



ERYTECH Announces Trading Suspension of Its Ordinary Shares on Euronext Paris

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Regulatory News:

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Trading in the ordinary shares of ERYTECH Pharma (Euronext Paris: ERYP) ("ERYTECH" or "the Company") (Paris:ERYP) (ADR:EYRY), was suspended at the request of the Company on November 10, 2017 from 9:00 am CET in connection with its previously announced global offering in order to allow for the confirmation of allocations to investors and for the commencement of trading of the Company's American Depositary Shares ("ADSs") on the Nasdaq Global Select Market.

This suspension will be effective until a new communication is released by the Company. Trading on the regulated market of Euronext Paris is expected to resume today, November 10, 2017, at approximately 4:00 pm (CET), which is approximately the same time as the ADSs are expected to begin trading on the Nasdaq Global Select Market (10:00 am (EST)) under the ticker symbol "ERYP". The Company expects to announce the terms of the global offering later today.

A registration statement on Form F-1 relating to the global offering and the securities offered thereby has been filed with the U.S. Securities and Exchange Commission and was declared effective on November 9, 2017.

The securities referred to in this press release are being offered only by means of a U.S. prospectus. When available, copies of the final U.S. prospectus relating to and describing the terms of the global offering may be obtained for free from: Jefferies LLC, Attention: Equity Syndicate Prospectus Department, 520 Madison Avenue, 2nd Floor, New York, NY 10022, or by telephone at (877) 821-7388, or by email at prospectus_department@jefferies.com (mail to: Prospectus_Department@Jefferies.com); or from Cowen and Company, LLC, c/o Broadridge Financial Solutions, 1155 Long Island Avenue, Edgewood, NY 11717, Attention: Prospectus Department, or by telephone at (631) 274-2806.

Application will be made to list the ordinary shares to be issued pursuant to the global offering on the regulated market of Euronext Paris pursuant to a listing prospectus, subject to a visa application with the Autorité des Marchés Financiers ("AMF") and comprising the 2016 Reference Document (*Document de Référence*) of the Company registered with the AMF on March 31, 2017 under number D. 17-0283, the Actualization of the 2016 Reference Document registered with the AMF on October 6, 2017 under number D. 17-0283-A01 (*Actualisation du Document de Référence*) and a Securities Note (*Note d'opération*), including a summary of the prospectus. Copies of the 2016 Reference Document and in its Actualization are available free of charge at the Company's principal offices located at Bâtiment Adénine, 60 Avenue Rockefeller, 69008 Lyon, France, on the Company's website and on the website of the AMF.

This press release does not constitute an offer to sell or a solicitation of an offer to buy securities, and shall not constitute an offer, solicitation or sale in any state or jurisdiction in which such an offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction.

About ERYTECH and eryaspase (GRASPA®)

Founded in Lyon, France in 2004, ERYTECH is a clinical-stage biopharmaceutical company developing innovative therapies for rare forms of cancer and orphan diseases. Leveraging its proprietary ERYCAPS platform, which uses a novel technology to encapsulate therapeutic drug substances inside red blood cells, ERYTECH has developed a pipeline of product candidates targeting markets with high unmet medical needs. ERYTECH's initial focus is on the development of products that target the amino acid metabolism of cancer, depriving them of nutrients necessary for their survival.

The company's lead product, eryaspase, also known under the trade name GRASPA®, consists of an enzyme, L-asparaginase, encapsulated inside donor-derived red blood cells. L-asparaginase depletes asparagine, a naturally occurring amino acid essential for the survival and proliferation of cancer cells. L-asparaginase has been a standard component of multi-agent chemotherapy for the treatment of acute lymphoblastic leukemia (ALL), but side effects limit treatment, especially in adults and patients with weak performance status. With its improved safety profile, eryaspase aims to provide L-asparaginase to patients who cannot tolerate current non-encapsulated asparaginases.

Eryaspase in combination with chemotherapy achieved positive efficacy and safety results in a Phase 2/3 study in children and adults with relapsed or refractory ALL and in a Phase 2b clinical study in second-line metastatic pancreatic cancer. ERYTECH also has an ongoing Phase 1 clinical study of eryaspase in the United States in adults with newly diagnosed ALL and a Phase 2b clinical study in Europe in elderly patients with newly diagnosed acute myeloid leukemia (AML), each in combination with chemotherapy.

