

OSE Immunotherapeutics to Present New Clinical and Preclinical Data on its Immuno-Oncology Portfolio at AACR 2019

- **Oral presentation will highlight early signs of activity of Tedopi® from the ongoing Phase 3 trial in Non-Small Cell Lung Cancer**

Nantes, France, Feb. 28, 2019, 7:00 a.m. CET – OSE Immunotherapeutics (ISIN: FR0012127173; Mnémo: OSE), today announced the presentation of new clinical data on Tedopi® and preclinical data on OSE-172 at the American Association of Cancer Research (AACR) Annual Meeting, taking place from March 29-April 3, in Atlanta.

In an oral presentation, Dr. Santiago Viteri* will provide details on early signs of activity of Tedopi®, a combination of neoepitopes, currently in an ongoing Phase 3 trial in advanced Non-Small Cell Lung Cancer in patients after failure to previous immune checkpoint inhibitors.

Presentation details:

Title: Early signs of activity of Tedopi® (OSE2101), a multiple neoepitope vaccine, in a phase 3 trial in advanced lung cancer patients after failure to previous immune checkpoint inhibitors (ATALANTE-1)

Session Category: Immunology

Session Title: Cancer Vaccines and Intratumoral Immunomodulation

Session Date and Time: Sunday, March 31, 2019, 3:00 p.m. - 5:00 p.m.

**Instituto Oncológico Dr Rosell, University Hospital Dexeus, QuironSalud Group, Barcelona*

In addition, the OSE team will present a poster with new preclinical data showing that selective SIRP α antagonist monoclonal antibody OSE-172, acting synergistically with T-cell checkpoint blockade, modifies the tumor microenvironment and therefore limits T-cell exclusion from tumor nest.

Presentation details:

Title: SIRP α blockade reinvigorates myeloid cells in the tumor microenvironment and reverses T-cell exclusion

Section Category: Immunology

Section Title: Immune Checkpoints 1

Section Date and Time: Tuesday, April 2, 2019, 8:00 a.m. – 12:00 p.m.

Location: Georgia World Congress Center, Exhibition Hall B, Poster Section 24

Poster board n° 18

Abstract n° 3238

ABOUT OSE Immunotherapeutics

OSE Immunotherapeutics is a clinical-stage biotechnology company focused on developing and partnering therapies to control the immune system for immuno-oncology and autoimmune diseases. The company has a diversified first-in-class clinical portfolio consisting of several scientific and technological platforms including neoepitopes and agonist or antagonist monoclonal antibodies, all ideally positioned to fight cancer and autoimmune diseases. The most advanced therapeutic-candidate, Tedopi[®], is a proprietary combination of 10 neo-epitopes aimed at stimulating T-lymphocytes and is currently in Phase 3 development in non-small cell lung cancer (NSCLC) after checkpoint inhibitor failure (anti PD-1 and anti PD-L1) and in Phase 2 testing in pancreatic cancer in combination with checkpoint inhibitor Opdivo[®]. FR104 (an anti-CD28 mAb) has successfully completed Phase 1 testing and has potential to treat autoimmune diseases. In April 2018, Boehringer Ingelheim and OSE signed a global license and collaboration agreement to develop preclinical checkpoint inhibitor OSE-172 (anti-SIRPa monoclonal antibody) in multiple cancer indications. OSE-127 (monoclonal antibody targeting the CD127 receptor, the alpha chain of the interleukin-7 receptor) is partnered with Servier under an option agreement up to the completion of a Phase 2 clinical trial planned in autoimmune bowel diseases; in parallel, Servier plans a development in the Sjögren syndrome. OSE-127 is currently under Phase 1 clinical trial.

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Forward-looking statements

This press release contains express or implied information and statements that might be deemed forward-looking information and statements in respect of OSE Immunotherapeutics. They do not constitute historical facts. These information and statements include financial projections that are based upon certain assumptions and assessments made by OSE Immunotherapeutics' management in light of its experience and its perception of historical trends, current economic and industry conditions, expected future developments and other factors they believe to be appropriate.

These forward-looking statements include statements typically using conditional and containing verbs such as "expect", "anticipate", "believe", "target", "plan", or "estimate", their declensions and conjugations and words of similar import. Although the OSE Immunotherapeutics management believes that the forward-looking statements and information are reasonable, the OSE Immunotherapeutics' shareholders and other investors are cautioned that the completion of such expectations is by nature subject to various risks, known or not, and uncertainties which are difficult to predict and generally beyond the control of OSE Immunotherapeutics. These risks could cause actual results and developments to differ materially from those expressed in or implied or projected by the forward-looking statements. These risks include those discussed or identified in the public filings made by OSE Immunotherapeutics with the AMF. Such forward-looking statements are not guarantees of future performance. This press release includes only summary information and should be read with the OSE Immunotherapeutics Reference Document filed with the AMF on 26 April 2018, including the annual financial report for the fiscal year 2017, available on the OSE Immunotherapeutics' website.

Other than as required by applicable law, OSE Immunotherapeutics issues this press release at the date hereof and does not undertake any obligation to update or revise the forward-looking information or statements.