1	A bill to be entitled
2	An act relating to the Department of Health; creating
3	s. 381.875, F.S.; defining terms; prohibiting certain
4	research in this state relating to enhanced potential
5	pandemic pathogens; requiring researchers applying for
6	state or local funding to disclose certain
7	information; requiring the Department of Health to
8	enjoin violations of specified provisions; providing
9	construction; amending s. 381.986, F.S.; defining the
10	term "attractive to children"; prohibiting medical
11	marijuana treatment centers from producing marijuana
12	products that are attractive to children or
13	manufactured in specified manners; prohibiting
14	marijuana packaging and labeling from including
15	specified wording; prohibiting medical marijuana
16	treatment centers from using certain content in their
17	advertising which is attractive to children or
18	promotes the recreational use of marijuana; revising
19	background screening requirements for certain
20	individuals; amending s. 381.988, F.S.; requiring
21	medical marijuana testing laboratories to subject
22	their employees to background screenings; revising
23	background screening requirements for certain
24	individuals; amending s. 382.005, F.S.; requiring
25	local registrars to electronically file all live

Page 1 of 77

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26 birth, death, and fetal death records in their 27 respective jurisdictions in the department's 28 electronic registration system; requiring the local 29 registrars to file a paper record with the department if the electronic system is unavailable; requiring 30 31 local registrars to make blank paper forms available 32 in such instances; providing requirements for such 33 paper records; amending s. 382.008, F.S.; conforming 34 provisions to changes made by the act; amending s. 382.009, F.S.; revising the types of health care 35 36 practitioners who may make certain determinations of 37 death; amending ss. 382.013 and 382.015, F.S.; 38 conforming provisions to changes made by the act; 39 amending ss. 382.021 and 382.023, F.S.; revising the 40 frequency with which circuit courts must transmit 41 marriage licenses and certain dissolution-of-marriage 42 records to the department; requiring that such records 43 be transmitted electronically; amending s. 382.025, 44 F.S.; extending the timeframe for the confidentiality of certain birth records; authorizing persons 45 46 appointed by the department to issue certified copies 47 of live birth, death, and fetal death certificates; 48 amending s. 401.27, F.S.; revising requirements for 49 applicants for certification or recertification as 50 emergency medical technicians or paramedics; deleting

Page 2 of 77

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51	a requirement that a certain certification examination
52	be offered monthly; deleting related duties of the
53	department; deleting a temporary certificate and
54	related provisions; amending s. 401.2701, F.S.;
55	exempting certain emergency medical services training
56	program applicants from the requirement to have a
57	certain affiliation agreement; amending s. 401.272,
58	F.S.; revising the purpose of certain provisions;
59	specifying requirements for the provision of specified
60	services by paramedics and emergency medical
61	technicians under certain circumstances; revising the
62	department's rulemaking authority; amending s. 401.34,
63	F.S.; deleting certain provisions and fees related to
64	the department's grading of a certain certification
65	examination; amending s. 401.435, F.S.; revising
66	provisions related to minimum standards for emergency
67	medical responder training; amending s. 464.203, F.S.;
68	exempting certain applicants for certification as a
69	certified nursing assistant from the skills-
70	demonstration portion of a certain competency
71	examination; amending ss. 468.1225 and 468.1245, F.S.;
72	revising the scope of practice for audiologists, as it
73	relates to hearing aids to apply to prescription
74	hearing aids only; amending s. 468.1246, F.S.;
75	conforming provisions to changes made by the act;

Page 3 of 77

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76 deleting obsolete language; amending ss. 468.1255, 77 468.1265, and 468.1275, F.S.; conforming provisions to 78 changes made by the act; amending s. 484.0401, F.S.; revising legislative findings and intent to conform to 79 changes made by the act; reordering and amending s. 80 484.041, F.S.; providing and revising definitions; 81 82 amending s. 484.042, F.S.; revising membership 83 requirements for members of the Board of Hearing Aid 84 Specialists; amending s. 484.044, F.S.; revising the board's rulemaking authority; deleting obsolete 85 language; amending ss. 484.0445, 484.045, 484.0501, 86 87 and 484.051, F.S.; revising the scope of practice for 88 hearing aid specialists and making conforming changes 89 to licensure and practice requirements; amending s. 484.0512, F.S.; conforming provisions to changes made 90 91 by the act; deleting obsolete language; amending ss. 484.0513, 484.053, and 484.054, F.S.; conforming 92 93 provisions to changes made by the act; amending s. 94 484.059, F.S.; conforming provisions to changes made 95 by the act; providing applicability; providing a 96 directive to the Division of Law Revision; providing 97 effective dates. 98 99 Be It Enacted by the Legislature of the State of Florida: 100

Page 4 of 77

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101	Section 1. Effective upon this act becoming law, section
102	381.875, Florida Statutes, is created to read:
103	381.875 Enhanced potential pandemic pathogen research
104	prohibited
105	(1) As used in this section, the term:
106	(a) "Enhanced potential pandemic pathogen" means a
107	potential pandemic pathogen that results from enhancing the
108	transmissibility or virulence of a pathogen. The term does not
109	include naturally occurring pathogens circulating in or
110	recovered from nature, regardless of their pandemic potential.
111	(b) "Enhanced potential pandemic pathogen research" means
112	research that may be reasonably anticipated to create, transfer,
113	or use potential pandemic pathogens that result from enhancing a
1 1 1	nothegenle twonemicsibility on winulance in humans
114	pathogen's transmissibility or virulence in humans.
114	(c) "Potential pandemic pathogen" means a bacterium,
115	(c) "Potential pandemic pathogen" means a bacterium,
115 116	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both:
115 116 117	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: 1. Highly transmissible and capable of wide,
115 116 117 118	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: <u>1. Highly transmissible and capable of wide,</u> uncontrollable spread in human populations; and
115 116 117 118 119	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: <u>1. Highly transmissible and capable of wide,</u> <u>uncontrollable spread in human populations; and</u> <u>2. Highly virulent, making it likely to cause significant</u>
115 116 117 118 119 120	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: <u>1. Highly transmissible and capable of wide,</u> <u>uncontrollable spread in human populations; and</u> <u>2. Highly virulent, making it likely to cause significant</u> <u>morbidity or mortality in humans.</u>
115 116 117 118 119 120 121	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: 1. Highly transmissible and capable of wide, uncontrollable spread in human populations; and 2. Highly virulent, making it likely to cause significant morbidity or mortality in humans. (2) Any research that is reasonably likely to create an
115 116 117 118 119 120 121 122	<pre>(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: 1. Highly transmissible and capable of wide, uncontrollable spread in human populations; and 2. Highly virulent, making it likely to cause significant morbidity or mortality in humans. (2) Any research that is reasonably likely to create an enhanced potential pandemic pathogen or that has been determined</pre>
115 116 117 118 119 120 121 122 123	(c) "Potential pandemic pathogen" means a bacterium, virus, or other microorganism that is likely to be both: <u>1. Highly transmissible and capable of wide,</u> <u>uncontrollable spread in human populations; and</u> <u>2. Highly virulent, making it likely to cause significant</u> <u>morbidity or mortality in humans.</u> <u>(2) Any research that is reasonably likely to create an</u> <u>enhanced potential pandemic pathogen or that has been determined</u> <u>by the United States Department of Health and Human Services,</u>

Page 5 of 77

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126 (3) Any researcher applying for state or local funding to 127 conduct research in this state must disclose in the application 128 to the funding source whether the research meets the definition 129 of enhanced potential pandemic pathogen research. 130 The Department of Health shall exercise its authority (4) under s. 381.0012 to enjoin violations of this section. 131 132 (5) This section does not affect research funded or 133 conducted before the effective date of this act. 134 Section 2. Present paragraphs (a) through (o) of 135 subsection (1) of section 381.986, Florida Statutes, are 136 redesignated as paragraphs (b) through (p), respectively, a new 137 paragraph (a) is added to that subsection, and paragraphs (a) 138 and (c) of subsection (3), paragraphs (e) and (h) of subsection 139 (8), and subsection (9) of that section are amended, to read: 140 381.986 Medical use of marijuana.-141 (1) DEFINITIONS.-As used in this section, the term: 142 "Attractive to children" means the use of any image or (a) 143 words designed or likely to appeal to persons younger than 18 144 years of age, including, but not limited to, cartoons, toys, 145 animals, food, or depictions of persons younger than 18 years of age; any other likeness to images, characters, or phrases that 146 147 are popularly used to advertise to persons younger than 18 years 148 of age; or any reasonable likeness to commercially available candy. 149 150 (3) QUALIFIED PHYSICIANS AND MEDICAL DIRECTORS. -Page 6 of 77

