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In accordance with the provisions of article 227 of the Spanish Securities Markets and Investment Services Act (*Ley de los Mercados de Valores y de los Servicios de Inversión*), approved by Law 6/2023, of 17 March, and concordant provisions, is hereby reported the following:

OTHER RELEVANT INFORMATION

Pharma Mar, S.A. announces that its licensing partner, Luye Pharma Group Ltd, has received marketing approval for Zepzelca® (lurbinectedin) by the Pharmaceutical Administration Bureau in Macao for the treatment of adult patients with metastatic Small-Cell Lung Cancer (SCLC), with disease progression on or after platinum-based chemotherapy.

Please find attached press release that Pharma Mar, S.A. will distribute to the media.

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PharmaMar gets the approval of Zepzelca® (lurbinectedin) for the treatment of metastatic Small-Cell Lung Cancer in Macao, China

Madrid, December 4nd, 2023.- PharmaMar (MSE:PHM) has announced today that its licensing partner, Luye Pharma Group Ltd, has received marketing approval for Zepzelca® (lurbinectedin) by the Pharmaceutical Administration Bureau in Macao for the treatment of adult patients with metastatic Small-Cell Lung Cancer (SCLC), with disease progression on or after platinum-based chemotherapy.

The approval of lurbinectedin in Macao is mainly based on data from two clinical trials. One based on the data from the open-label, multi-center, single-arm monotherapy clinical trial in 105 adult patients with relapsed SCLC (including patients with platinum-sensitive and platinum-resistant disease), that the Food and Drug Administration (FDA) used to grant accelerated approval for lurbinectedin. The other, conducted in China, was a single-arm, dose-escalation, and dose-expansion clinical trial aiming to evaluate the safety, tolerability, pharmacokinetics and preliminary efficacy of lurbinectedin in Chinese patients with advanced tumors, including relapsed SCLC.

In April 2019, PharmaMar and Luye Pharma signed an agreement for the development and commercialization of lurbinectedin in SCLC.

Lurbinectedin is now approved in 14 countries around the world plus Macao. In the meantime, lurbinectedin Marketing Authorisation applications are under review for approval in Hong Kong and mainland China.

Lung cancer is the most common of all cancers in China in terms of incidence and mortality. In 2020, there were about 815,000 new cases of lung cancer and 714,000 deaths[1] caused by it, of which SCLC accounted for 13-17%[2].

Legal warning

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

About PharmaMar

PharmaMar is a biopharmaceutical company focused on the research and development of new oncology treatments, whose mission is to improve the healthcare outcomes of patients afflicted by serious diseases with our innovative medicines. The Company is inspired by the sea, driven by science, and motivated by patients with serious diseases to improve their lives by delivering novel medicines to them. PharmaMar intends to continue to be the world leader in marine medicinal discovery, development and innovation.
PharmaMar has developed and now commercializes Yondelis® in Europe by itself, as well as Zepzelca® (lurbinectedin), in the US; and Aplidin® (plitidepsin), in Australia, with different partners. In addition, it has a pipeline of drug candidates and a robust R&D oncology program. PharmaMar has other clinical-stage programs under development for several types of solid cancers: lurbinectedin, ecubectedin, PM534 and PM54. It also has a preclinical and clinical program in virology. Headquartered in Madrid (Spain), PharmaMar has subsidiaries in Germany, France, Italy, Belgium, Austria, Switzerland and The United States. PharmaMar also wholly owns Sylentis, a company dedicated to researching therapeutic applications of gene silencing (RNAi). To learn more about PharmaMar, please visit us at www.pharmamar.com

About Zepzelca®

Zepzelca® (lurbinectedin), also known as PM1183, is an analog of the marine compound ET-736 isolated from the sea squirt *Ecteinascidia turbinata* in which a hydrogen atom has been replaced by a methoxy group. It is a selective inhibitor of the oncogenic transcription programs on which many tumors are particularly dependent. Together with its effect on cancer cells, lurbinectedin inhibits oncogenic transcription in tumor-associated macrophages, downregulating the production of cytokines that are essential for the growth of the tumor. Transcriptional addiction is an acknowledged target in those diseases, many of them lacking other actionable targets.

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[1] International Agency for Research on Cancer – Population Fact Sheets – China. Available at https://gco.iarc.fr/today/data/factsheets/populations/160-china-fact-sheets.pdf. Accessed in October 2023
[2] The Working Committee of Chinese Society of Clinical Oncology Guidelines. The 2023 Chinese Society of Clinical Oncology Guidelines for Small Cell Lung Cancer. People's Medical Publishing House. 2023.

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