

PharmaMar receives Orphan Drug Designation for lurbinectedin from the Swiss Agency for Therapeutic Products for Small Cell Lung Cancer

Madrid, November 26th, 2019.- PharmaMar (PHM:MSE) has announced today that the Swiss Agency for Therapeutic Products (Swissmedic) has granted Orphan Drug designation to lurbinectedin for the treatment of Small Cell Lung Cancer (SCLC).

This decision is based on the orphan drug recognition granted by the European Medicines Agency on February 26th, 2019.

SCLC is PharmaMar's priority research area. On August 19th it was announced that the Company will file lurbinectedin's NDA for the treatment of relapsed SCLC with the FDA in the United States under the "accelerated approval" program during the last quarter of this year. This would open up the possibility for the FDA to approve lurbinectedin in the U.S. for the treatment of relapsed SCLC within the next year.

The ATLANTIS phase III trial of lurbinectedin in combination with doxorubicin for the treatment of this same type of tumor ended patient recruitment in July 2018 and is expected to have Overall Survival results in 2020.

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

Headquartered in Madrid, PharmaMar is a biopharmaceutical company, focused on oncology and committed to research and development which takes its inspiration from the sea to discover molecules with antitumor activity. It is a company that seeks innovative products to provide healthcare professionals with new tools to treat cancer. Its commitment to patients and to research has made it one of the world leaders in the discovery of antitumor drugs of marine origin.

PharmaMar has a pipeline of drug candidates and a robust R&D oncology program. It develops and commercializes Yondelis® in Europe and has other clinical-stage programs under development for several types of solid cancers: lurbinectedin (PM1183), PM184 and PM14. With subsidiaries in Germany, Italy, France, Switzerland, Belgium, Austria and the United States. PharmaMar wholly owns other companies:

GENOMICA, a molecular diagnostics company; Sylentis, dedicated to researching therapeutic applications of gene silencing (RNAi). To learn more about PharmaMar, please visit us at www.pharmamar.com.

About lurbinectedin

Lurbinectedin (PM1183) is a synthetic compound currently under clinical investigation. 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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