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151 Before being approved as a qualified physician $\frac{1}{1}$ as (a) 152 defined in paragraph (1) (m), and before each license renewal, a 153 physician must successfully complete a 2-hour course and 154 subsequent examination offered by the Florida Medical 155 Association or the Florida Osteopathic Medical Association which 156 encompass the requirements of this section and any rules adopted 157 hereunder. The course and examination must shall be administered at least annually and may be offered in a distance learning 158 159 format, including an electronic, online format that is available 160 upon request. The price of the course may not exceed \$500. A 161 physician who has met the physician education requirements of former s. 381.986(4), Florida Statutes 2016, before June 23, 162 163 2017, shall be deemed to be in compliance with this paragraph 164 from June 23, 2017, until 90 days after the course and 165 examination required by this paragraph become available. 166 (C) Before being employed as a medical director, as 167 defined in paragraph (1)(i), and before each license renewal, a 168 medical director must successfully complete a 2-hour course and 169 subsequent examination offered by the Florida Medical 170 Association or the Florida Osteopathic Medical Association which 171 encompass the requirements of this section and any rules adopted 172 hereunder. The course and examination must shall be administered 173 at least annually and may be offered in a distance learning 174 format, including an electronic, online format that is available

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Page 7 of 77

upon request. The price of the course may not exceed \$500.

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2023

176 (8) MEDICAL MARIJUANA TREATMENT CENTERS.-177 A licensed medical marijuana treatment center shall (e) 178 cultivate, process, transport, and dispense marijuana for medical use. A licensed medical marijuana treatment center may 179 180 not contract for services directly related to the cultivation, processing, and dispensing of marijuana or marijuana delivery 181 182 devices, except that a medical marijuana treatment center 183 licensed pursuant to subparagraph (a)1. may contract with a 184 single entity for the cultivation, processing, transporting, and 185 dispensing of marijuana and marijuana delivery devices. A 186 licensed medical marijuana treatment center must, at all times, 187 maintain compliance with the criteria demonstrated and 188 representations made in the initial application and the criteria 189 established in this subsection. Upon request, the department may 190 grant a medical marijuana treatment center a variance from the 191 representations made in the initial application. Consideration 192 of such a request shall be based upon the individual facts and 193 circumstances surrounding the request. A variance may not be 194 granted unless the requesting medical marijuana treatment center 195 can demonstrate to the department that it has a proposed 196 alternative to the specific representation made in its 197 application which fulfills the same or a similar purpose as the 198 specific representation in a way that the department can 199 reasonably determine will not be a lower standard than the specific representation in the application. A variance may not 200

Page 8 of 77

201 be granted from the requirements in subparagraph 2. and 202 subparagraphs (b)1. and 2.

1. A licensed medical marijuana treatment center may transfer ownership to an individual or entity who meets the requirements of this section. A publicly traded corporation or publicly traded company that meets the requirements of this section is not precluded from ownership of a medical marijuana treatment center. To accommodate a change in ownership:

a. The licensed medical marijuana treatment center shall
notify the department in writing at least 60 days before the
anticipated date of the change of ownership.

212 b. The individual or entity applying for initial licensure 213 due to a change of ownership must submit an application that 214 must be received by the department at least 60 days before the 215 date of change of ownership.

c. Upon receipt of an application for a license, the department shall examine the application and, within 30 days after receipt, notify the applicant in writing of any apparent errors or omissions and request any additional information required.

d. Requested information omitted from an application for licensure must be filed with the department within 21 days after the department's request for omitted information or the application shall be deemed incomplete and shall be withdrawn from further consideration and the fees shall be forfeited.

Page 9 of 77

e. Within 30 days after the receipt of a completeapplication, the department shall approve or deny theapplication.

229 2. A medical marijuana treatment center, and any 230 individual or entity who directly or indirectly owns, controls, 231 or holds with power to vote 5 percent or more of the voting 232 shares of a medical marijuana treatment center, may not acquire 233 direct or indirect ownership or control of any voting shares or 234 other form of ownership of any other medical marijuana treatment 235 center.

3. A medical marijuana treatment center may not enter into
any form of profit-sharing arrangement with the property owner
or lessor of any of its facilities where cultivation,
processing, storing, or dispensing of marijuana and marijuana
delivery devices occurs.

4. All employees of a medical marijuana treatment center
must be 21 years of age or older and have passed a background
screening pursuant to subsection (9).

5. Each medical marijuana treatment center must adopt and enforce policies and procedures to ensure employees and volunteers receive training on the legal requirements to dispense marijuana to qualified patients.

248 6. When growing marijuana, a medical marijuana treatment 249 center:

250

a. May use pesticides determined by the department, after

Page 10 of 77

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251 consultation with the Department of Agriculture and Consumer 252 Services, to be safely applied to plants intended for human 253 consumption, but may not use pesticides designated as 254 restricted-use pesticides pursuant to s. 487.042.

255 b. Must grow marijuana within an enclosed structure and in256 a room separate from any other plant.

c. Must inspect seeds and growing plants for plant pests that endanger or threaten the horticultural and agricultural interests of the state in accordance with chapter 581 and any rules adopted thereunder.

261 d. Must perform fumigation or treatment of plants, or
262 remove and destroy infested or infected plants, in accordance
263 with chapter 581 and any rules adopted thereunder.

264 7. Each medical marijuana treatment center must produce 265 and make available for purchase at least one low-THC cannabis 266 product.

267 8. A medical marijuana treatment center that produces 268 edibles must hold a permit to operate as a food establishment 269 pursuant to chapter 500, the Florida Food Safety Act, and must 270 comply with all the requirements for food establishments 271 pursuant to chapter 500 and any rules adopted thereunder. 272 Edibles may not contain more than 200 milligrams of 273 tetrahydrocannabinol, and a single serving portion of an edible 274 may not exceed 10 milligrams of tetrahydrocannabinol. Edibles may have a potency variance of no greater than 15 percent. 275

Page 11 of 77

276 Marijuana products, including edibles, may not be attractive to 277 children; be manufactured in the shape of humans, cartoons, or 278 animals; be manufactured in a form that bears any reasonable 279 resemblance to products available for consumption as 280 commercially available candy; or contain any color additives. To 281 discourage consumption of edibles by children, the department 282 shall determine by rule any shapes, forms, and ingredients allowed and prohibited for edibles. Medical marijuana treatment 283 284 centers may not begin processing or dispensing edibles until 285 after the effective date of the rule. The department shall also 286 adopt sanitation rules providing the standards and requirements for the storage, display, or dispensing of edibles. 287

288 9. Within 12 months after licensure, a medical marijuana 289 treatment center must demonstrate to the department that all of 290 its processing facilities have passed a Food Safety Good 291 Manufacturing Practices, such as Global Food Safety Initiative 292 or equivalent, inspection by a nationally accredited certifying 293 body. A medical marijuana treatment center must immediately stop 294 processing at any facility which fails to pass this inspection 295 until it demonstrates to the department that such facility has 296 met this requirement.

297 10. A medical marijuana treatment center that produces 298 prerolled marijuana cigarettes may not use wrapping paper made 299 with tobacco or hemp.

300

11. When processing marijuana, a medical marijuana

Page 12 of 77

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301 treatment center must:

302 a. Process the marijuana within an enclosed structure and303 in a room separate from other plants or products.

b. Comply with department rules when processing marijuana with hydrocarbon solvents or other solvents or gases exhibiting potential toxicity to humans. The department shall determine by rule the requirements for medical marijuana treatment centers to use such solvents or gases exhibiting potential toxicity to humans.

