



XVIVO Perfusion intends to carry out a directed issue of shares

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XVIVO Perfusion AB (publ) ("XVIVO" or the "Company") (Nasdaq Stockholm: XVIVO) intends to carry out a new issue of shares of approximately SEK 500 million directed to Swedish and international institutional investors (the "Directed Issue"). XVIVO has appointed Carnegie Investment Bank AB (publ) ("Carnegie") to act as Sole Global Coordinator and Bookrunner in connection with the Directed Issue.

XVIVO announced earlier today that the Company has entered into an agreement to acquire the Dutch medtech company Organ Assist B.V. ("Organ Assist"), for a total cash consideration of up to EUR 24 million (see the separate press release on the Company's website). The completion of the acquisition is conditional upon XVIVO obtaining financing through the Directed Issue.

XVIVO intends to carry out the Directed Issue based on the authorization to issue shares granted by the annual general meeting on March 31, 2020. The subscription price and the total number of new shares in the Directed Issue will be determined through an accelerated bookbuilding procedure, which will commence immediately following the publication of this press release. Pricing and allocation of the new shares are expected to take place before the commencement of trading on Nasdaq Stockholm at 09:00 CEST on September 24, 2020. By establishing the subscription price in the Directed Issue through an accelerated bookbuilding procedure, it is the assessment of the Board of Directors that the subscription price will be set on market terms and conditions. The closing, pricing and allocation in the bookbuilding procedure are determined at the discretion of the Company and may be cancelled at any time, meaning the Company may refrain from carrying out the Directed Issue. The Company will announce the outcome of the Directed Issue in a subsequent press release after the bookbuilding procedure has been completed.

The reason for deviating from the shareholders' preferential rights in the Directed Issue is to raise capital in a time and cost-effective manner as well as to further diversify the shareholder base with Swedish and international institutional investors.

The Company intends to use the net proceeds from the Directed Issue for:

- 1. financing of the acquisition of Organ Assist;
- 2. funding the FDA (US Food and Drug Administration) 510k[1] regulatory approval process in the US of Organ Assist's Kidney Assist Transport device;
- 3. funding the regulatory approval processes of the Liver Assist in combination with the STEEN Solution™

technology in the US and other key markets; and

4. continue to build and strengthen the organization to support the Company's growth strategy and for general corporate purposes.

In connection with the Directed Issue, the Company has agreed to a lock-up undertaking, with customary exceptions, on future share issuances for a period of 180 calendar days after the settlement date of the Directed Issue. In addition, the members of the Board of Directors and certain members of the senior management of the Company, have agreed not to sell any shares in XVIVO for a period of 180 calendar days after the settlement date of the Directed Issue, subject to customary exceptions.

Advisers

Carnegie Investment Bank AB (publ) is Sole Global Coordinator and Bookrunner in connection with the Directed Issue. Gernandt & Danielsson Advokatbyrå is legal adviser to the Company and Baker McKenzie is legal adviser to Carnegie in connection with the Directed Issue.

September 23, 2020 Gothenburg XVIVO Perfusion AB (publ)

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This press release does not identify or suggest, or purport to identify or suggest, the risks (direct or indirect) that may be associated with an investment in the new shares. Any investment decision in connection with the directed share issue must be made on the basis of all publicly available information relating to the Company and the Company's shares. Such information has not been independently verified by Carnegie. Carnegie acting for the Company in connection with the transaction and no one else and will not be responsible to anyone other than the Company for providing the protections afforded to its clients nor for giving advice in relation to the transaction or any other matter referred to herein.

