AM1154
LB332
MMM - 04/23/2025

AMENDMENTS TO LB332

(Amendments to E&R amendments, ER42)

Introduced by Hardin, 48.

1. Strike sections 1, 11 to 13, 15 to 17, 19 to 29, 32, 34, 40, and 41 and insert the following new sections:

Section 1. Section 38-511, Reissue Revised Statutes of Nebraska, is amended to read:

38-511 Nothing in the Audiology and Speech-Language Pathology Practice Act shall be construed to prevent or restrict:

(1) The practice of audiology or speech-language pathology or the use of the official title of such practice by a person employed as a speech-language pathologist or audiologist by the federal government;

(2) A physician from engaging in the practice of medicine and surgery or any individual from carrying out any properly delegated responsibilities within the normal practice of medicine and surgery under the supervision of a physician;

(3) A person licensed as a hearing instrument specialist in this state from engaging in the fitting, selling, ordering, and servicing of hearing instruments or performing such other duties as defined in the Hearing Instrument Specialists Practice Act;

(4) The practice of audiology or speech-language pathology or the use of the official title of such practice by a person who holds a valid and current credential as a speech-language pathologist or audiologist issued by the State Department of Education, if such person performs speech-language pathology or audiology services solely as a part of his or her duties within an agency, institution, or organization for which no fee is paid directly or indirectly by the recipient of such service and under the jurisdiction of the State Department of Education, but such person may elect to be within the jurisdiction of the Audiology and

AM1154
LB332
MMM - 04/23/2025

Speech-Language Pathology Practice Act;

(5) The clinical practice in audiology or speech-language pathology required for students enrolled in an accredited college or university pursuing a major in audiology or speech-language pathology, if such clinical practices are supervised by a person licensed to practice audiology or speech-language pathology and if the student is designated by a title such as student clinician or other title clearly indicating the training status; or

(6) The utilization of a speech aide or other personnel employed by a public school, educational service unit, or other private or public educational institution working under the direct supervision of a licensed ~~credentialed~~ speech-language pathologist.

Sec. 11. Section 38-1501, Reissue Revised Statutes of Nebraska, is amended to read:

38-1501 Sections 38-1501 to 38-1518 and sections 14 and 23 to 32 of this act shall be known and may be cited as the Hearing Instrument Specialists Practice Act.

Sec. 12. Section 38-1502, Reissue Revised Statutes of Nebraska, is amended to read:

38-1502 For purposes of the Hearing Instrument Specialists Practice Act and elsewhere in the Uniform Credentialing Act, unless the context otherwise requires, the definitions found in sections 38-1503 to 38-1507 and section 14 of this act apply.

Sec. 13. Section 38-1504, Reissue Revised Statutes of Nebraska, is amended to read:

38-1504 Hearing instrument means any wearable instrument or device designed for, ~~or~~ offered for the purpose of, or represented as aiding persons with or compensating for impaired ~~human~~ hearing ~~and any parts, attachments, or accessories, including earmold, but excluding batteries and cords~~. Hearing instrument does not include a wearable instrument or device with an implantable component such as a wearable processor for a

AM1154
LB332
MMM - 04/23/2025

cochlear implant or bone-anchored implant.

Sec. 15. Section 38-1505, Reissue Revised Statutes of Nebraska, is amended to read:

38-1505 (1) Practice of ordering the use and fitting of hearing instruments includes the following activities:

(a) Eliciting patient case histories, including medical history, otological history, pharmacological history, amplification history, and patient attitudes and expectations;

(b) Administering otoscopy and, if required, cerumen removal for the purpose of identifying possible otological conditions, including, but not limited to, any of the conditions related to warnings found in the regulations of the federal Food and Drug Administration, 21 C.F.R. 801.422, as such regulations existed on January 1, 2025, which may indicate the need for a medical referral or which may have a bearing on outcomes or recommendations;

(c) Administering and interpreting tests of human hearing performed with an audiometer, including other appropriate objective and subjective methodology and measures, for purposes of ordering and fitting hearing aids;