310 c. Comply with federal and state laws and regulations and 311 department rules for solid and liquid wastes. The department 312 shall determine by rule procedures for the storage, handling, 313 transportation, management, and disposal of solid and liquid 314 waste generated during marijuana production and processing. The 315 Department of Environmental Protection shall assist the 316 department in developing such rules.

317 Test the processed marijuana using a medical marijuana d. 318 testing laboratory before it is dispensed. Results must be 319 verified and signed by two medical marijuana treatment center 320 employees. Before dispensing, the medical marijuana treatment 321 center must determine that the test results indicate that low-THC cannabis meets the definition of low-THC cannabis, the 322 concentration of tetrahydrocannabinol meets the potency 323 requirements of this section, the labeling of the concentration 324 325 of tetrahydrocannabinol and cannabidiol is accurate, and all

Page 13 of 77

326 marijuana is safe for human consumption and free from 327 contaminants that are unsafe for human consumption. The 328 department shall determine by rule which contaminants must be tested for and the maximum levels of each contaminant which are 329 330 safe for human consumption. The Department of Agriculture and 331 Consumer Services shall assist the department in developing the 332 testing requirements for contaminants that are unsafe for human 333 consumption in edibles. The department shall also determine by 334 rule the procedures for the treatment of marijuana that fails to 335 meet the testing requirements of this section, s. 381.988, or 336 department rule. The department may select samples of marijuana 337 from a medical marijuana treatment center facility which shall 338 be tested by the department to determine whether the marijuana 339 meets the potency requirements of this section, is safe for 340 human consumption, and is accurately labeled with the 341 tetrahydrocannabinol and cannabidiol concentration or to verify 342 the result of marijuana testing conducted by a marijuana testing 343 laboratory. The department may also select samples of marijuana 344 delivery devices from a medical marijuana treatment center to 345 determine whether the marijuana delivery device is safe for use 346 by qualified patients. A medical marijuana treatment center may 347 not require payment from the department for the sample. A medical marijuana treatment center must recall marijuana, 348 349 including all marijuana and marijuana products made from the same batch of marijuana, that fails to meet the potency 350

Page 14 of 77

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351 requirements of this section, that is unsafe for human 352 consumption, or for which the labeling of the 353 tetrahydrocannabinol and cannabidiol concentration is 354 inaccurate. The department shall adopt rules to establish 355 marijuana potency variations of no greater than 15 percent using 356 negotiated rulemaking pursuant to s. 120.54(2)(d) which accounts 357 for, but is not limited to, time lapses between testing, testing 358 methods, testing instruments, and types of marijuana sampled for 359 testing. The department may not issue any recalls for product 360 potency as it relates to product labeling before issuing a rule 361 relating to potency variation standards. A medical marijuana 362 treatment center must also recall all marijuana delivery devices 363 determined to be unsafe for use by qualified patients. The 364 medical marijuana treatment center must retain records of all 365 testing and samples of each homogenous batch of marijuana for at 366 least 9 months. The medical marijuana treatment center must 367 contract with a marijuana testing laboratory to perform audits 368 on the medical marijuana treatment center's standard operating 369 procedures, testing records, and samples and provide the results 370 to the department to confirm that the marijuana or low-THC 371 cannabis meets the requirements of this section and that the 372 marijuana or low-THC cannabis is safe for human consumption. A 373 medical marijuana treatment center shall reserve two processed 374 samples from each batch and retain such samples for at least 9 375 months for the purpose of such audits. A medical marijuana

Page 15 of 77

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376 treatment center may use a laboratory that has not been 377 certified by the department under s. 381.988 until such time as 378 at least one laboratory holds the required certification, but in 379 no event later than July 1, 2018. 380 Package the marijuana in compliance with the United e. 381 States Poison Prevention Packaging Act of 1970, 15 U.S.C. ss. 382 1471 et seq. Package the marijuana in a receptacle that has a firmly 383 f. 384 affixed and legible label stating the following information: 385 The marijuana or low-THC cannabis meets the (I) 386 requirements of sub-subparagraph d. 387 The name of the medical marijuana treatment center (II)from which the marijuana originates. 388 389 (III) The batch number and harvest number from which the 390 marijuana originates and the date dispensed. 391 (IV) The name of the physician who issued the physician 392 certification. 393 (V) The name of the patient. 394 The product name, if applicable, and dosage form, (VI) 395 including concentration of tetrahydrocannabinol and cannabidiol. 396 The product name may not contain wording commonly associated 397 with products that are attractive to children or which promote 398 the recreational use of marijuana marketed by or to children. 399 (VII) The recommended dose. 400 (VIII) A warning that it is illegal to transfer medical

Page 16 of 77

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401 marijuana to another person. 402 A marijuana universal symbol developed by the (IX) 403 department. 404 12. The medical marijuana treatment center shall include 405 in each package a patient package insert with information on the 406 specific product dispensed related to: 407 a. Clinical pharmacology. 408 Indications and use. b. 409 с. Dosage and administration. Dosage forms and strengths. 410 d. Contraindications. 411 e. 412 Warnings and precautions. f. 413 Adverse reactions. g. 414 In addition to the packaging and labeling requirements 13. 415 specified in subparagraphs 11. and 12., marijuana in a form for 416 smoking must be packaged in a sealed receptacle with a legible 417 and prominent warning to keep away from children and a warning 418 that states marijuana smoke contains carcinogens and may 419 negatively affect health. Such receptacles for marijuana in a 420 form for smoking must be plain, opaque, and white without 421 depictions of the product or images other than the medical 422 marijuana treatment center's department-approved logo and the 423 marijuana universal symbol. 424 14. The department shall adopt rules to regulate the 425 types, appearance, and labeling of marijuana delivery devices

Page 17 of 77

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426 dispensed from a medical marijuana treatment center. The rules 427 must require marijuana delivery devices to have an appearance 428 consistent with medical use.

429 15. Each edible must shall be individually sealed in 430 plain, opaque wrapping marked only with the marijuana universal 431 symbol. Where practical, each edible must shall be marked with 432 the marijuana universal symbol. In addition to the packaging and 433 labeling requirements in subparagraphs 11. and 12., edible 434 receptacles must be plain, opaque, and white without depictions 435 of the product or images other than the medical marijuana 436 treatment center's department-approved logo and the marijuana 437 universal symbol. The receptacle must also include a list of all the edible's ingredients, storage instructions, an expiration 438 439 date, a legible and prominent warning to keep away from children 440 and pets, and a warning that the edible has not been produced or 441 inspected pursuant to federal food safety laws.

442 16. When dispensing marijuana or a marijuana delivery443 device, a medical marijuana treatment center:

a. May dispense any active, valid order for low-THC
cannabis, medical cannabis and cannabis delivery devices issued
pursuant to former s. 381.986, Florida Statutes 2016, which was
entered into the medical marijuana use registry before July 1,
2017.

b. May not dispense more than a 70-day supply of marijuana within any 70-day period to a qualified patient or caregiver.

Page 18 of 77

May not dispense more than one 35-day supply of marijuana in a form for smoking within any 35-day period to a qualified patient or caregiver. A 35-day supply of marijuana in a form for smoking may not exceed 2.5 ounces unless an exception to this amount is approved by the department pursuant to paragraph (4)(f).

456 c. Must have the medical marijuana treatment center's 457 employee who dispenses the marijuana or a marijuana delivery 458 device enter into the medical marijuana use registry his or her 459 name or unique employee identifier.

