Press release

Sensorion to present at the Gene Therapy Development and Manufacturing conference in June 2022

Montpellier, June 9, 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 in-person of a member of Sensorion's senior management in the Gene Therapy Development and Manufacturing conference being held in London, UK, in June 2022.

Otmane Boussif, Sensorion's Chief Technical Officer, will give a presentation on "**Gene Therapy Approach For Hearing Disorders: AAV Manufacturing & Control Strategy**" on June 13, 2022, at the **Gene Therapy Development and Manufacturing** conference taking place in London, UK, June 13-14, 2022.

Date: Monday, June 13, 2022
Presentation: 4.50 am – 5.15 am EDT (9.50 am – 10.15 UK time)
Event: Gene Therapy Development and Manufacturing

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 study of SENS-401 clinical study 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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Disclaimer

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