(d) Determining candidacy for hearing instruments, and discussing the results of a human hearing test with the individual to inform the individual about potential options for addressing the individual's hearing loss, including hearing instruments, hearing-assistive devices, or other medical interventions, and facilitating appropriate referrals, if needed;

(e) Ordering, selecting, or fitting appropriate hearing instruments and assistive devices, including appropriate technology, programming parameters, and special custom earpiece applications, as indicated;

(f) Assessing hearing instrument efficacy utilizing appropriate fitting verification methodology and equipment, which may include real-ear measures or speech mapping, and electroacoustic analysis equipment;

AM1154
LB332
MMM - 04/23/2025

(g) Assessing hearing instrument benefits through appropriate validation measures, which may include communication assessment questionnaires or speech audiometry;

(h)(i) Taking ear impressions or electronic scans by any method used for the purpose of creating earmolds and (ii) preparing earmolds for hearing instruments, assistive devices, telecommunications applications, ear protection, and other related applications;

(i) Ordering and modifying earmolds and auditory equipment, excluding FM transmitters, to meet a patient's needs;

(j) Providing services in the use and care of hearing instruments and assistive devices, including listening strategies and other approaches to foster optimal patient results;

(k) Providing supervision and inservice training of those entering the dispensing profession;

(l) Providing post-fitting care and services and hearing instrument care and repair services; or

(m) Any other act of hearing assessment pertaining to hearing testing, ordering the use of hearing instruments, or the selling, renting, leasing, and delivery of hearing instruments.

(2) Practice of ordering the use and fitting of hearing instruments does not include:

(a) Evaluation, diagnosis, management, or treatment of auditory or vestibular conditions;

(b) Provision of tinnitus evaluation, treatment, or management;

(c) Interpretation of tests of human hearing for any purpose beyond the selection and fitting of hearing aids;

(d) Removal of foreign bodies from the ear; and

(e) Testing and treatment of auditory processing disorders, including the provision of aural rehabilitation or auditory training.

~~Practice of fitting hearing instruments means the measurement of human hearing by means of an audiometer or by other means approved by the~~

AM1154
LB332
MMM - 04/23/2025

~~board solely for the purpose of making selections, adaptations, or sale of hearing instruments. The term also includes the making of impressions for earmolds. A dispenser, at the request of a physician or a member of related professions, may make audiograms for the professional's use in consultation with the hard-of-hearing.~~

Sec. 17. Section 38-1509, Revised Statutes Cumulative Supplement, 2024, is amended to read:

38-1509 (1)(a) ~~(1)~~ Except as otherwise provided in this section, it shall be unlawful for any person to engage in the practice of ordering the use and fitting of ~~no person shall engage in the sale of or practice of fitting~~ hearing instruments or display a sign or in any other way advertise or represent that the person is engaged in the practice of ordering the use and fitting ~~himself or herself as a person who practices the fitting and sale or dispensing~~ of hearing instruments unless such person ~~he or she~~ holds a current, ~~an~~ unsuspended, and unrevoked hearing instrument specialist license issued by the department as provided in the Hearing Instrument Specialists Practice Act.

(b) A hearing instrument specialist license shall confer upon the holder the right to select, fit, and sell hearing instruments. ~~A person holding a license issued under the act prior to August 30, 2009, may continue to practice under such license until it expires under the terms of the license.~~

(2) A licensed audiologist ~~who maintains a practice pursuant to (a) licensure as an audiologist, or (b) a privilege to practice audiology under the Audiology and Speech-Language Pathology Interstate Compact, in which hearing instruments are regularly dispensed, or who intends to maintain such a practice,~~ shall be exempt from the requirement to be licensed as a hearing instrument specialist.

(3) A hearing instrument specialist or audiologist may order the use of devices pursuant to 21 C.F.R. 801.109, as such regulation existed on January 1, 2025.

AM1154
LB332
MMM - 04/23/2025

(4)(a) ~~(3)~~ Nothing in the Hearing Instrument Specialists Practice Act ~~act~~ shall prohibit a corporation, partnership, limited liability company, trust, association, or other like organization maintaining an established business address from engaging in the business of selling or offering for sale hearing instruments at retail without a license if it employs only properly licensed natural persons in the direct sale and fitting of such products.