460 Must verify that the qualified patient and the d. 461 caregiver, if applicable, each have an active registration in 462 the medical marijuana use registry and an active and valid 463 medical marijuana use registry identification card, the amount 464 and type of marijuana dispensed matches the physician 465 certification in the medical marijuana use registry for that 466 qualified patient, and the physician certification has not 467 already been filled.

e. May not dispense marijuana to a qualified patient who
is younger than 18 years of age. If the qualified patient is
younger than 18 years of age, marijuana may only be dispensed to
the qualified patient's caregiver.

f. May not dispense or sell any other type of cannabis, alcohol, or illicit drug-related product, including pipes or wrapping papers made with tobacco or hemp, other than a marijuana delivery device required for the medical use of

Page 19 of 77

2023

476 marijuana and which is specified in a physician certification. 477 Must, upon dispensing the marijuana or marijuana q. 478 delivery device, record in the registry the date, time, 479 quantity, and form of marijuana dispensed; the type of marijuana 480 delivery device dispensed; and the name and medical marijuana 481 use registry identification number of the qualified patient or 482 caregiver to whom the marijuana delivery device was dispensed. 483 Must ensure that patient records are not visible to h. 484 anyone other than the qualified patient, his or her careqiver, 485 and authorized medical marijuana treatment center employees. 486 (h) A medical marijuana treatment center may not engage in 487 advertising that is visible to members of the public from any 488 street, sidewalk, park, or other public place, except: 489 The dispensing location of a medical marijuana 1. 490 treatment center may have a sign that is affixed to the outside 491 or hanging in the window of the premises which identifies the 492 dispensary by the licensee's business name, a department-493 approved trade name, or a department-approved logo. A medical 494 marijuana treatment center's trade name and logo may not contain 495 wording or images that are attractive to children commonly 496 associated with marketing targeted toward children or which 497 promote recreational use of marijuana. A medical marijuana treatment center may engage in 498 2.

498 2. A medical marijuana treatment center may engage in 499 Internet advertising and marketing under the following 500 conditions:

Page 20 of 77

501 All advertisements must be approved by the department. a. 502 An advertisement may not have any content that is b. 503 attractive to children or which promotes the recreational use of 504 marijuana specifically targets individuals under the age of 18, 505 including cartoon characters or similar images. 506 An advertisement may not be an unsolicited pop-up с. 507 advertisement. 508 d. Opt-in marketing must include an easy and permanent 509 opt-out feature. 510 (9) BACKGROUND SCREENING. - An individual required to undergo a background screening pursuant to this section must 511 512 pass a level 2 background screening as provided under chapter 513 435, which, in addition to the disqualifying offenses provided in s. 435.04, shall exclude an individual who has an arrest 514 515 awaiting final disposition for, has been found quilty of, 516 regardless of adjudication, or has entered a plea of nolo 517 contendere or guilty to an offense under chapter 837, chapter 518 895, or chapter 896 or similar law of another jurisdiction. 519 Exemptions from disgualification as provided under s. 435.07 do not apply to this subsection. 520 Such individual must submit a full set of fingerprints 521 (a) to the department or to a vendor, entity, or agency authorized 522 523 by s. 943.053(13). The department, vendor, entity, or agency 524 shall forward the fingerprints to the Department of Law 525 Enforcement for state processing, and the Department of Law

Page 21 of 77

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526 Enforcement shall forward the fingerprints to the Federal Bureau 527 of Investigation for national processing. 528 (b) Fees for state and federal fingerprint processing and retention shall be borne by the medical marijuana treatment 529 530 center or caregiver, as applicable individual. The state cost 531 for fingerprint processing shall be as provided in s. 532 943.053(3)(e) for records provided to persons or entities other 533 than those specified as exceptions therein. 534 (C) Fingerprints submitted to the Department of Law 535 Enforcement pursuant to this subsection shall be retained by the 536 Department of Law Enforcement as provided in s. 943.05(2)(g) and 537 (h) and, when the Department of Law Enforcement begins 538 participation in the program, enrolled in the Federal Bureau of 539 Investigation's national retained print arrest notification program. Any arrest record identified shall be reported to the 540 541 department. 542 Section 3. Paragraph (d) of subsection (1) of section 543 381.988, Florida Statutes, is amended to read: 544 381.988 Medical marijuana testing laboratories; marijuana 545 tests conducted by a certified laboratory.-546 (1)A person or entity seeking to be a certified marijuana 547 testing laboratory must: 548 Require all employees, owners, and managers to submit (d) 549 to and pass a level 2 background screening pursuant to chapter 435. The department s. 435.04 and shall deny certification if 550

Page 22 of 77

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551 the person or entity seeking certification has a disqualifying 552 offense as provided in s. 435.04 or has an arrest awaiting final 553 disposition for, has been found guilty of, or has entered a plea 554 of quilty or nolo contendere to, regardless of adjudication, any 555 offense listed in chapter 837, chapter 895, or chapter 896 or 556 similar law of another jurisdiction. Exemptions from 557 disqualification as provided under s. 435.07 do not apply to 558 this paragraph.

1. Such <u>employees</u>, owners, and managers must submit a full set of fingerprints to the department or to a vendor, entity, or agency authorized by s. 943.053(13). The department, vendor, entity, or agency shall forward the fingerprints to the Department of Law Enforcement for state processing, and the Department of Law Enforcement shall forward the fingerprints to the Federal Bureau of Investigation for national processing.

2. Fees for state and federal fingerprint processing and retention shall be borne by <u>the certified marijuana testing</u> <u>laboratory</u> such owners or managers. The state cost for fingerprint processing shall be as provided in s. 943.053(3)(e) for records provided to persons or entities other than those specified as exceptions therein.

572 3. Fingerprints submitted to the Department of Law
573 Enforcement pursuant to this paragraph shall be retained by the
574 Department of Law Enforcement as provided in s. 943.05(2)(g) and
575 (h) and, when the Department of Law Enforcement begins

Page 23 of 77

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576 participation in the program, enrolled in the Federal Bureau of 577 Investigation's national retained print arrest notification 578 program. Any arrest record identified shall be reported to the 579 department.

580 Section 4. Section 382.005, Florida Statutes, is amended 581 to read:

582

382.005 Duties of local registrars.-

(1) Each local registrar is charged with the strict and thorough enforcement of the provisions of this chapter and rules adopted hereunder in his or her registration district, and shall make an immediate report to the department of any violation or apparent violation of this law or rules adopted hereunder.

588 (2) Each local registrar must electronically file all live 589 birth, death, and fetal death records within their respective 590 jurisdictions in the department's electronic registration 591 system. If the department's electronic registration system is 592 unavailable, the local registrar must file a paper record with 593 the department.

594 <u>(3)</u> Each local registrar <u>must</u> shall make available blank 595 forms <u>available if the department's electronic registration</u> 596 <u>system is unavailable</u>, as necessary and <u>must</u> shall examine each 597 <u>paper</u> certificate of live birth, death, or fetal death when 598 presented for registration in order to ascertain whether or not 599 it has been completed in accordance with the provisions of this 600 chapter and adopted rules. All paper birth, death, and fetal

Page 24 of 77

601 death certificates <u>must</u> shall be typewritten in permanent black 602 ink, and a <u>paper</u> certificate is not complete and correct if it 603 does not supply each item of information called for or 604 satisfactorily account for its omission.

605 (4) (4) (3) The local registrar or his or her deputy, if 606 authorized by the department, shall sign as registrar in 607 attestation of the date of registration of any paper records 608 filed, and may also make and preserve a local paper record of 609 each birth, death, and fetal death certificate registered by him or her, in such manner as directed by the department. The local 610 611 registrar shall transmit daily to the department all original 612 paper certificates registered. If no births, deaths, or fetal 613 deaths occurred in any month, the local registrar or deputy 614 shall, on the 7th day of the following month, report that fact 615 to the department on a form provided for such purpose.

616 <u>(5)(4)</u> Each local registrar, immediately upon appointment, 617 shall designate one or more deputy registrars to act on behalf 618 of the local registrar.

619 Section 5. Subsection (2) of section 382.008, Florida 620 Statutes, is amended to read:

621 382.008 Death, fetal death, and nonviable birth 622 registration.-

(2) (a) The funeral director who first assumes custody of a
dead body or fetus shall <u>electronically</u> file the certificate of
death or fetal death. In the absence of the funeral director,

Page 25 of 77

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2023

626 the physician, physician assistant, advanced practice registered 627 nurse registered under s. 464.0123, or other person in 628 attendance at or after the death or the district medical 629 examiner of the county in which the death occurred or the body 630 was found shall <u>electronically</u> file the certificate of death or 631 fetal death. The person who files the certificate shall obtain 632 personal data from a legally authorized person as described in 633 s. 497.005 or the best qualified person or source available. The 634 medical certification of cause of death must shall be furnished 635 to the funeral director, either in person or via certified mail 636 or electronic transfer, by the physician, physician assistant, advanced practice registered nurse registered under s. 464.0123, 637 638 or medical examiner responsible for furnishing such information. 639 For fetal deaths, the physician, physician assistant, advanced 640 practice registered nurse registered under s. 464.0123, midwife, 641 or hospital administrator shall provide any medical or health 642 information to the funeral director within 72 hours after 643 expulsion or extraction.

