Press release

Sensorion to participate in the LifeSci Partners 2nd Annual Genetic Medicines Symposium on June 29, 2022

Montpellier, June 24, 2022 – Sensorion (FR0012596468 – ALSEN) a pioneering clinical-stage biotechnology company which specializes in the development of novel therapies to restore, treat and prevent within the field of hearing loss disorders, is pleased to announce the participation of CEO Nawal Ouzren in the LifeSci Partners 2nd Annual Genetic Medicines Symposium being held virtually on June 28-29, 2022.

CEO Nawal Ouzren will participate in a fireside chat with Jeff Goodenbour as part of the **LifeSci Partners 2nd Annual Genetic Medicines Symposium**, on June 29, 2022. To attend the discussion, please register here. A replay of the fireside chat will be available shortly after the presentation is over on the same link.

Date: Wednesday, June 29, 2022
Fireside Chat: 8.30 am – 9 am EST (2.30 pm – 3 pm CEST)
Event: LifeSci Partners 2nd Annual Genetic Medicines Symposium

About Sensorion

Sensorion is a pioneering clinical-stage biotech company, which specializes in the development of novel therapies to restore, treat and prevent hearing loss disorders, a significant global unmet medical need.
Sensorion has built a unique R&D technology platform to expand its understanding of the pathophysiology and etiology of inner ear related diseases, enabling it to select the best targets and modalities for drug candidates. Its portfolio combines both small molecule programs and a preclinical portfolio of inner ear gene therapies.
Its clinical-stage portfolio includes one Phase 2 product: SENS-401 (Arazasetron) progressing in a planned Phase 2 Proof of Concept clinical study of SENS-401 in Cisplatin-Induced Ototoxicity (CIO) and, with partner Cochlear Limited, a study of SENS-401 in patients scheduled for cochlear implantation.
Sensorion has entered into a broad strategic collaboration with Institut Pasteur focused on the genetics of hearing. It has two gene therapy programs aimed at correcting hereditary monogenic forms of deafness including deafness caused by a mutation of the gene encoding for Otoferlin, and hearing loss related to mutation in GJB2 gene to potentially address important hearing loss segments in adults and children. The Company is also working on the identification of biomarkers to improve diagnosis of these underserved illnesses.
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Press release

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Mnemonic: **ALSEN**

Disclaimer

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