(b) Each such organization shall file annually with the department, on a form provided by the department, a list of the licensed hearing instrument specialists employed by the organization. The department may adopt and promulgate rules and regulations as necessary to carry out this section.

~~(4) Nothing in the act shall prohibit the holder of a hearing instrument specialist license from the fitting and sale of wearable instruments or devices designed for or offered for the purpose of conservation or protection of hearing.~~

Sec. 18. Section 38-1510, Reissue Revised Statutes of Nebraska, is amended to read:

38-1510 (1) A licensed hearing instrument specialist shall only provide services to an individual who is eighteen years of age or older unless prohibited by federal law.

(2) The Hearing Instrument Specialists Practice Act does not change the scope of practice of a licensed audiologist.

(3) ~~(1)~~ The Hearing Instrument Specialists Practice Act is not intended to prevent any person from engaging in the practice of measuring human hearing for the purpose of selection of hearing instruments if such person or organization employing such person does not sell hearing instruments or the accessories thereto.

(4) ~~(2)~~ The Hearing Instrument Specialists Practice Act does ~~act shall~~ not apply to a person who is a physician or audiologist licensed to practice in this state, except that such physician or audiologist shall

AM1154
LB332
MMM - 04/23/2025

not delegate the authority to fit and dispense hearing instruments unless the person to whom the authority is delegated is licensed as a hearing instrument specialist under the act.

Sec. 19. Section 38-1511, Reissue Revised Statutes of Nebraska, is amended to read:

38-1511 (1) A licensed hearing instrument specialist shall enter into a written contract for each sale of a hearing instrument which states the terms of the sale.

(2)(a) A licensed hearing instrument specialist shall, at the time of delivery of the hearing instrument, provide the patient with a receipt containing (i) the signature, regular business address, and license number of the licensee, (ii) the brand, model, manufacturer or manufacturer identification code, and serial number of the hearing instrument, and (iii) the amount charged for the hearing instrument.

(b) The receipt shall indicate that the hearing device is classified as programmed with one of the following:

(i) Locked software - this device utilizes locked software that is available to limited providers. The purchase of this device will require the user to have the device programmed by a provider or chain store that has been granted proprietary access to the software. In addition, the availability of any part or service for this device is limited to the provider or chain store that has such proprietary access; or

(ii) Unlocked software - this device utilizes unlocked software that is readily available to any provider or location licensed to provide hearing health care.

(c) The receipt shall also specify (i) whether the hearing instrument is new, used, or rebuilt, as provided in 21 C.F.R. 801.422, as such regulation existed on January 1, 2025, (ii) the length of time and other terms of the guarantee, and (iii) by whom the hearing instrument is guaranteed.

(3) No hearing instrument may be sold to any person unless both the

AM1154
LB332
MMM - 04/23/2025

packaging containing the hearing instrument and the itemized receipt are in compliance with all applicable state and federal laws and regulations.

(4) Upon delivery of the hearing instrument to any person, the licensed hearing instrument specialist shall confirm the physical and operational performance of the hearing instrument. If a patient purchases a hearing instrument from a licensed hearing instrument specialist outside of the licensee's regular place of business and the regular place of business is not within a reasonable distance, as determined by the board, the licensed hearing instrument specialist shall provide the patient with the address of an affiliate location with which the licensee is associated that is within a reasonable distance, at which a licensed hearing instrument specialist or audiologist is available for fitting services.

(5) Any seller offering for sale or selling a hearing instrument in this state or to a resident of this state shall make available in this state an in-person fitting of the hearing instrument by a licensed hearing instrument specialist in this state prior to the sale.

~~(1) Any person who practices the fitting and sale of hearing instruments shall deliver to each person supplied with a hearing instrument a receipt which shall contain the licensee's signature and show his or her business address and the number of his or her certificate, together with specifications as to the make and model of the hearing instrument furnished, and clearly stating the full terms of sale. If a hearing instrument which is not new is sold, the receipt and the container thereof shall be clearly marked as used or reconditioned, whichever is applicable, with terms of guarantee, if any.~~

(6) A receipt provided pursuant to this section ~~(2) Such receipt~~ shall bear in no smaller type than the largest used in the body copy portion the following: The purchaser has been advised at the outset of the ~~his or her~~ relationship with the hearing instrument specialist that any examination or representation made by a licensed hearing instrument

AM1154
LB332
MMM - 04/23/2025

specialist in connection with the fitting and selling of this hearing instrument is not an examination, diagnosis, or prescription by a person licensed to practice medicine in this state and therefor must not be regarded as medical opinion or advice.