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Forward-looking statements

This press release contains forward-looking statements that reflect the Company's intentions, beliefs, or current expectations about and targets for the Company's future results of operations, financial condition, liquidity, performance, prospects, anticipated growth, strategies and opportunities and the markets in which the Company operates. Forward-looking statements are statements that are not historical facts and may be identified by words such as "believe", "expect", "anticipate", "intend", "may", "plan", "estimate", "will", "should", "could", "aim" or "might", or, in each case, their negative, or similar expressions. The forwardlooking statements in this press release are based upon various assumptions, many of which are based, in turn, upon further assumptions. Although the Company believes that the expectations reflected in these forward-looking statements are reasonable, it can give no assurances that they will materialize or prove to be correct. Because these statements are based on assumptions or estimates and are subject to risks and uncertainties, the actual results or outcome could differ materially from those set out in the forward-looking statements as a result of many factors. Such risks, uncertainties, contingencies and other important factors could cause actual events to differ materially from the expectations expressed or implied in this release by such forward-looking statements. The Company does not guarantee that the assumptions underlying the forward-looking statements in this press release are free from errors and readers of this press release should not place undue reliance on the forward-looking statements in this press release. The information, opinions and forward-looking statements that are expressly or implicitly contained herein speak only as of its date and

are subject to change without notice. Neither the Company nor anyone else undertake to review, update, confirm or to release publicly any revisions to any forward-looking statements to reflect events that occur or circumstances that arise in relation to the content of this press release, unless it is not required by law or Nasdaq Stockholm's rule book for issuers.

Information to distributors

Solely for the purposes of the product governance requirements contained within: (a) EU Directive 2014/65 /EU on markets in financial instruments, as amended ("MiFID II"); (b) Articles 9 and 10 of Commission Delegated Directive (EU) 2017/593 supplementing MiFID II; and (c) local implementing measures (together, the "MiFID II Product Governance Requirements"), and disclaiming all and any liability, whether arising in tort, contract or otherwise, which any "manufacturer" (for the purposes of the MiFID II Product Governance Requirements) may otherwise have with respect thereto, the shares in XVIVO have been subject to a product approval process, which has determined that such shares are: (i) compatible with an end target market of retail investors and investors who meet the criteria of professional clients and eligible counterparties, each as defined in MiFID II; and (ii) eligible for distribution through all distribution channels as are permitted by MiFID II (the "Target Market Assessment"). Notwithstanding the Target Market Assessment, Distributors should note that: the price of the shares in XVIVO may decline and investors could lose all or part of their investment; the shares in XVIVO offer no guaranteed income and no capital protection; and an investment in the shares in XVIVO is compatible only with investors who do not need a guaranteed income or capital protection, who (either alone or in conjunction with an appropriate financial or other adviser) are capable of evaluating the merits and risks of such an investment and who have sufficient resources to be able to bear any losses that may result therefrom. The Target Market Assessment is without prejudice to the requirements of any contractual, legal or regulatory selling restrictions in relation to the directed share issue. Furthermore, it is noted that, notwithstanding the Target Market Assessment, Carnegie will only procure investors who meet the criteria of professional clients and eligible counterparties.

For the avoidance of doubt, the Target Market Assessment does not constitute: (a) an assessment of suitability or appropriateness for the purposes of MiFID II; or (b) a recommendation to any investor or group of investors to invest in, or purchase, or take any other action whatsoever with respect to the shares in XVIVO.

Each distributor is responsible for undertaking its own target market assessment in respect of the shares in XVIVO and determining appropriate distribution channels.

This is a translation of the Swedish version of the press release. In case of discrepancies, the Swedish wording shall prevail.

[1] A 510k is a premarket submission made to the FDA to demonstrate that the device to be marketed is as safe and effective (i.e., substantially equivalent) as a legally approved marketed device.

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About Us

XVIVO Perfusion AB is a medical technology company which develops solutions and systems for assessing and preserving organs outside the body and for selecting usable organs and maintaining them in optimal condition pending transplantation. The company is headquartered in Gothenburg, Sweden, and has one office in Lund, Sweden and one office in the USA. The XVIVO share is listed on Nasdaq Stockholm and has the ticker symbol XVIVO. More information can be found on the website www.xvivoperfusion.com.

This information is information that XVIVO Perfusion AB is obliged to make public pursuant to the EU Market Abuse Regulation. The information was submitted for publication, through the agency of the contact persons set out above, at 2020-09-23 17:31 CEST.

Attachments

XVIVO Perfusion intends to carry out a directed issue of shares