Nyxoah Joins the Euronext Tech Leaders Initiative, Included in the Euronext Tech Leaders Index

Mont-Saint-Guibert, Belgium – June 7, 2022 4:30pm ET / 10:30pm CET– Nyxoah SA (Euronext Brussels/Nasdaq: NYXH) ("Nyxoah" or the "Company"), a medical technology company focused on the development and commercialization of innovative solutions to treat Obstructive Sleep Apnea (OSA), is proud to announce that it is part of the newly-formed Euronext Tech Leaders initiative, which is composed of 100+ innovative and high-growth technology companies with greater than 1 trillion Euros in aggregate market capitalization.

Companies participating in the Euronext Tech Leaders initiative will be included in the Euronext Tech Leaders Index and benefit from a suite of exclusive programs, such as dedicated Euronext programs targeting improved trading conditions for retail investors, greater international visibility through marketing and communication initiatives, and access to the C-level club offering exclusive networking events.

"We are proud that our patient-first mission has led to Nyxoah being one of 100 Tech companies to be recognized for innovation and growth through inclusion in the Euronext Tech Leaders initiative and index," commented Olivier Taelman, Nyxoah's Chief Executive Officer.

About Nyxoah

Nyxoah is a medical technology company focused on the development and commercialization of innovative solutions to treat Obstructive Sleep Apnea (OSA). Nyxoah's lead solution is the Genio® system, a patient-centered, leadless and battery-free hypoglossal neurostimulation therapy for OSA, the world's most common sleep disordered breathing condition that is associated with increased mortality risk and cardiovascular comorbidities. Nyxoah is driven by the vision that OSA patients should enjoy restful nights and feel enabled to live their life to its fullest.

Following the successful completion of the BLAST OSA study, the Genio® system received its European CE Mark in 2019. Nyxoah completed two successful IPOs: on Euronext Brussels in September 2020 and NASDAQ in July 2021. Following the positive outcomes of the BETTER SLEEP study, Nyxoah received CE mark approval for the expansion of its therapeutic indications to Complete Concentric Collapse (CCC) patients, currently contraindicated in competitors' therapy. Additionally, the Company is currently conducting the DREAM IDE pivotal study for FDA and US commercialization approval.

For more information, please visit http://www.nyxoah.com/.

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