Sec. 20. Section 38-1512, Revised Statutes Cumulative Supplement, 2024, is amended to read:

38-1512 (1) Any person may obtain a hearing instrument specialist license under the Hearing Instrument Specialists Practice Act by successfully passing a qualifying examination pursuant to section 38-1514 if the applicant provides verification to the department, on a form provided by the department, that such person:

(a) Is at least twenty-one years of age; ~~and~~

(b) Has an education equivalent to a four-year course in an accredited high school; and ~~.~~

(c)(i) Has completed the minimum number of practicum hours prescribed by the board;

(ii) Has a two-year degree in hearing instrument sciences or an equivalent as determined by the board;

(iii) Has held a current, unsuspended, and unrevoked license to dispense hearing instruments from another jurisdiction for at least twelve of the last eighteen months prior to taking the examination;

(iv) Is certified by the National Board for Certification in Hearing Instrument Sciences at the time of taking the examination; or

(v) Holds an advanced credential offered by the International Hearing Society at the time of taking the examination.

(2) The department, with the recommendation of the board, may determine whether a person who has completed a licensure program outside of the United States may take the examination.

(3) The department, upon recommendation of the board, may waive components of the examination pursuant to section 38-1514 for licensure as a hearing instrument specialist if the person has passed the same

AM1154
LB332
MMM - 04/23/2025

examination as provided in section 38-1514 or a substantially equivalent examination as determined by the board.

(4) The department, with the recommendation of the board, shall determine whether a person has met the requirements to be eligible to take the examination pursuant to the Hearing Instrument Specialists Practice Act.

~~(2) The qualifying examination shall consist of written and practical tests. The examination shall not be conducted in such a manner that college training is required in order to pass. Nothing in this examination shall imply that the applicant is required to possess the degree of medical competence normally expected of physicians.~~

~~(3) The department shall give examinations approved by the board. A minimum of two examinations shall be offered each calendar year.~~

Sec. 21. Section 38-1513, Revised Statutes Cumulative Supplement, 2024, is amended to read:

38-1513 (1) The department, with the recommendation of the board, shall issue a temporary training license to any person who has met the requirements for licensure as a hearing instrument specialist pursuant to subdivisions (1)(a) and (b) ~~subsection (1)~~ of section 38-1512. Previous experience or a waiting period shall not be required to obtain a temporary training license.

(2) Any person who desires a temporary training license shall make application to the department. The temporary training license shall be issued for a period of one year. A person holding a valid license as a hearing instrument specialist or an audiologist shall be responsible for the supervision and training of such applicant and shall maintain adequate personal contact with him or her.

(3) If a person who holds a temporary training license under this section has not successfully passed the licensing examination within twelve months of the date of issuance of the temporary training license, the temporary training license may be renewed or reissued for a twelve-

AM1154
LB332
MMM - 04/23/2025

month period. In no case may a temporary training license be renewed or reissued more than once. A renewal or reissuance may take place any time after the expiration of the first twelve-month period.

Sec. 22. Section 38-1514, Reissue Revised Statutes of Nebraska, is amended to read:

38-1514 (1) The examination required by section 38-1512 for licensure as a hearing instrument specialist shall be comprised of:

(a) A written or computer-based, psychometrically valid, competency examination approved by the board that tests the examinee for knowledge fundamental to the practice of ordering the use and fitting of hearing instruments;

(b) A practical examination approved by the board that requires the examinee to demonstrate competence in the practice of ordering the use and fitting of hearing instruments; and

(c) A jurisprudence examination approved by the board.

(2)(a) If an examinee fails more than one portion of the practical examination, the examinee shall retake the entire practical examination upon payment of the examination fee.

