

# Sequana Medical announces September 2020 Investor Conference Schedule

**Ghent, BELGIUM – 09 September 2020 – Sequana Medical NV (Euronext Brussels: SEQUA)**, an innovator in the management of fluid overload in liver disease, malignant ascites and heart failure, today announces that it will participate in two virtual investor events in September 2020.

### H.C. Wainwright & Co 22<sup>nd</sup> Annual Global Investment Conference, 14-16 September 2020

### KBC Life Sciences Conference, 22-23 September 2020

Company presentation by Ian Crosbie, CEO, *Tuesday 22 September 2020 at 10:00 am CEST* The presentation will be available on the Sequana Medical <u>Investors website</u> shortly after the event.

To request a one on one meeting with Sequana Medical management at one of these events, contact us at IR@sequanamedical.com.

### For more information, please contact:

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### About Sequana Medical

Sequana Medical is a commercial stage medical device company developing the **alfa**pump platform for the management of fluid overload in liver disease, malignant ascites and heart failure where diuretics are no longer effective. Fluid overload is a fast growing complication of advanced liver disease driven by NASH (non-alcoholic steatohepatitis) related cirrhosis and a common complication in heart failure. 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 the **alfa**pump DSR (Direct Sodium Removal) is estimated to be over €5 billion annually in the U.S. and EU5 by 2026. Both indications leverage Sequana Medical's **alfa**pump, a unique, fully implanted wireless device that automatically pumps fluid from the abdomen into the bladder, where it is naturally eliminated through urination.

In the U.S., the company's key growth market, the **alfa**pump has been granted breakthrough device designation by the FDA. The North American pivotal study (POSEIDON) in recurrent and refractory ascites due to liver cirrhosis is currently underway, and is intended to support a commercial 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 800 **alfa**pump systems have been implanted to date. Building on its proven **alfa**pump platform, Sequana Medical is developing the **alfa**pump DSR, a breakthrough, proprietary approach to fluid overload due to heart failure. Clinical proofof-concept was achieved in a first-in-human single dose DSR study and a repeated dose **alfa**pump DSR study (RED DESERT) in heart failure patients is currently underway. Sequana Medical is headquartered in Ghent, Belgium. For further information, please visit <u>www.sequanamedical.com</u>.

### Important Regulatory Disclaimers

The **alfa**pump<sup>®</sup> system is not currently approved in the United States or Canada. In the United States and Canada, the **alfa**pump<sup>®</sup> 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<sup>®</sup> system in Europe.

### Forward-looking statements

This press release may contain predictions, estimates or other information that might be considered forwardlooking 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.