(b) The State Registrar <u>shall</u> may receive electronically a
certificate of death, fetal death, or nonviable birth which is
required to be filed with the registrar under this chapter
through facsimile or other electronic transfer for the purpose
of filing the certificate. The receipt of a certificate of
death, fetal death, or nonviable birth by electronic transfer
constitutes delivery to the State Registrar as required by law.

Page 26 of 77

651 Section 6. Subsection (2) of section 382.009, Florida 652 Statutes, is amended to read: 653 382.009 Recognition of brain death under certain 654 circumstances.-655 Determination of death pursuant to this section must (2) 656 shall be made in accordance with currently accepted reasonable 657 medical standards. 658 (a) If the patient's treating health care practitioner is 659 a physician licensed under chapter 458 or chapter 459, the 660 determination must be made by that physician and a second 661 physician two physicians licensed under chapter 458 or chapter 662 459 who is. One physician shall be the treating physician, and 663 the other physician shall be a board-eligible or board-certified 664 neurologist, neurosurgeon, internist, pediatrician, surgeon, or 665 anesthesiologist. 666 (b) If the patient's treating health care practitioner is 667 an autonomous advanced practice registered nurse registered 668 under s. 464.0123, the determination must be made by that 669 practitioner and two physicians licensed under chapter 458 or 670 chapter 459. Each physician must be a board-eligible or boardcertified neurologist, neurosurgeon, internist, pediatrician, 671 672 surgeon, or anesthesiologist. 673 Section 7. Section 382.013, Florida Statutes, is amended 674 to read: 675 382.013 Birth registration. - A certificate for each live

Page 27 of 77

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676 birth that occurs in this state shall be filed within 5 days 677 after such birth in the department's electronic registration 678 system with the local registrar of the district in which the 679 birth occurred and shall be registered by the local registrar if 680 the certificate has been completed and filed in accordance with 681 this chapter and adopted rules. The information regarding 682 registered births shall be used for comparison with information 683 in the state case registry, as defined in chapter 61.

684

(1) FILING.-

685 If a birth occurs in a hospital, birth center, or (a) 686 other health care facility, or en route thereto, the person in 687 charge of the facility is shall be responsible for preparing the 688 certificate, certifying the facts of the birth, and filing the 689 certificate in the department's electronic registration system 690 with the local registrar. Within 48 hours after the birth, the 691 physician, midwife, or person in attendance during or 692 immediately after the delivery shall provide the facility with 693 the medical information required by the birth certificate.

(b) If a birth occurs outside a facility and a physician
licensed in this state, a certified nurse midwife, a midwife
licensed in this state, or a public health nurse employed by the
department was in attendance during or immediately after the
delivery, that person shall prepare and file the certificate.

(c) If a birth occurs outside a facility and the deliveryis not attended by one of the persons described in paragraph

Page 28 of 77

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(b), the person in attendance, the mother, or the father shall report the birth to the registrar and provide proof of the facts of birth. The department may require such documents to be presented and such proof to be filed as it deems necessary and sufficient to establish the truth of the facts to be recorded by the certificate and may withhold registering the birth until its requirements are met.

(d) If a birth occurs in a moving conveyance and the child is first removed from the conveyance in this state, the birth shall be filed and registered in this state and the place to which the child is first removed shall be considered the place of birth.

(e) The mother or the father of the child shall attest to the accuracy of the personal data entered on the certificate in time to permit the timely registration of the certificate.

(f) If a certificate of live birth is incomplete, the local registrar shall immediately notify the health care facility or person filing the certificate and shall require the completion of the missing items of information if they can be obtained <u>before</u> prior to issuing certified copies of the birth certificate.

(g) Regardless of any plan to place a child for adoption after birth, the information on the birth certificate as required by this section must be as to the child's birth parents unless and until an application for a new birth record is made

Page 29 of 77

726 under s. 63.152.

(h) The State Registrar may receive electronically a birth certificate for each live birth which is required to be filed with the registrar under this chapter through facsimile or other electronic transfer for the purpose of filing the birth certificate. The receipt of a birth certificate by electronic transfer constitutes delivery to the State Registrar as required by law.

734

(2) PATERNITY.-

(a) If the mother is married at the time of birth, the
name of the husband shall be entered on the birth certificate as
the father of the child, unless paternity has been determined
otherwise by a court of competent jurisdiction.

(b) Notwithstanding paragraph (a), if the husband of the mother dies while the mother is pregnant but before the birth of the child, the name of the deceased husband shall be entered on the birth certificate as the father of the child, unless paternity has been determined otherwise by a court of competent jurisdiction.

(c) If the mother is not married at the time of the birth, the name of the father may not be entered on the birth certificate without the execution of an affidavit signed by both the mother and the person to be named as the father. The facility shall give notice orally or through the use of video or audio equipment, and in writing, of the alternatives to, the

Page 30 of 77

751 legal consequences of, and the rights, including, if one parent 752 is a minor, any rights afforded due to minority status, and 753 responsibilities that arise from signing an acknowledgment of 754 paternity, as well as information provided by the Title IV-D 755 agency established pursuant to s. 409.2557, regarding the 756 benefits of voluntary establishment of paternity. Upon request 757 of the mother and the person to be named as the father, the 758 facility shall assist in the execution of the affidavit, a 759 notarized voluntary acknowledgment of paternity, or a voluntary 760 acknowledgment of paternity that is witnessed by two individuals 761 and signed under penalty of perjury as specified by s. 762 92.525(2).

763 (d) If the paternity of the child is determined by a court 764 of competent jurisdiction as provided under s. 382.015 or there 765 is a final judgment of dissolution of marriage which requires 766 the former husband to pay child support for the child, the name 767 of the father and the surname of the child shall be entered on the certificate in accordance with the finding and order of the 768 769 court. If the court fails to specify a surname for the child, 770 the surname shall be entered in accordance with subsection (3).

(e) If the paternity of the child is determined pursuant to s. 409.256, the name of the father and the surname of the child shall be entered on the certificate in accordance with the finding and order of the Department of Revenue.

775

(f) If the mother and father marry each other at any time

Page 31 of 77

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776 after the child's birth, upon receipt of a marriage license that 777 identifies any such child, the department shall amend the 778 certificate with regard to the parents' marital status as though 779 the parents were married at the time of birth.

(g) If the father is not named on the certificate, no other information about the father shall be entered on the certificate.

783

(3) NAME OF CHILD.-

(a) If the mother is married at the time of birth, the mother and father whose names are entered on the birth certificate shall select the given names and surname of the child if both parents have custody of the child, otherwise the parent who has custody shall select the child's name.

789 If the mother and father whose names are entered on (b) 790 the birth certificate disagree on the surname of the child and 791 both parents have custody of the child, the surname selected by 792 the father and the surname selected by the mother shall both be 793 entered on the birth certificate, separated by a hyphen, with 794 the selected names entered in alphabetical order. If the parents 795 disagree on the selection of a given name, the given name may 796 not be entered on the certificate until a joint agreement that 797 lists the agreed upon given name and is notarized by both 798 parents is submitted to the department, or until a given name is 799 selected by a court.

800

(c) If the mother is not married at the time of birth, the

Page 32 of 77

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801 parent who will have custody of the child shall select the 802 child's given name and surname.

(d) If multiple names of the child exceed the space provided on the face of the birth certificate they shall be listed on the back of the certificate. Names listed on the back of the certificate shall be part of the official record.