(b) If an examinee fails only one portion of the practical examination, the examinee may retake that portion of the examination without payment of a fee.

(c) If an examinee fails the jurisprudence examination or competency examination, the examinee shall retake the entire examination upon payment of the examination fee.

(d) If an examinee fails either the practical or competency component of the examination and fails two subsequent reexaminations, the examinee shall be disqualified from retaking the examination a fourth time until the examinee meets with the board, presents an acceptable written training plan to the board for passing the components of the examination, and successfully completes that plan.

(3) The qualifying examination provided in section 38-1512 shall be

AM1154
LB332
MMM - 04/23/2025

designed to demonstrate the applicant's adequate technical qualifications by:

(a) ~~(1)~~ Tests of knowledge in the following areas as they pertain to the practice of ordering the use and fitting ~~and sale~~ of hearing instruments:

(i) ~~(a)~~ Basic physics of sound;

(ii) ~~(b)~~ The anatomy and physiology of the ear; and

(iii) ~~(c)~~ The function of hearing instruments; and

(b) ~~(2)~~ Practical tests of proficiency in the following techniques as they pertain to the fitting of hearing instruments:

(i) ~~(a)~~ Pure tone audiometry, including air conduction testing and bone conduction testing;

(ii) ~~(b)~~ Live voice or recorded voice speech audiometry;

(iii) ~~(c)~~ Masking when indicated;

(iv) ~~(d)~~ Recording and evaluation of audiograms and speech audiometry to determine proper selection and adaptation of a hearing instrument; and

(v) ~~(e)~~ Taking earmold impressions.

Sec. 23. (1) A licensed hearing instrument specialist shall not engage in the practice of ordering the use and fitting of hearing instruments with respect to a patient without having conducted a face-to-face hearing assessment for the patient or having conducted or reviewed a valid and current hearing assessment for the patient that is dated within six months and signed by a licensed hearing instrument specialist or audiologist. Such hearing assessment shall include the following procedures, or modified procedures as required by the patient's cognitive function or health and appropriate to technological developments as determined by the board:

(a) Completion of a patient history questionnaire;

(b) Otoscopic examination;

(c) Testing to determine the type and degree of hearing loss that

AM1154
LB332
MMM - 04/23/2025

includes (i) pure-tone air conduction testing at two hundred fifty hertz, five hundred hertz, one thousand hertz, two thousand hertz, four thousand hertz, and eight thousand hertz, (ii) bone conduction testing at five hundred hertz, one thousand hertz, two thousand hertz, and four thousand hertz, and (iii) appropriate inter-octave testing when needed if the octave to adjacent octave threshold difference is greater than fifteen decibels;

(d) Effective masking when indicated;

(e) Appropriate testing to determine speech reception thresholds, word recognition scores, most comfortable listening levels, uncomfortable loudness levels, frequency-specific loudness discomfort levels, ability to understand speech in noise, and the selection of the best fitting arrangement for maximum hearing instrument benefit when indicated; and

(f) Other speech tests commonly used to assess human hearing acuity for ordering the use and fitting of hearing instruments.

(2) Each component of a hearing instrument shall be adapted to the needs of the patient. A licensed hearing instrument specialist shall conduct a final fitting to ensure physical fit and operational comfort of the hearing instrument.

Sec. 24. A licensed hearing instrument specialist shall demonstrate the benefit of a hearing instrument fitting by using objective measures.

Sec. 25. A licensed hearing instrument specialist shall determine a patient's benefit with the hearing instrument fitting using validation measures, such as speech audiometry and validated communication assessment questionnaires, or any other method approved by the board for ordering the use and fitting of hearing instruments.

Sec. 26. (1) A licensed hearing instrument specialist shall use the following equipment as part of any hearing testing conducted for the purpose of dispensing of hearing instruments:

(a) An audiometer that has been calibrated within the twelve months preceding the test and that meets the specifications set forth under this

AM1154
LB332
MMM - 04/23/2025

section; and

(b) A speech audiometer that has been calibrated within the twelve months preceding the test and that meets the specifications set forth under this section.

