Press Release

Nicox Announces Presentations of Additional NCX 470 Data at the Upcoming World Glaucoma Congress

March 21, 2023 – release at 7:30 am CET
Sophia Antipolis, France

Nicox SA (Euronext Paris: FR0013018124, COX), an international ophthalmology company, today announced presentations of additional NCX 470 data at the 10th World Glaucoma Congress (WGC) which will be held from June 28 to July 1st, 2023 in Rome, Italy.

Details of the presentations:

Poster Title: Effects of NCX 470, a Nitric Oxide (NO)-Donating Bimatoprost, in *in vitro* 3D-Human Trabecular Meshwork (TM) / Schlemm's Canal (SC) Co-Culture Tissue Model
Type: Poster Walk Presentation
Presenter: Corinna Galli, PhD, Nicox Research Institute, Italy

Poster Title: NCX 470, a Nitric Oxide Donating Bimatoprost versus Latanoprost has Greater Proportion of Subjects Achieving ≥10 mmHg IOP Decrease in Phase 3 Trial
Type: Poster presentation
Presenter: Robert Fechtner, MD, Chair of the Department of Ophthalmology at SUNY Upstate Medical University in Syracuse, NY, U.S.

Poster Title: NCX 470, a Nitric Oxide Donating Bimatoprost Compared with Latanoprost - Adaptive Design Period Results from the Phase 3 Mont Blanc Clinical Trial
Type: Poster Walk Presentation
Presenter: Steven Mansberger, MD, Vice-Chair, Director of Glaucoma Services and Ophthalmic Clinical Trials for the Devers Eye Institute in Portland, OR, U.S.

NCX 470 is a novel nitric oxide (NO)-donating bimatoprost currently in Phase 3 clinical development for the reduction of intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension. The first Phase 3 trial Mont Blanc, a randomized, double-masked, multi-center, parallel group trial conducted in the U.S., comparing NCX 470 (0.1%) to latanoprost (0.005%) was completed in October 2022. The second Phase 3 trial, Denali, similarly designed to the Mont Blanc trial, and which includes a long-term safety extension, is ongoing.

About Nicox

Nicox SA is an international ophthalmology company developing innovative solutions to help maintain vision and improve ocular health. Nicox's lead program in clinical development is NCX 470, a novel nitric oxide-donating bimatoprost, for lowering intraocular pressure in patients with open-angle glaucoma or ocular hypertension. The company is also conducting research on NCX 1728, a nitric oxide-donating phosphodiesterase 5 inhibitor, in retinal conditions. NCX 4251, a novel, patented, ophthalmic suspension of fluticasone propionate nanocrystals for topical ocular application for dry eye disease, is being developed by Ocumension Therapeutics in China under an exclusive license agreement and is available for partnering elsewhere. Nicox generates revenue from VYZULTA® in glaucoma, licensed exclusively worldwide to Bausch + Lomb, and ZERVIATE® in allergic conjunctivitis, licensed in multiple geographies, including to Eyevance Pharmaceuticals, LLC (a wholly-owned subsidiary of Santen Pharmaceutical Co., Ltd.), in the U.S. and Ocumension Therapeutics in the Chinese and in the majority of Southeast Asian markets.

Nicox is headquartered in Sophia Antipolis, France, is listed on Euronext Paris (Compartment C: Mid Caps; Ticker symbol: COX) and is part of the CAC Healthcare, CAC Pharma & Bio and Next 150 indexes.

For more information on Nicox, its products or pipeline, please visit: www.nicox.com.

Analyst coverage

Bryan, Garnier & Co	Eric Yoo	Paris, France
Edison Investment Research	Pooya Hemami	London, UK
H.C. Wainwright & Co	Yi Chen	New York, U.S.
Kepler Cheuvreux	Arsene Guekam	Paris, France

The views expressed by analysts in their coverage of Nicox are those of the author and do not reflect the views of Nicox. Additionally, the information contained in their reports may not be correct or current. Nicox disavows any obligation to correct or to update the information contained in analyst reports.

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Forward-Looking Statements

The information contained in this document may be modified without prior notice. This information includes forward-looking statements. Such forward-looking statements are not guarantees of future performance. These statements are based on current expectations or beliefs of the management of Nicox S.A. and are subject to a number of factors and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. Nicox S.A. and its affiliates, directors, officers, employees, advisers or agents, do not undertake, nor do they have any obligation, to provide updates or to revise any forward-looking statements.

Risks factors which are likely to have a material effect on Nicox's business are presented in the 3rd chapter of the '*Document d'enregistrement universel, rapport financier annuel et rapport de gestion 2021*' filed with the French *Autorité des Marchés Financiers* (AMF) on April 29, 2022 whose first amendment has been filed with the AMF on May 19, 2022, in the 2nd chapter of the second amendment filed with the AMF on November 22, 2022 and in the 2nd chapter of the Securities noted filed with the AMF on November 22, 2022 which are available on Nicox's website (www.nicox.com)

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