807 (4) UNDETERMINED PARENTAGE.-The person having custody of a 808 child of undetermined parentage shall register a birth 809 certificate showing all known or approximate facts relating to 810 the birth. To assist in later determination, information 811 concerning the place and circumstances under which the child was 812 found shall be included on the portion of the birth certificate relating to marital status and medical details. In the event the 813 814 child is later identified, a new birth certificate shall be 815 prepared which shall bear the same number as the original birth 816 certificate, and the original certificate shall be sealed and 817 filed, shall be confidential and exempt from the provisions of 818 s. 119.07(1), and shall not be opened to inspection by, nor 819 shall certified copies of the same be issued except by court 820 order to, any person other than the registrant if of legal age. 821 (5)DISCLOSURE.-The original certificate of live birth shall contain all the information required by the department for 822 823 legal, social, and health research purposes. However, all

824 information concerning parentage, marital status, and medical825 details shall be confidential and exempt from the provisions of

Page 33 of 77

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826 s. 119.07(1), except for health research purposes as approved by 827 the department, nor shall copies of the same be issued except as 828 provided in s. 382.025.

829 Section 8. Section 382.015, Florida Statutes, is amended 830 to read:

831 382.015 New certificates of live birth; duty of clerks of 832 court and department.-The clerk of the court in which any 833 proceeding for adoption, annulment of an adoption, affirmation 834 of parental status, or determination of paternity is to be 835 registered, shall within 30 days after the final disposition, 836 forward electronically to the department a certified copy of the 837 court order, or a report of the proceedings upon a form to be furnished by the department, together with sufficient 838 839 information to identify the original birth certificate and to 840 enable the preparation of a new birth certificate. The clerk of 841 the court shall implement a monitoring and quality control plan 842 to ensure that all judicial determinations of paternity are 843 reported to the department in compliance with this section. The 844 department shall track paternity determinations reported monthly 845 by county, monitor compliance with the 30-day timeframe, and 846 report the data to the clerks of the court quarterly.

847

(1) ADOPTION AND ANNULMENT OF ADOPTION.-

848 (a) Upon receipt of the report or certified copy of an
849 adoption decree, together with the information necessary to
850 identify the original certificate of live birth, and establish a

Page 34 of 77

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851 new certificate, the department shall prepare and file a new 852 birth certificate, absent objection by the court decreeing the 853 adoption, the adoptive parents, or the adoptee if of legal age. 854 The certificate shall bear the same file number as the original 855 birth certificate. All names and identifying information 856 relating to the adoptive parents entered on the new certificate 857 shall refer to the adoptive parents, but nothing in the 858 certificate shall refer to or designate the parents as being 859 adoptive. All other items not affected by adoption shall be 860 copied as on the original certificate, including the date of 861 registration and filing.

862 Upon receipt of the report or certified copy of an (b) 863 annulment-of-adoption decree, together with the sufficient 864 information to identify the original certificate of live birth, 865 the department shall, if a new certificate of birth was filed 866 following an adoption report or decree, remove the new 867 certificate and restore the original certificate to its original 868 place in the files, and the certificate so removed shall be 869 sealed by the department.

(c) Upon receipt of a report or certified copy of an adoption decree or annulment-of-adoption decree for a person born in another state, the department shall forward the report or decree to the state of the registrant's birth. If the adoptee was born in Canada, the department shall send a copy of the report or decree to the appropriate birth registration authority

Page 35 of 77

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876 in Canada.

877 DETERMINATION OF PATERNITY.-Upon receipt of the (2) 878 report, a certified copy of a final decree of determination of 879 paternity, or a certified copy of a final judgment of 880 dissolution of marriage which requires the former husband to pay 881 child support for the child, together with sufficient 882 information to identify the original certificate of live birth, 883 the department shall prepare and file a new birth certificate, 884 which shall bear the same file number as the original birth 885 certificate. The registrant's name shall be entered as decreed 886 by the court or as reflected in the final judgment or support 887 order. The names and identifying information of the parents 888 shall be entered as of the date of the registrant's birth.

889 AFFIRMATION OF PARENTAL STATUS.-Upon receipt of an (3) 890 order of affirmation of parental status issued pursuant to s. 891 742.16, together with sufficient information to identify the 892 original certificate of live birth, the department shall prepare 893 and file a new birth certificate which shall bear the same file 894 number as the original birth certificate. The names and 895 identifying information of the registrant's parents entered on 896 the new certificate shall be the commissioning couple, but the 897 new certificate may not make reference to or designate the 898 parents as the commissioning couple.

899 (4) SUBSTITUTION OF NEW CERTIFICATE OF BIRTH FOR900 ORIGINAL.-When a new certificate of birth is prepared, the

Page 36 of 77

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901 department shall substitute the new certificate of birth for the 902 original certificate on file. All copies of the original 903 certificate of live birth in the custody of a local registrar or 904 other state custodian of vital records shall be forwarded to the 905 State Registrar. Thereafter, when a certified copy of the 906 certificate of birth or portion thereof is issued, it shall be a 907 copy of the new certificate of birth or portion thereof, except 908 when a court order requires issuance of a certified copy of the 909 original certificate of birth. In an adoption, change in 910 paternity, affirmation of parental status, undetermined 911 parentage, or court-ordered substitution, the department shall 912 place the original certificate of birth and all papers 913 pertaining thereto under seal, not to be broken except by order 914 of a court of competent jurisdiction or as otherwise provided by 915 law.

916 (5) FORM.-Except for certificates of foreign birth which 917 are registered as provided in s. 382.017, and delayed 918 certificates of birth which are registered as provided in ss. 919 382.019 and 382.0195, all original, new, or amended certificates 920 of live birth shall be identical in form, regardless of the 921 marital status of the parents or the fact that the registrant is 922 adopted or of undetermined parentage.

923 (6) RULES.—The department shall adopt and enforce all
924 rules necessary for carrying out the provisions of this section.
925 Section 9. Section 382.021, Florida Statutes, is amended

Page 37 of 77

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926 to read:

927 382.021 Department to receive marriage licenses.-Weekly On 928 or before the 5th day of each month, the county court judge or 929 clerk of the circuit court shall electronically transmit all 930 original marriage licenses, with endorsements, received during 931 the preceding calendar week month, to the department. Any 932 marriage licenses issued and not returned or any marriage 933 licenses returned but not recorded shall be reported by the 934 issuing county court judge or clerk of the circuit court to the 935 department at the time of transmitting the recorded licenses on 936 the forms to be prescribed and furnished by the department. If 937 during any month no marriage licenses are issued or returned, 938 the county court judge or clerk of the circuit court shall 939 report such fact to the department upon forms prescribed and 940 furnished by the department.

941 Section 10. Section 382.023, Florida Statutes, is amended 942 to read:

943 382.023 Department to receive dissolution-of-marriage 944 records; fees.-Clerks of the circuit courts shall collect for 945 their services at the time of the filing of a final judgment of 946 dissolution of marriage a fee of up to \$10.50, of which 43 947 percent shall be retained by the clerk of the circuit court as a 948 part of the cost in the cause in which the judgment is granted. 949 The remaining 57 percent shall be remitted to the Department of Revenue for deposit to the Department of Health to defray part 950

Page 38 of 77

951 of the cost of maintaining the dissolution-of-marriage records. 952 A record of each and every judgment of dissolution of marriage 953 granted by the court during the preceding calendar month, giving 954 names of parties and such other data as required by forms 955 prescribed by the department, shall be electronically 956 transmitted to the department weekly, on or before the 10th day 957 of each month, along with an accounting of the funds remitted to 958 the Department of Revenue pursuant to this section. 959 Section 11. Subsections (1) and (4) of section 382.025, 960 Florida Statutes, are amended to read: 382.025 Certified copies of vital records; 961 962 confidentiality; research.-963 BIRTH RECORDS.-Except for birth records over 125 100 (1)964 years old which are not under seal pursuant to court order, all 965 birth records of this state shall be confidential and are exempt 966 from the provisions of s. 119.07(1). 967 Certified copies of the original birth certificate or (a) 968 a new or amended certificate, or affidavits thereof, are 969 confidential and exempt from the provisions of s. 119.07(1) and, 970 upon receipt of a request and payment of the fee prescribed in 971 s. 382.0255, shall be issued only as authorized by the 972 department and in the form prescribed by the department, and 973 only: 974 To the registrant, if the registrant is of legal age, 1. 975 is a certified homeless youth, or is a minor who has had the

