

## **Oxurion Announces Upcoming Preclinical Presentation on THR-149 at KININ2022**

**Leuven, BELGIUM, Boston, MA, US – MAY 24, 2022** – [Oxurion NV](#) (Euronext Brussels: **OXUR**), a biopharmaceutical company developing next generation standard of care ophthalmic therapies, with clinical stage assets in vascular retinal disorders, today announced that an abstract evaluating the pharmacokinetic properties of THR-149 will be presented at KININ2022 to be held June 6-8, 2022 in Annecy, France.

THR-149 is a potent plasma kallikrein inhibitor being developed as a potential new standard of care for the up to 50% of diabetic macular edema (DME) patients showing suboptimal response to anti-VEGF therapy. Oxurion is studying THR-149 in Part B of its two-part Phase 2 KALAHARI trial evaluating THR-149 for the treatment of DME.

### **Details of the presentation:**

**TITLE:** Pharmacokinetics of THR-149, a potent and specific plasma kallikrein bicyclic peptide inhibitor

**PRESENTER:** Marc Vanhove, Head, Biochemistry & in vitro Pharmacology, Oxurion NV

**PRESENTATION DATE AND TIME:** Tuesday, June 7, 2022, 2.30 p.m. CET

### **About Oxurion**

Oxurion (Euronext Brussels: OXUR) is a biopharmaceutical company developing next generation standard of care ophthalmic therapies, which are designed to improve and better preserve vision in patients with retinal disorders including diabetic macular edema (DME), the leading cause of vision loss in working-age people, as well as other conditions. Oxurion intends to play an important role in the treatment of retinal disorders, including the successful development of THR-149, its novel therapeutic for the treatment of DME. THR-149 is a potent plasma kallikrein inhibitor being developed as a potential new standard of care for the up to 50% of DME patients showing suboptimal response to anti-VEGF therapy. Oxurion is headquartered in Leuven, Belgium, with corporate operations in Boston, MA. More information is available at [www.oxurion.com](http://www.oxurion.com).

**For more information, please contact:**

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