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# Xbrane Biopharma announces intention to carry out a directed share issue

Xbrane Biopharma AB ("Xbrane Biopharma" or the "Company") hereby announces its intention to carry out a directed share issue of approximately SEK 350 million to institutional investors through an accelerated book-building procedure, pursuant to the authorization granted by the annual general meeting held on May 6, 2021 (the "Directed Share Issue").

The Directed Share Issue is intended to be resolved by the board of directors pursuant to the authorization granted by the annual general meeting held on May 6, 2021 and is intended to be carried out with deviation from the shareholders' preferential rights. The Directed Share Issue will be directed to Swedish and international institutional investors. The price of the new shares in the Directed Share Issue will be determined through an accelerated book-building procedure, which will commence immediately after publication of this press release and end before commencement of trading on Nasdaq Stockholm on July 1, 2021. The book-building procedure may, at the discretion of the Company, close earlier or later and may be cancelled at any time.

The purpose of the Directed Share Issue and the reason for the deviation from the shareholders' preferential rights, is to diversify the shareholder base among international institutional investors and at the same time raise capital on favorable terms in a time and cost-efficient manner.

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

- i. Submission of Marketing Authorization Application for Xlucane™ in the US and Europe and production of launch material
- ii. Scale-up of Xcimzane™ production process to commercial scale and initiation of Phase I clinical trial together with commercialization partner
- iii. Continued pre-clinical development of Xdivane™ and initiation of additional biosimilar development programs
- iv. General corporate purposes

In connection with the Directed Share Issue, the Company has, subject to customary exceptions, agreed to a lock-up undertaking on future share issuances for a period of 180 days following completion of the Directed Share Issue. Members of the Company's board of directors and management, have, subject to customary exceptions, agreed to not sell their shares in the Company for a period of 180 days after the date of registration of the shares issued in connection with the Directed Share Issue with the Swedish Companies Registration Office. The existing shareholders STADA Arzneimittel AG and Serendipity Group AB have, subject to customary



exceptions, agreed to not sell their shares in the Company for a period of 90 calendar days after the date of registration of the shares issued in connection with the Directed Share Issue with the Swedish Companies Registration Office.

#### **Advisers**

Kempen & Co and Pareto Securities have been appointed Joint Global Coordinators and Joint Bookrunners. Handelsbanken Capital Markets has been appointed Co-Manager. Baker McKenzie Advokatbyrå KB is acting as legal adviser to the Company. White & Case Advokat AB is acting as legal adviser to the Joint Bookrunners and Co-Manager in connection with the Directed Share Issue.

# Important information

The release, announcement or distribution of this press release may, in certain jurisdictions, be subject to restrictions. The recipients of this press release in jurisdictions where this press release has been published or distributed shall inform themselves of and follow such restrictions. The recipient of this press release is responsible for using this press release, and the information contained herein, in accordance with applicable rules in each jurisdiction. This press release does not constitute an offer, or a solicitation of any offer, to buy or subscribe for any securities in Xbrane Biopharma in any jurisdiction, neither from Xbrane Biopharma nor from someone else.

This announcement 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 Joint Global Coordinators and Joint Bookrunners. The information contained in this announcement is for background purposes only and does not purport to be full or complete. No reliance may be placed for any purpose on the information contained in this announcement or its accuracy or completeness. Joint Global Coordinators and Joint Bookrunners is 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.

This announcement does not constitute a recommendation concerning any investor's option with respect to the Directed Share Issue. Each investor or prospective investor should conduct his, her or its own investigation, analysis and evaluation of the business and data described in this announcement and publicly available information. The price and value of securities can go down as well as up. Past performance is not a guide to future performance.

This press release does not constitute or form part of an offer or solicitation to purchase or subscribe for securities in the United States. The securities referred to herein may not be sold in the United States absent registration or an exemption from registration under the US Securities Act of 1933, as amended (the "Securities Act"), and may not be offered or sold within the United States absent registration or an applicable exemption from, or in a transaction not subject to, the



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This press release is not a prospectus for the purposes of Regulation (EU) 2017/1129 of the European Parliament and of the Council of June 14, 2017 (the "**Prospectus Regulation**") and has not been approved by any regulatory authority in any jurisdiction. Xbrane Biopharma has not authorized any offer to the public of shares or rights in any member state of the EEA and no prospectus for an offering has been or will be prepared in connection with the Directed Share Issue. In any EEA Member State, this communication is only addressed to and is only directed at qualified investors in that Member State within the meaning of the Prospectus Regulation.

In the United Kingdom, this document and any other materials in relation to the securities described herein is only being distributed to, and is only directed at, and any investment or investment activity to which this document relates is available only to, and will be engaged in only with, "qualified investors" who are (i) persons having professional experience in matters relating to investments who fall within the definition of "investment professionals" in Article 19(5) of the Financial Services and Markets Act 2000 (Financial Promotion) Order 2005 (the "**Order**"); (ii) high net worth entities falling within Article 49(2)(a) to (d) of the Order; or (iii) any other person to whom it may otherwise lawfully be communicated (all such persons together being referred to as "relevant persons"). In the United Kingdom, any investment or investment activity to which this communication relates is available only to, and will be engaged in only with, relevant persons. Persons who are not relevant persons should not take any action on the basis of this press release and should not act or rely on it.

# 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 forward-looking 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 Nasdag 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 Xbrane Biopharma 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 Xbrane Biopharma may decline and investors could lose all or part of their investment; the shares in Xbrane Biopharma offer no guaranteed income and no capital protection; and an investment in the shares in Xbrane Biopharma 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, Joint Global Coordinators and Joint Bookrunners 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 Xbrane Biopharma.



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

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#### **About Us**

Xbrane Biopharma AB develops biological drugs based on a platform technology that provides significantly lower production costs compared to competing systems. Xbrane's leading product Xlucane™, a Lucentis® biosimilar candidate, addresses the € 10.4bn ophthalmic VEGFa inhibitor market. Marketing authorization of Xlucane™ is expected for the second half of 2022. Xbrane has additionally two biosimilars in its pipeline targeting € 7.9bn in originator sales. Xbrane's head office is in Solna, just outside Stockholm. Xbrane is listed on Nasdaq Stockholm under the ticker XBRANE. For more information, visit www.xbrane.com.

This information is information that Xbrane Biopharma 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 2021-06-30 17:31 CEST.

# **Attachments**

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