Oxford Vacmedix announces major milestone for lead cancer vaccine OVM-200

OVM-200 Drug Product, manufactured to Good Manufacturing Process (GMP), has been certified stable and sterile four years post manufacture – a major milestone for OVM-200 development.

Oxford 23rd January 2025

Oxford Vacmedix (OVM), the UK-based biopharma company developing therapeutic cancer vaccines, announced today that OVM-200 which was manufactured on contract by Eurofins Amatsi to exacting GMP standards, has been tested and certified as both stable and sterile a full four years post-manufacture. The GMP drug substance for OVM-200 is dispensed into vials for use in clinical trials and is being used in the ongoing Phase 1 trial of OVM-200. Four-year stability is an important milestone in the development of cost-efficient cancer vaccines using the novel Recombinant Overlapping Peptide (ROP) technology pioneered by Professor Shisong Jiang.

OVM-200 is an ROP based vaccine that targets-survivin, over-expressed in many solid tumours. It is being trialled in the UK in four leading hospitals to treat ovarian cancer, prostate cancer and non-small cell lung cancer. Phase 1a, the first part of a Phase 1 trial, has been completed and has demonstrated both excellent safety and a strong immune response as well as allowing the selection of the optimal dose of OVM-200 for use to treat late-stage cancer. Phase 1b is ongoing.

Dr Debra Nevin, CEO and Senior Consultant at Amethyst Pharma commented;

“The Recombinant Overlapping Peptide platform technology is a completely new approach for making vaccines to treat cancer. Manufacturing these complex peptides is not straightforward and new techniques had to be developed to manufacture the first ROP vaccine, OVM-200, to GMP standard. The robustness of the manufacturing process is demonstrated in the drug product for OVM-200 showing both stability and sterility at four years after manufacture”.

William Finch, CEO of Oxford Vacmedix added;

“We are delighted to have reached this significant milestone for OVM-200. Stable drug product means that existing stock can be used for continuing progress with OVM-200 and that future manufacturing of ROP based vaccines is likely to have at least five-year stability. This will contribute towards cost efficient manufacturing and lower cost of goods. With ongoing testing, we plan to extend the stability profile of our ROP vaccines.”

ENDS

Oxford Vacmedix UK
The Magdalen Centre, Oxford Science Park, Oxford OX4 4GA, UK
T +44 (0)1865 742087 **E** enquiries@oxfordvacmedix.com www.oxfordvacmedix.com

For more information or to express an interest in investing in Series B please contact:

William Finch, CEO, Oxford Vacmedix
T: +44 (0)1865 742087 | M: +44(0)7769 903711 | E: wfinch@oxfordvacmedix.com

Notes to Editor

About Oxford Vacmedix

Oxford Vacmedix UK Ltd, based at the Oxford Science Park, UK, is a clinical stage bio-pharma company that was spun out from the University of Oxford's Department of Oncology and is utilising the novel proprietary platform technology of recombinant overlapping peptides (ROPs) invented by Professor Shisong Jiang. ROPs have been validated as a technology to stimulate broad and strong T cell immunity therefore forming a good platform for therapeutic vaccines and diagnostics in cancer and infectious diseases.

The technology uses the novel, proprietary platform of ROPs to design and develop therapeutic cancer vaccines and diagnostics with the potential for increased efficacy, lower costs, simpler regulatory pathways and synergy when used in combination with other immune oncology (IO) agents. The company has extensive contacts and collaborations in China through Changzhou Bioscience Group (CBIG) that is using the ROP platform for diagnostics in both cancer and in infectious diseases.

OVM is developing two lead vaccines, OVM-100 and OVM-200, focusing on unmet clinical need. OVM-100 is an HPV vaccine targeted at cervical cancer and head & neck cancer, and OVM-200 represents a new type of vaccine utilising survivin to target solid tumours including prostate, ovarian and non-small cell lung cancer (NSCLC). Both vaccines will be tested as single agents and in combination with IO agents. OVM has a strong pipeline, with a diagnostic for anti-microbial resistance being tested and two other cancer vaccines is preclinical development.

OVM secured Series A investment from Dx&Vx (formerly Cancer ROP), a leading South Korean biotech company, and from existing shareholders in China in 2018. The lead investment for Series B was recently made by Dx&Vx who have licensed OVM-200 for South Korea, China and India. Additional Series B funding is sought to advance OVM-200 to Phase 2 and OVM-100 into Phase 1 trials, as monotherapy and also in combination. In addition, the option of using mRNA delivery with the ROP technology will be explored.

For more information: http://www.oxfordvacmedix.com