ERYTECH produces eryaspase at its own GMP-approved and operational manufacturing site in Lyon (France), and at a site for clinical production in Philadelphia (USA). ERYTECH has entered into licensing and distribution partnership agreements for eryaspase for ALL and AML in Europe with Orphan Europe (Recordati Group) and for ALL in Israel with TEVA, which will market the product under the GRASPA® brand name. The European Medicines Agency (EMA) and the U.S. Food and Drug Administration (FDA) have granted orphan drug designations for eryaspase for the treatment of ALL, AML and pancreatic cancer.

In addition to eryaspase, ERYTECH is developing two other product candidates, erymethionase and eryminase, that focus on using encapsulated enzymes to target cancer metabolism and induce tumor starvation. ERYTECH is also exploring the use of its ERYCAPS platform for developing cancer immunotherapies (ERYMMUNE) and enzyme replacement therapies (ERYZYME).

ERYTECH is listed on Euronext regulated market in Paris (ISIN code: FR0011471135, ticker: ERYP) and is part of the CAC Healthcare, CAC Pharma & Bio, CAC Mid & Small, CAC All Tradable, EnterNext PEA-PME 150 and Next Biotech indexes.

Forward-looking information

This press release contains forward-looking statements, forecasts and estimates with respect to the global offering, ERYTECH's clinical development plans, business and regulatory strategy, and the anticipated future performance of ERYTECH and of the market in which it operates. Certain of these statements, forecasts and estimates can be recognized by the use of words such as, without limitation, "believes", "anticipates", "expects", "intends", "plans", "seeks", "estimates", "may", "will" and "continue" and similar expressions. They include all matters that are not historical facts. Such statements, forecasts and estimates are based on various assumptions and assessments of known and unknown risks, uncertainties and other factors, which were deemed reasonable when made but may or may not prove to be correct. Actual events are difficult to predict and may depend upon factors that are beyond ERYTECH's control. There can be no guarantees with respect to pipeline product candidates that the candidates will receive the necessary regulatory approvals or that they will prove to be commercially successful. Therefore, actual results may turn out to be materially different from the anticipated future results, performance or achievements expressed or implied by such statements, forecasts and estimates. Documents filed by ERYTECH with the U.S. Securities and Exchange Commission and the AMF, also available on ERYTECH's website (www.erytech.com) describe such risks and uncertainties. Given these uncertainties, no representations are made as to the accuracy or fairness of such forward-looking statements, forecasts and estimates. Furthermore, forward-looking statements, forecasts and estimates only speak as of the date of this press release. Readers are cautioned not to place undue reliance on any of these forward-looking statements. ERYTECH disclaims any obligation to update any such forward-looking statement, forecast or estimates to reflect any change in ERYTECH's expectations with regard thereto, or any change in events, conditions or circumstances on which any such statement, forecast or estimate is based, except to the extent required by law.

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This announcement does not, and shall not, in any circumstances constitute a public offering nor an invitation to solicit the interest of the public in France, the United States, or in any other jurisdiction, in connection with any offer.

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This announcement is not an advertisement and not a prospectus within the meaning of Directive 2003/71/EC of the European Parliament and of the Council of 4 November 2003, as amended (the "Prospectus Directive").

With respect to the member States of the European Economic Area, no action has been undertaken or will be undertaken to make an offer to the public of the securities referred to herein requiring a publication of a prospectus in any relevant member State. As a result, the securities may not and will not be offered in any relevant member State except in accordance with the exemptions set forth in Article 3(2) of the Prospectus Directive or under any other circumstances which do not require the publication by the Company of a prospectus pursuant to Article 3 of the Prospectus Directive and/or to applicable regulations of that relevant member State.

This document does not constitute an offer to the public in France and the securities referred to in this document can only be offered or sold in France pursuant to article L. 411-2-II of the French Monetary and Financial Code to (i) providers of third party portfolio management investment services, (ii) qualified investors (*investisseurs qualifiés*) acting for their own account and/or (iii) a limited group of investors (*cercle restreint d'investisseurs*) acting for their own account, all as defined in and in accordance with articles L. 411-1, L. 411-2 and D. 411-1 to D. 411-4 and D. 754-1 and D. 764-1 of the French Monetary and Financial Code.

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