

# Idiopathic Sudden Sensorineural Hearing Loss

A WINDOW OF DAYS TO WEEKS — in treated patients the odds of complete recovery fall about 12% per day

An adjunct to corticosteroids in acute idiopathic sudden hearing loss, given early, to improve audiometric recovery.

| TREATMENT PROTOCOL |                                                                                                                                             |
|--------------------|---------------------------------------------------------------------------------------------------------------------------------------------|
| Pressure           | 2.0–2.5 ATA                                                                                                                                 |
| Duration           | 90 min                                                                                                                                      |
| Frequency          | Daily                                                                                                                                       |
| Sessions           | 10–20 sessions. Benefit is greatest with a total exposure of 1,200 min or more                                                              |
| Review             | UHMS sets no review point; continue ENT follow-up. Bagli 2020 model: audiometry after 10, continue to 20 only if PTA improved 11 dB or more |

| PATIENT SELECTION                 |                                                                                                                                                                                                                                                                                                                             |
|-----------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Who to treat                      | Idiopathic acute sudden hearing loss, treated as early as possible alongside steroids. Benefit was greatest in severe-to-profound loss and in salvage after steroids had failed in Rhee 2018 (mostly non-randomised); Han 2026 found no overall effect in profound loss. UHMS: consider for loss over 40 dB within 14 days. |
| Who not to treat, or when to stop | Chronic loss or isolated tinnitus — Cochrane found no benefit. One trial found the tinnitus response tracked patient expectation ( $p<0.05$ ), which the authors attributed largely to psychological mechanisms.                                                                                                            |
| Mechanism                         | The cochlea has a high metabolic demand and poor collateral supply. Raising perilymph oxygen tension can recover hair-cell function that is ischaemic but still viable, if applied early.                                                                                                                                   |
| Outcome measure                   | Pure-tone audiometry gain in dB; 25% and 50% threshold-improvement criteria; Siegel's criteria; word discrimination score.                                                                                                                                                                                                  |

| WHAT THE NUMBERS SHOW                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |  |
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| <ul style="list-style-type: none"><li>Newth 2025 Cochrane-lineage update (7 trials, <math>n=562</math>, GRADE moderate by author judgement; AMSTAR-2 critically low): <b>RR 1.5 (95% CI 1.2–1.7)</b>, <b>NNT 4</b>, and <b>15.1 dB (95% CI 8.2–22.0)</b> of additional gain.</li><li>Bagli 2020 (<math>n=96</math>, all HBOT-treated, no comparator): the onset-to-treatment interval was <b>the only independent predictor</b> of complete recovery — <b>OR 0.88 per day of delay (95% CI 0.82–0.95, <math>p=0.002</math>)</b>, about 12% of the odds lost per day.</li><li>Lun 2026 (<math>n=70</math>, four timing groups): median gain <b>33.8 dB early, 6.9 dB late, 0.0 dB salvage, 11.9 dB with steroids but no HBOT</b>.</li><li>Alde 2023 (<math>n=49</math>, treated within 3 days, no steroids): complete recovery <b>71.4% (35/49)</b>, uncontrolled.</li><li>Chaushu 2023: pooled <b>spontaneous</b> recovery is about <b>60.3%</b> — the baseline every figure above must be read against.</li></ul> |  |

| EVIDENCE BASE                                     |                                                                                                                             |                                                                 |
|---------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| Shape and certainty                               | Very large — 40 or more records, at least 8 randomised trials, multiple Cochrane-lineage reviews. Certainty GRADE moderate. |                                                                 |
| Study                                             | Design                                                                                                                      | Result and risk of bias                                         |
| Newth 2025, Diving Hyperb Med — PMID 41364864     | Cochrane-lineage update, 7 trials, $n=562$                                                                                  | <b>Positive</b> ; GRADE moderate; AMSTAR-2 critically low       |
| Bennett 2012, Cochrane — PMID 23076907            | SR, 7 RCTs, $n=392$                                                                                                         | Positive acute; no benefit chronic                              |
| Rhee 2018, JAMA Otolaryngol — PMID 30267033       | Meta-analysis, 19 studies, $n=2,401$                                                                                        | Positive                                                        |
| Xie 2026, Braz J Otorhinolaryngol — PMID 42143851 | Propensity-matched cohort, $n=270$ matched                                                                                  | <b>No benefit</b> ; estimates favour control; ROBINS-I moderate |
| Lun 2026, Diving Hyperb Med — PMID 41875440       | Retrospective cohort by timing, $n=70$                                                                                      | Positive for early; ROBINS-I serious                            |

| WHAT THE EVIDENCE DOES NOT SUPPORT |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           |
|------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Cautions                           | <b>There is no 48-hour threshold in this literature.</b> The only trial enrolling inside 48 h (Fattori 2001) used that as an entry criterion, not a tested cut-off, and its comparator was a vasodilator rather than steroids. <b>Xie 2026, the best-controlled study, found no added benefit when full intravenous and intratympanic steroids are already given early.</b> Lun's timing gradient is confounded — its own steroid-only (no-HBOT) group improved more (11.9 dB) than its late-treated group (6.9 dB), which is internal evidence the groups are not exchangeable. Absolute effects are modest against a 60% spontaneous-recovery baseline. |

| STATUS AND SOCIETY POSITIONS |                                                                                                                                                                               |
|------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Status                       | <b>Not a PMB condition</b> — H91.2 is not on the CMS coded list. Relies on the clinical and SAUHMA Level B argument plus NRPL recognition, within the early-treatment window. |
| ICD-10 (SA)                  | H91.2, H90.3, H90.4, H90.5, H91.9                                                                                                                                             |
| ECHM 2017 (Type, Level)      | 1B with medical therapy within 2 weeks; not after 6 months                                                                                                                    |
| AAO-HNSF 2019                | Option, with steroids: within 2 weeks of onset, or as salvage within 1 month                                                                                                  |
| NICE NG98                    | Does not address HBO <sub>2</sub> (WHO guideline review, 2025)                                                                                                                |

Compiled from the UHMS Hyperbaric Oxygen Therapy Indications Manual, 15th edition (2023) and the appraised trial evidence held in an HBOT evidence library. All treatments are 100% oxygen. US Navy Treatment Table profiles are named, not reproduced — the US Navy Diving Manual is the operative document at the chamber. Session counts are largely guideline convention rather than dose-finding evidence. This card summarises published parameters and does not replace those source documents or clinical judgement for the individual patient. Generated 28 October 2026. Not a SAUHMA publication and carries no society endorsement.