(2) A licensed hearing instrument specialist shall provide for the calibration of the equipment utilized for hearing assessments required under section 23 of this act and in the dispensing of hearing instruments at least annually in conformance with current standards of the American National Standards Institute for ordering the use and fitting of hearing instruments. A licensed hearing instrument specialist shall annually ensure that audiometric equipment has been evaluated electrically and acoustically, that the equipment has been adjusted or repaired if necessary, and that conformity with such standards was determined at that time. A licensed hearing instrument specialist shall maintain calibration records for ten years and shall make the records available for inspection by the department at any time. A licensed hearing instrument specialist shall also use routine procedures for the daily inspection of audiometric equipment, or prior to use if used less often than daily, to generally determine that the equipment is in normal working order.

(3) A licensed hearing instrument specialist shall provide the following care of the equipment used in the licensee's practice of ordering the use and fitting of hearing instruments:

(a) Hearing instruments, assistive-listening devices, and electronic equipment shall be maintained according to the manufacturer's specifications;

(b) Instrumental technology shall be maintained in proper working order and be properly calibrated according to accepted standards; and

(c) Proper infection control and sanitation procedures shall be utilized.

Sec. 27. (1) Prior to engaging in cerumen removal, a licensed hearing instrument specialist shall have held a valid, undisciplined

AM1154
LB332
MMM - 04/23/2025

license as a licensed hearing instrument specialist for a minimum of two consecutive years and provide the board with evidence of (a) successful completion of a cerumen removal course pursuant to subsection (3) of this section and (b) professional liability insurance pursuant to subsection (5) of this section. If the licensed hearing instrument specialist continues to engage in cerumen removal, the licensee shall annually provide evidence to the board of professional liability insurance.

(2) If the patient exhibits contraindications to cerumen removal requiring medical consultation or medical intervention, a licensed hearing instrument specialist shall refer the patient to an otolaryngologist or another physician licensed to practice medicine and surgery under the Uniform Credentialing Act. If a licensed hearing instrument specialist engaged in routine cerumen removal discovers any trauma, including, but not limited to, continuous uncontrolled bleeding, lacerations, or other traumatic injuries, the licensee shall, as soon as practicable, seek immediate medical attention for the patient.

(3)(a) Prior to engaging in cerumen removal, a licensed hearing instrument specialist shall complete a cerumen removal course recommended by a national medical or audiology organization and approved by the board and provide the board with evidence of such successful completion and attestation of competence. In order to be approved by the board as a cerumen removal course, the course shall be approved by a national medical or audiology organization and shall:

(i) Be overseen by a physician or an audiologist, preferably an otolaryngologist;

(ii) Consist of at least six hours of practice of cerumen removal from an ear canal model using a variety of safe techniques;

(iii) Include in-person practice of cerumen removal techniques;

(iv) Include an infectious control component; and

(v) Result in a certificate of successful completion and attestation of competence signed by such physician or audiologist.

AM1154
LB332
MMM - 04/23/2025

(b) The board may, only after consultation with the Board of Medicine and Surgery and the Board of Audiology and Speech-Language Pathology, adopt rules and regulations as provided in section 38-126 to provide requirements for the initial cerumen removal course.

(4) The licensed hearing instrument specialist shall maintain documentation evidencing the satisfactory completion of the training.

(5) A licensed hearing instrument specialist shall carry appropriate professional liability insurance before engaging in cerumen removal.

(6) A licensed hearing instrument specialist shall perform cerumen removal using the customary removal techniques that are commensurate with the licensee's training and experience. Performance of cerumen removal is limited to the patient's cartilaginous outer one-third portion of the external auditory canal.

