NOXXON TO PRESENT TOP-LINE EFFICACY DATA FROM NOX-A12 / KEYTRUDA® COMBINATION TRIAL IN DECEMBER AS PLANNED AT ESMO IMMUNO-ONCOLOGY CONFERENCE

Berlin, Germany, October 26, 2018, 08.00 a.m. CEST – NOXXON Pharma N.V. (Euronext Growth Paris: ALNOX), a biotechnology company focused on improving cancer treatments by targeting the tumor microenvironment (TME), announced today that it will report top-line efficacy data from the ongoing clinical trial (NCT03168139) testing NOX-A12 (olaptesed pegol) in combination with Merck & Co./MSD's PD-1 inhibitor Keytruda® in metastatic, microsatellite stable pancreatic and colorectal cancer patients at the ESMO Immuno-Oncology Conference taking place in Geneva, Switzerland, December 13-16, 2018.

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About NOXXON

NOXXON's oncology-focused pipeline acts on the tumor microenvironment (TME) and the cancer immunity cycle by breaking the tumor protection barrier, blocking tumor repair and exposing hidden tumor cells. Through neutralizing chemokines in the tumor microenvironment, NOXXON's approach works in combination with other forms of treatment to weaken tumor defenses against the immune system and enable greater therapeutic impact. Building on extensive clinical experience and safety data, the lead program NOX-A12 will deliver top-line data from a Keytruda® combination trial in metastatic colorectal and pancreatic cancer patients in 2018. The company plans to initiate further studies with NOX-A12 in brain cancer in combination with radiotherapy, for which an orphan drug status has been granted in the US and EU. The company's second asset, NOX-E36 is a Phase 2 TME asset targeting the innate immune system. NOXXON plans to test NOX-E36 in pancreatic cancer patients both as a monotherapy and in combination. Further information can be found at: www.noxxon.com

Keytruda® is a registered trademark of Merck Sharp & Dohme Corp.

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