

Sensorion announces its financial calendar for the first half of 2023

Montpellier, February 20, 2023 – 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, today announces its financial calendar for the first half of 2023.

Event	Date*
2022 Full-Year Financial Results	March 16, 2023
General Assembly	May 24, 2023

In addition to these periodic publications, investors are invited to consult the Company's website (www.sensorion-pharma.com), where information is regularly updated. All corporate and financial information on the Company is available in the "Investor" section of the website.

*This tentative agenda is subject to change. Publications will take place before Paris market open.

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 mechanisms of action for drug candidates. Its portfolio combines both small molecule and inner ear gene therapies programs.

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, in a study of SENS-401 in patients scheduled for cochlear implantation. A Phase 2 study of SENS-401 was also completed in Sudden Sensorineural Hearing Loss (SSNHL) in January 2022.

Sensorion pursues its 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 OTOF-GT, targeting deafness caused by mutations of the gene encoding for otoferlin, and hearing loss related to mutations in *GJB2* gene to potentially address important hearing loss segments in adults and children (GJB2-GT). The Company is also working on the identification of biomarkers to improve diagnosis of these underserved illnesses.

www.sensorion.com

Contacts

Investor Relations

Noemie Djokovic
Investor Relations and Communications
+33 6 76 67 98 31
ir.contact@sensorion-pharma.com

International Media Relations

Consilium Strategic Communications
Matthew Cole/Jessica Hodgson
+44 7593 572720
+44 7561 424788
Sensorion@consilium-comms.com

Press release

Label: **SENSORION**
ISIN: **FR0012596468**
Mnemonic: **ALSEN**



Disclaimer

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