Press release

Sensorion announces attendance and presentations at upcoming conferences

Montpellier, September 1, 2020 - 7:30AM CEST - 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, announces its attendance and presentations at a number of upcoming conferences.

- September 9: Fireside chat in the SunTrust Robinson Humphrey Virtual Meeting Series – Sensorion will present at 10:00am ET
- September 8-11: Advanced Therapies (Virtual) - Christine Le Bec, Head of CMC Gene Therapy will present "Challenges and Issues in Dual AAV Vectors Approach" on September 8th in the Viral Vector Manufacturing track at 11:30am BST
- October 1-2: Jefferies Gene Therapy/Editing Summit (Virtual)
- October 5-6: Chardan 4th Annual Genetic Medicines Conference (Virtual) - Sensorion's presentation time will be announced closer to the date
- October 5-6: Healthtech Innovation Days (In-person Paris and Virtual)
- October 12-16: Cell and Gene Therapy Meeting on the Mesa (Virtual) - Sensorion's presentation date and time will be announced closer to the date

About Sensorion

Sensorion is a pioneering clinical-stage biotech company, which specializes in the development of novel therapies to restore, treat and prevent within the field of hearing loss disorders. Its clinical-stage portfolio includes one Phase 2 product: SENS-401 (Arazasetron) for sudden sensorineural hearing loss (SSNHL). 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. The Company is also working on the identification of biomarkers to improve diagnosis of these underserved illnesses. In the second half of 2019, Sensorion launched two preclinical gene therapy programs aiming at correcting hereditary monogenic forms of deafness including Usher Type 1 and deafness caused by a mutation of the gene encoding for Otoferlin. The Company is uniquely placed, through its platforms and pipeline of potential therapeutics, to make a lasting positive impact on hundreds of thousands of people with inner ear related disorders, a significant global unmet medical need.
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Market Authority) on September 7th, 2017 under n°R.17-062 and to the development of economic conditions, financial markets and the markets in which Sensorion operates. The forward-looking statements contained in this press release are also subject to risks not yet known to Sensorion or not currently considered material by Sensorion. The occurrence of all or part of such risks could cause actual results, financial conditions, performance or achievements of Sensorion to be materially different from such forward-looking statements.

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