Page 39 of 77

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976 disabilities of nonage removed under s. 743.01 or s. 743.015; 977 2. To the registrant's parent or guardian or other legal 978 representative; 979 3. Upon receipt of the registrant's death certificate, to 980 the registrant's spouse or to the registrant's child, 981 grandchild, or sibling, if of legal age, or to the legal 982 representative of any of such person persons; 983 To any person if the birth record is more than 125 over 4. 984 100 years old and not under seal pursuant to court order; 985 To a law enforcement agency for official purposes; 5. 986 6. To any agency of the state or the United States for 987 official purposes upon approval of the department; or 988 Upon order of any court of competent jurisdiction. 7. 989 To protect the integrity of vital records and prevent (b) 990 the fraudulent use of the birth certificates of deceased 991 persons, the department shall match birth and death certificates 992 and post the fact of death to the appropriate birth certificate. 993 Except for a commemorative birth certificate, any certification 994 of a birth certificate of a deceased registrant shall be marked "deceased." In the case of a commemorative birth certificate, 995 996 such indication of death shall be made on the back of the 997 certificate. 998 (C) The department shall issue, upon request and upon 999 payment of an additional fee as prescribed under s. 382.0255, a 1000 commemorative birth certificate representing that the birth of

Page 40 of 77

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1001 the person named thereon is recorded in the office of the 1002 registrar. The certificate issued under this paragraph shall be 1003 in a form consistent with the need to protect the integrity of 1004 vital records but shall be suitable for display. It may bear the 1005 seal of the state printed thereon and may be signed by the 1006 Governor.

(4) CERTIFIED COPIES OF ORIGINAL CERTIFICATES.—Only the
state registrar, and local registrars, and those persons
appointed by the department are authorized to issue any
certificate which purports to be a certified copy of an original
certificate of live birth, death, or fetal death. Except as
provided in this section, preparing or issuing certificates is
exempt from the provisions of s. 119.07(1).

1014Section 12. Subsections (3), (4), and (5) of section1015401.27, Florida Statutes, are amended to read:

1016

401.27 Personnel; standards and certification.-

1017 Any person who desires to be certified or recertified (3) 1018 as an emergency medical technician or paramedic must apply to 1019 the department under oath on forms provided by the department 1020 which shall contain such information as the department 1021 reasonably requires, which may include affirmative evidence of 1022 ability to comply with applicable laws and rules. The department 1023 shall determine whether the applicant meets the requirements 1024 specified in this section and in rules of the department and shall issue a certificate to any person who meets such 1025

Page 41 of 77

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1026 requirements.

1027(4) An applicant for certification or recertification as1028an emergency medical technician or paramedic must:

1029 (a) Have completed an appropriate training program as1030 follows:

1031 1. For an emergency medical technician, an emergency 1032 medical technician training program approved by the department 1033 as equivalent to the most recent EMT-Basic National Standard 1034 Curriculum or the National EMS Education Standards of the United 1035 States Department of Transportation;

1036 2. For a paramedic, a paramedic training program approved 1037 by the department as equivalent to the most recent EMT-Paramedic 1038 National Standard Curriculum or the National EMS Education 1039 Standards of the United States Department of Transportation;

1040 (b) <u>Attest Certify under oath</u> that he or she is not 1041 addicted to alcohol or any controlled substance;

(c) <u>Attest Certify under oath</u> that he or she is free from any physical or mental defect or disease that might impair the applicant's ability to perform his or her duties;

(d) Within 2 years after program completion have passed anexamination developed or required by the department;

(e)1. For an emergency medical technician, hold a current American Heart Association cardiopulmonary resuscitation course card or an American Red Cross cardiopulmonary resuscitation course card or its equivalent as defined by department rule;

Page 42 of 77

1051 2. For a paramedic, hold a certificate of successful 1052 course completion in advanced cardiac life support from the 1053 American Heart Association or its equivalent as defined by 1054 department rule;

(f) Submit the certification fee and the nonrefundable examination fee prescribed in s. 401.34, which examination fee will be required for each examination administered to an applicant; and

(g) Submit a completed application to the department, which application documents compliance with paragraphs (a), (b), (c), (e), (f), and this paragraph, and, if applicable, paragraph (d). The application must be submitted so as to be received by the department at least 30 calendar days before the next regularly scheduled examination for which the applicant desires to be scheduled.

1066 (5) The certification examination must be offered monthly. 1067 The department shall issue an examination admission notice to 1068 the applicant advising him or her of the time and place of the 1069 or examination for which ho she is scheduled 1070 achieving a passing score on the certification examination may 1071 be issued a temporary certificate with their examination grade 1072 report. The department must issue an original certification 1073 within 45 days after the examination. Examination guestions and 1074 answers are not subject to discovery but may be introduced into evidence and considered only in camera in any administrative 1075

Page 43 of 77

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1084

1076 proceeding under chapter 120. If an administrative hearing is 1077 held, the department shall provide challenged examination 1078 questions and answers to the administrative law judge. The 1079 department shall establish by rule the procedure by which an 1080 applicant, and the applicant's attorney, may review examination 1081 questions and answers in accordance with s. 119.071(1)(a).

1082Section 13. Paragraph (a) of subsection (1) of section1083401.2701, Florida Statutes, is amended to read:

401.2701 Emergency medical services training programs.-

1085 (1) Any private or public institution in Florida desiring 1086 to conduct an approved program for the education of emergency 1087 medical technicians and paramedics shall:

1088 (a) Submit a completed application on a form provided by1089 the department, which must include:

1090 1. Evidence that the institution is in compliance with all 1091 applicable requirements of the Department of Education.

1092 2. Evidence of an affiliation agreement with a hospital 1093 that has an emergency department staffed by at least one 1094 physician and one registered nurse.

3. Evidence of an affiliation agreement with a current emergency medical services provider that is licensed in this state. Such agreement shall include, at a minimum, a commitment by the provider to conduct the field experience portion of the education program. <u>An applicant licensed as an advanced life</u> support service under s. 401.25 with permitted transport

Page 44 of 77

1101 <u>vehicles pursuant to s. 401.26 is exempt from the requirements</u> 1102 <u>of this subparagraph and need not submit evidence of an</u> 1103 <u>affiliation agreement with a current emergency medical services</u>

1104 provider.

1105

4. Documentation verifying faculty, including:

a. A medical director who is a licensed physician meeting
the applicable requirements for emergency medical services
medical directors as outlined in this chapter and rules of the
department. The medical director shall have the duty and
responsibility of certifying that graduates have successfully
completed all phases of the education program and are proficient
in basic or advanced life support techniques, as applicable.

b. A program director responsible for the operation, organization, periodic review, administration, development, and approval of the program.

1116

5. Documentation verifying that the curriculum:

a. Meets the most recent Emergency Medical TechnicianBasic National Standard Curriculum or the National EMS Education
Standards approved by the department for emergency medical
technician programs and Emergency Medical Technician-Paramedic
National Standard Curriculum or the National EMS Education
Standards approved by the department for paramedic programs.

b. Includes 2 hours of instruction on the trauma scorecard
methodologies for assessment of adult trauma patients and
pediatric trauma patients as specified by the department by

Page 45 of 77

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1126 rule.

1127 6. Evidence of sufficient medical and educational
1128 equipment to meet emergency medical services training program
1129 needs.

1130 Section 14. Section 401.272, Florida Statutes, is amended 1131 to read:

1132

401.272 Emergency medical services community health care.-

(1) The purpose of this section is to encourage more effective utilization of the skills of emergency medical technicians and paramedics by enabling them to perform, in partnership with local county health departments, specific additional health care tasks that are consistent with the public health and welfare.

1139 (2) Notwithstanding any other provision of law to the 1140 contrary:

1141 (a) Paramedics or emergency medical technicians shall 1142 operate under the medical direction of a physician through two-1143 way voice communication or pursuant to established standing orders or protocols and within the scope of their training when 1144 1145 providing basic life support, advanced life support, and may 1146 perform health promotion and wellness activities and blood 1147 pressure screenings in a nonemergency environment, within the 1148 scope of their training, and under the direction of a medical 1149 director. As used in this paragraph, the term "health promotion and wellness" means the provision of public health programs 1150

Page 46 of 77

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395.002(12).

1151

pertaining to the prevention of illness and injury.

