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In accordance with the provisions of article 226 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:

INSIDE INFORMATION

Pharma Mar, S.A. announces that the U.S. Food and Drug Administration (FDA) has granted approval for Zepzelca® (lurbinectedin) in combination with atezolizumab (Tecentriq®) as a maintenance treatment for adults with extensive-stage small cell lung cancer (ES-SCLC) whose disease has not progressed after first-line induction therapy with atezolizumab, carboplatin and etoposide. The approval marks the first combination therapy for first-line maintenance treatment of ES-SCLC, a fast-growing and aggressive cancer with limited treatment options.

Please find attached press release that will be distribute to the media.

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Press Release

FDA approves PharmaMar's Zepzelca® (lurbinectedin) and Atezolizumab (Tecentriq®) Combination as First-line Maintenance Therapy for Extensive-stage Small Cell Lung Cancer

- **Combination reduced the risk of disease progression or death by 46% and risk of death by 27% in the pivotal Phase 3 IMforte trial.**
- **Lurbinectedin and atezolizumab combination added to National Comprehensive Cancer Network® Guidelines for SCLC.**

Madrid, October 3rd, 2025.- PharmaMar (MSE:PHM) has announced that the U.S Food and Drug Administration (FDA) has granted approval for Zepzelca® (lurbinectedin) in combination with atezolizumab (Tecentriq®) as a maintenance treatment for adults with extensive-stage small cell lung cancer (ES-SCLC) whose disease has not progressed after first-line induction therapy with atezolizumab, carboplatin and etoposide[i]. The approval marks the first combination therapy for first-line maintenance treatment of ES-SCLC, a fast growing and aggressive cancer with limited treatment options.

The National Comprehensive Cancer Network® (NCCN®) recently updated the NCCN Clinical Practice Guidelines in Oncology® (NCCN Guidelines®) for SCLC to include the combination as preferred regimen for patients whose disease has not progressed following four cycles of platinum-based chemotherapy and atezolizumab induction.

The FDA approval is based on results from the Phase 3 IMforte trial, which showed that the lurbinectedin and atezolizumab combination reduced the risk of disease progression or death by 46% and the risk of death by 27%, compared to atezolizumab maintenance therapy alone. Following four cycles of induction therapy, from the point of randomization the median overall survival (OS) for the combination regimen was 13.2 months versus 10.6 months (stratified hazard ratio [HR]=0.73; 95% CI: 0.57–0.95; p=0.0174). From the point of randomization, median progression-free survival (PFS) by independent assessment was 5.4 months versus 2.1 months, respectively (stratified HR=0.54, 95% CI: 0.43–0.67; p<0.0001). Safety was consistent with the known safety profiles of both treatments. The results were presented at the 2025 American Society of Clinical Oncology (ASCO) annual meeting and simultaneously published in The Lancet.

PharmaMar has also submitted a Marketing Authorisation Application (MAA) to the European Medicines Agency (EMA), which is currently under review.

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. 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.

Tecentriq (atezolizumab) is a registered trademark of Genentech, a member of the Roche Group.

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[i] ZEPZELCA (lurbinectedin) Prescribing Information. Palo Alto, CA: Jazz Pharmaceuticals, Inc.