

## Hyloris welcomed for Exclusive Company Visit

Liège, Belgium – 20 May 2026 – 03:30 PM CET – Non Regulated Information - Hyloris Pharmaceuticals SA (“Hyloris”) (Euronext Brussels: HYL), the specialty pharma group committed to addressing unmet medical needs through reinventing existing medicines and smart NCE developments, today hosted an exclusive company visit for members of VFB (Vlaamse Federatie van Beleggers) at its headquarters on the LégiaPark pharma campus in Liège, together with neighbouring listed company Nyxoah.

The afternoon programme brought together up to 40 visitors and invited press for a comprehensive insight into both companies. Attendees were welcomed by CEO Stijn Van Rompay, who presented Hyloris' strategy, broad pipeline of 30 innovative drug candidates and its 2025 highlights - including 6 new products added to the portfolio and 8 out-licensing deals concluded across key global markets.

A separate presentation by Olivier Taelman, CEO of Nyxoah, followed, covering that company's advances in obstructive sleep apnea treatment through neuromodulation, including its newly announced cleanroom production unit at LégiaPark.

Guests were then invited for a guided tour of the Hyloris R&D laboratory, featuring the Formulation Lab, Analytical Lab and Stability Chambers - the facilities where Hyloris builds, characterizes and stress-tests its drug candidates.

LégiaPark, located at Boulevard de Patience et Beaujonc 31, 4000 Liège, serves as a unique life sciences innovation ecosystem in Wallonia, hosting two Belgian-listed companies on a single campus.

### About Hyloris Pharmaceuticals

Hyloris is a specialty pharma group focused on reinventing and optimizing existing medications through reformulation and repurposing to address important healthcare needs and deliver meaningful improvements for patients, healthcare professionals, and payers. The Company's development strategy primarily focuses on leveraging established regulatory pathways, such as the FDA's 505(b)(2) pathway in the U.S., or equivalent regulatory frameworks in other regions, which are specifically designed for pharmaceuticals for which safety and efficacy of the molecule have already been established. This approach can reduce the clinical and regulatory burden required for market entry, and significantly shorten development timelines, leading to reduced costs and risks. Hyloris has announced a broad development portfolio of 28 products, including 25 value-added medicines of which two are currently in early stages of commercialization in collaboration with commercial partners: Sotalol IV for the treatment of atrial fibrillation, and Maxigesic® IV, a non-opioid post-operative pain treatment. In addition to its core strategic focus, the Company has two high-barrier generic products approved in the U.S. and one additional high-barrier generic product in development. Beyond its announced portfolio, Hyloris has initiated several additional internal early-stage development activities, bringing the total pipeline to more than 30 products and product candidates, and continues to evaluate further product opportunities to support future growth. **Where others stop, Hyloris starts and adds value through Reinventing Existing Medicines for unmet patient needs worldwide.** Hyloris is based in Liège, Belgium and since 2020 listed on Euronext Brussels (EBR: HYL) with about 55 people across 13 nationalities, targeting time-to-market of 7 years at an average cost of EUR 7 million per product.

For more information, visit [www.hyloris.com](http://www.hyloris.com) and follow-us on [LinkedIn](#).

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**Disclaimer and forward-looking statements**

Hyloris means “high yield, lower risk”, which relates to the 505(b)(2) regulatory pathway for product approval on which the Company focuses, but in no way relates or applies to an investment in the Shares. Certain statements in this press release are “forward-looking statements.” These forward-looking statements can be identified using forward-looking terminology, including the words “believes”, “estimates”, “anticipates”, “expects”, “intends”, “may”, “will”, “plans”, “continue”, “ongoing”, “potential”, “predict”, “project”, “target”, “seek” or “should”, and include statements the Company makes concerning the intended results of its strategy. These statements relate to future events or the Company’s future financial performance and involve known and unknown risks, uncertainties, and other factors, many of which are beyond the Company’s control, that may cause the actual results, levels of activity, performance or achievements of the Company or its industry to be materially different from those expressed or implied by any forward-looking statements. The Company undertakes no obligation to publicly update or revise forward-looking statements, except as may be required by law. The Company undertakes no obligation to publicly update or revise forward-looking statements, except as may be required by law.