(7) A licensed hearing instrument specialist engaged in cerumen removal shall comply with the following requirements:

(a) The indications for cerumen removal for a licensed hearing instrument specialist shall include:

(i) Enabling audiometric testing;

(ii) Making ear impressions;

(iii) Fitting ear protection or prosthetic devices; and

(iv) Monitoring continuous use of hearing aids;

(b) The licensed hearing instrument specialist shall refer a patient to an otolaryngologist or another physician licensed under the Uniform Credentialing Act for medical consultation or medical intervention if the patient exhibits any of the following contraindications to cerumen removal:

(i) The patient is younger than eighteen years of age;

(ii) The patient has a perforated tympanic membrane;

(iii) The patient has a history of pain or active drainage or bleeding from the ear;

(iv) There is evidence of congenital or traumatic deformity of the

AM1154
LB332
MMM - 04/23/2025

ear;

(v) The patient has had previous ear surgery;

(vi) The patient has tympanostomy tubes, such that irrigation should not be used;

(vii) The patient has a bleeding disorder;

(viii) The patient has an actual or suspected foreign body in the ear;

(ix) The patient has a stenosis or bony exostosis of the ear canal;

(x) The patient has a tympanic membrane that the licensed hearing instrument specialist is unable to see; or

(xi) There is any other contraindication to cerumen removal that requires medical consultation or medical intervention; and

(c) If the patient, while undergoing cerumen removal that did not present contraindications, complains of significant pain, exhibits uncontrolled bleeding or a laceration of the external auditory canal, or experiences the acute onset of dizziness or vertigo or sudden hearing loss, the licensed hearing instrument specialist shall immediately stop the procedure and refer the patient to an otolaryngologist or another physician licensed under the Uniform Credentialing Act.

(8) The licensed hearing instrument specialist shall maintain the following proper infection control practices:

(a) Universal health precautions;

(b) Decontamination;

(c) Cleaning, disinfection, and sterilization of multiple-use equipment; and

(d) Universal precautions for prevention of the transmission of human immunodeficiency virus, hepatitis B virus, and other bloodborne pathogens, as defined by occupational safety and health standards promulgated pursuant to 29 C.F.R. 1910, as such regulations existed on January 1, 2025.

(9) The licensed hearing instrument specialist who performs cerumen

AM1154
LB332
MMM - 04/23/2025

removal shall maintain a case history for every patient and informed consent signed by the patient as part of the patient's records. A licensed hearing instrument specialist shall include in the patient's record video-otoscopy pictures of the patient's ear canal showing cerumen that must be removed and video-otoscopy pictures after the removal of the cerumen.

(10) The licensed hearing instrument specialist shall carry appropriate professional liability insurance before performing cerumen removal.

(11) The licensed hearing instrument specialist is prohibited from requiring patients to sign any form that eliminates liability if the patient is harmed.

(12) A licensed hearing instrument specialist who passes the initial training in cerumen removal shall take one additional hour of continuing education specific to cerumen removal annually, by any approved means, in addition to the required continuing education requirements for the license as a licensed hearing instrument specialist.

Sec. 30. A purchaser of a hearing instrument shall not be required to obtain a medical evaluation for the repurchase of a hearing instrument after a medical evaluation has been obtained for certain otologic conditions that are permanent and would be reidentified at each hearing assessment. Such conditions shall include, but not be limited to:

(1) Visible congenital or traumatic deformity of the ear;

(2) Unilateral or asymmetric hearing loss, assuming no change in thresholds; and

(3) Audiometric air-bone gap equal to or greater than fifteen decibels at five hundred hertz, one thousand hertz, and two thousand hertz.

Sec. 32. A licensed hearing instrument specialist who is certified by the National Board for Certification in Hearing Instrument Sciences or has an advanced credential recognized by the board may work for a company

AM1154
LB332
MMM - 04/23/2025

or organization as a trainer and provide specialized training in the practical application of hearing instrument sciences.

Sec. 38. Sections 1, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 35, 36, 37, 39, and 41 of this act become operative three calendar months after the adjournment of this legislative session. The other sections of this act become operative on their effective date.

Sec. 39. Original sections 38-511, 38-1501, 38-1502, 38-1504, 38-1505, 38-1508, 38-1510, 38-1511, 38-1514, 38-2849, and 38-2884, Reissue Revised Statutes of Nebraska, and sections 38-1509, 38-1512, 38-1513, 38-28,104, and 68-911, Revised Statutes Cumulative Supplement, 2024, are repealed.

Sec. 41. The following sections are outright repealed: Sections 38-512 and 38-1506, Reissue Revised Statutes of Nebraska.

2. Renumber the remaining sections accordingly.