Regulatory press release



## Enzymatica publishes supplementary prospectus

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The Board of Directors of Enzymatica AB ("Enzymatica" or "Company") has issued a supplement (the "Supplementary Prospectus") in addition to the EU growth prospectus that has been approved and registered by the Swedish Financial Supervisory Authority (Finansinspektionen) and was published on 5 August 2022 (the "Prospectus"). The Supplementary Prospectus has today been approved by Finansinspektionen and has been published on the Company's website, as well as in other channels.

The Supplementary Prospectus has been prepared as Enzymatica on 16 August 2022 announced through a press release that a new study from the Medical University of Innsbruck shows that the Company's CE marked product ColdZyme protects airway epithelium from infection by the widespread omicron variants BA.4 and BA.5. The press release is available on the Company's website, <a href="www.enzymatica.com">www.enzymatica.com</a>.

Investors who, prior to the publication of this Supplementary Prospectus, have made a subscription or otherwise agreed to subscribe for shares in the rights issue that Enzymatica issues on 19 July 2022 (the "Rights Issue") are entitled under Article 23 (2) of Regulation (EU) 2017/1129 to withdraw their subscription or consent within three working days of the publication of the Supplementary Prospectus, i.e. until 22 August 2022. Withdrawal must be made in writing to Erik Penser Bank, Box 7405, 103 91 Stockholm, or via e-mail to emission@penser.se. Investors who have subscribed for shares in the rights issue through a nominee must contact their nominee for withdrawal. Subscriptions that have not been withdrawn within the specified time will remain binding and investors who wish to remain with their subscription for shares in the rights issue do not need to take any action.

The Prospectus and the Supplementary Prospectus are available on <a href="www.penser.se">www.penser.se</a> and <a href="www.enzymatica.com">www.enzymatica.com</a>. For terms and conditions and other information about the Rights Issue, please refer to the Prospectus.

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Enzymatica AB is a life science company that develops and sells health products mainly to treat diseases and symptoms in the upper respiratory tract. The products are based on a barrier technology that includes marine enzymes with unique properties. The company's first product is the medical device product ColdZyme®, a mouth spray for colds. The product has been launched in about 30 markets on four continents. The strategy is to continue to grow by developing more health products, strengthening the company's position in existing markets and expanding into new geographic markets through established partners. The company is headquartered in Lund and is listed on Nasdaq First North Growth Market. For more information, please visit www.enzymatica.com.



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A prospectus regarding the Rights issue described in this release has been published by the Company on 5 August 2022. This release is however not a prospectus in accordance to the definition in the Prospectus Regulation. In accordance with article 2 k of the Prospectus Regulation this press release constitutes an advertisement. Complete information regarding the Rights issue can only be obtained through the Prospectus. Enzymatica has not authorized any offer to the public of shares or rights in any other member state of the EEA. 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. 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 Rights 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 the financial adviser. The financial adviser 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.



## 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 Enzymatica 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 Enzymatica may decline and investors could lose all or part of their investment; the shares in Enzymatica offer no guaranteed income and no capital protection; and an investment in the shares in Enzymatica 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 Rights issue.

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 Enzymatica.

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