthrough device designation by the FDA for recurrent or refractory ascites due to liver cirrhosis. Interim data from the ongoing North American pivotal study (POSEIDON) showed positive outcomes against all primary endpoints, rapid and persistent clinically important improvement in quality of life as well as a mean survival probability of 70% at 12 months post-implantation (compared to 50% survival rate for refractory ascites patients in the published literature). All patients have been implanted with the **alfa**pump and primary endpoint reporting is planned for Q4 2022. This study is intended to support a future marketing application of the **alfa**pump in the U.S. and Canada. In Europe, the **alfa**pump is CE-marked for the management of refractory ascites due to liver cirrhosis and malignant ascites and is included in key clinical practice guidelines. Over 900 **alfa**pump systems have been implanted to date.

Sequana Medical has combined its proven **alfa**pump and proprietary DSR therapy, and is developing the **alfa**pump DSR, a breakthrough approach to fluid overload due to heart failure. Top-line results of the RED DESERT study and interim results of the SAHARA DESERT study indicate that repeated DSR therapy in diuretic-resistant heart failure patients is able to safely, effectively and rapidly eliminate persistent congestion and restore euvolemia, improve cardio-renal status and restore diuretic response for months post-treatment. Reporting of top-line data for SAHARA DESERT is planned for H2 2022.

Sequana Medical is headquartered in Ghent, Belgium. For further information, please visit www.sequanamedical.com.

Important Regulatory Disclaimers

*The **alfa**pump® system is not currently approved in the United States or Canada. In the United States and Canada, the **alfa**pump system is currently under clinical investigation (POSEIDON Study) and is being studied in adult patients with refractory or recurrent ascites due to cirrhosis. For more information regarding the POSEIDON clinical study see www.poseidonstudy.com. The DSR® 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. The DSR therapy is not currently approved for clinical research in the United States or Canada. There is no link between the DSR therapy and ongoing investigations with the **alfa**pump system in Europe, the United States or Canada.*

***Note: alfa**pump® is a registered trademark. DSR® is a registered trademark in Australia, the Benelux, the EU, United Kingdom, Hong Kong, Israel, Norway, and Switzerland. **alfa**pump DSR® is a registered trademarks in Australia, the Benelux, China, the EU, United Kingdom, Hong Kong, Israel, New Zealand, and Norway.*

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.