1152 Paramedics and emergency medical technicians shall (b) 1153 operate under the medical direction of a physician through two-1154 way communication or pursuant to established standing orders or protocols and within the scope of their training when a patient 1155 1156 is not transported to an emergency department or is transported 1157 to a facility other than a hospital as defined in s. 1158

1159 (c) Paramedics may administer immunizations in a nonemergency environment, within the scope of their training, 1160 1161 and under the medical direction of a physician through two-way communication or pursuant to established standing orders or 1162 protocols medical director. There must be a written agreement 1163 1164 between the physician providing medical direction paramedic's medical director and the department or the county health 1165 1166 department located in each county in which the paramedic 1167 administers immunizations. This agreement must establish the 1168 protocols, policies, and procedures under which the paramedic 1169 must operate.

1170 (d) (c) Paramedics may provide basic life support services 1171 and advanced life support services to patients receiving acute 1172 and postacute hospital care at home as specified in the 1173 paramedic's supervisory relationship with a physician or 1174 standing orders as described in s. 401.265, s. 458.348, or s. 459.025. A physician who supervises or provides medical 1175

Page 47 of 77

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1176 direction to a paramedic who provides basic life support 1177 services or advanced life support services to patients receiving 1178 acute and postacute hospital care at home pursuant to a formal 1179 supervisory relationship or standing orders is liable for any act or omission of the paramedic acting under the physician's 1180 1181 supervision or medical direction when providing such services. 1182 The department may adopt and enforce rules necessary to 1183 implement this paragraph.

1184 (3)Each physician providing medical direction to medical 1185 director under whose direction a paramedic who administers 1186 immunizations must verify and document that the paramedic has received sufficient training and experience to administer 1187 immunizations. The verification must be documented on forms 1188 1189 developed by the department, and the completed forms must be 1190 maintained at the service location of the licensee and made 1191 available to the department upon request.

(4) The department may adopt and enforce all rules necessary to enforce the provisions relating to a paramedic's administration of immunizations and the performance of health promotion and wellness activities and blood pressure screenings by a paramedic or emergency medical technician in a nonemergency environment.

 1198
 Section 15.
 Subsections (5), (6), and (7) of section

 1199
 401.34, Florida Statutes, are amended to read:

 1200
 401.34
 Fees.

Page 48 of 77

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1201	(5) The department may provide same-day grading of the
1202	examination for an applicant for emergency medical technician or
1203	paramedic certification.
1204	(6) The department may offer walk-in eligibility
1205	determination and examination to applicants for emergency
1206	medical technician or paramedic certification who pay to the
1207	department a nonrefundable fee to be set by the department not
1208	to exceed \$65. The fee is in addition to the certification fee
1209	and examination fee. The department must establish locations and
1210	times for eligibility determination and examination.
1211	(7) The cost of emergency medical technician or paramedic
1212	certification examination review may not exceed \$50.
1213	Section 16. Section 401.435, Florida Statutes, is amended
1214	to read:
1215	401.435 Emergency medical First responder agencies and
1216	training
1217	(1) The department must adopt by rule the United States
1218	Department of Transportation National Emergency Medical Services
1219	Education Standards for the Emergency Medical Services: First
1220	Responder <u>level</u> Training Course as the minimum standard for
1221	<u>emergency medical</u> first responder training. In addition, the
1222	department must adopt rules establishing minimum <u>emergency</u>
1223	<u>medical</u> first responder instructor qualifications. For purposes
1224	of this section, <u>an emergency medical</u> a first responder includes
1225	any individual who receives training to render initial care to
	Dage 40 of 77

Page 49 of 77

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an ill or injured person, other than an individual trained and certified pursuant to s. 943.1395(1), but who does not have the primary responsibility of treating and transporting ill or injured persons.

1230 (2)Each emergency medical first responder agency must 1231 take all reasonable efforts to enter into a memorandum of 1232 understanding with the emergency medical services licensee 1233 within whose territory the agency operates in order to 1234 coordinate emergency services at an emergency scene. The 1235 department must provide a model memorandum of understanding for 1236 this purpose. The memorandum of understanding should include 1237 dispatch protocols, the roles and responsibilities of emergency 1238 medical first responder personnel at an emergency scene, and the 1239 documentation required for patient care rendered. For purposes 1240 of this section, the term "emergency medical first responder 1241 agency" includes a law enforcement agency, a fire service agency 1242 not licensed under this part, a lifeguard agency, and a 1243 volunteer organization that renders, as part of its routine 1244 functions, on-scene patient care before emergency medical 1245 technicians or paramedics arrive.

1246 Section 17. Paragraph (a) of subsection (1) of section 1247 464.203, Florida Statutes, is amended to read:

1248 464.203 Certified nursing assistants; certification 1249 requirement.-

1250

(1) The board shall issue a certificate to practice as a

Page 50 of 77

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1251 certified nursing assistant to any person who demonstrates a 1252 minimum competency to read and write and successfully passes the 1253 required background screening pursuant to s. 400.215. If the 1254 person has successfully passed the required background screening 1255 pursuant to s. 400.215 or s. 408.809 within 90 days before 1256 applying for a certificate to practice and the person's 1257 background screening results are not retained in the 1258 clearinghouse created under s. 435.12, the board shall waive the 1259 requirement that the applicant successfully pass an additional 1260 background screening pursuant to s. 400.215. The person must 1261 also meet one of the following requirements:

1262 Has successfully completed an approved training (a) program and achieved a minimum score, established by rule of the 1263 1264 board, on the nursing assistant competency examination, which 1265 consists of a written portion and skills-demonstration portion approved by the board and administered at a site and by 1266 1267 personnel approved by the department. Any person who has 1268 successfully completed an approved training program within 6 1269 months before filing an application for certification is not 1270 required to take the skills-demonstration portion of the 1271 competency examination. 1272 Section 18. Section 468.1225, Florida Statutes, is amended 1273 to read: 1274 468.1225 Procedures, equipment, and protocols.-1275 The following minimal procedures shall be used when a (1)

Page 51 of 77

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1276 licensed audiologist fits and sells a <u>prescription</u> hearing aid: 1277 (a) Pure tone audiometric testing by air and bone to 1278 determine the type and degree of hearing deficiency when 1279 indicated.

1280

(b) Effective masking when indicated.

(c) Appropriate testing to determine speech reception thresholds, speech discrimination scores, the most comfortable listening levels, uncomfortable loudness levels, and the selection of the best fitting arrangement for maximum hearing aid benefit when indicated.

1286

(2) The following equipment shall be used:

(a) A wide range audiometer <u>that</u> which meets the
specifications of the American National Standards Institute for
diagnostic audiometers when indicated.

(b) A speech audiometer or a master hearing aid in order
to determine the most comfortable listening level and speech
discrimination when indicated.

(3) A final fitting ensuring physical and operational comfort of the <u>prescription</u> hearing aid shall be made when indicated.

(4) A licensed audiologist who fits and sells <u>prescription</u>
hearing aids shall obtain the following medical clearance: If,
upon inspection of the ear canal with an otoscope in the common
procedure of fitting a <u>prescription</u> hearing aid and upon
interrogation of the client, there is any recent history of

Page 52 of 77

1301 infection or any observable anomaly, the client shall be 1302 instructed to see a physician, and a prescription hearing aid 1303 may shall not be fitted until medical clearance is obtained for 1304 the condition noted. If, upon return, the condition noted is no 1305 longer observable and the client signs a medical waiver, a 1306 prescription hearing aid may be fitted. Any person with a 1307 significant difference between bone conduction hearing and air 1308 conduction hearing must be informed of the possibility of 1309 medical or surgical correction.

(5) (a) A licensed audiologist's office must have available, or have access to, a selection of <u>prescription</u> hearing aid models, hearing aid supplies, and services complete enough to accommodate the various needs of the hearing aid wearers.

(b) At the time of the initial examination for fitting and sale of a <u>prescription</u> hearing aid, the attending audiologist must notify the prospective purchaser of the benefits of telecoil, also known as "t" coil or "t" switch, technology, including increased access to telephones and noninvasive access to assistive listening systems required under the Americans with Disabilities Act of 1990.

(6) Unless otherwise indicated, each audiometric test
conducted by a licensee or a certified audiology assistant in
the fitting and selling of prescription hearing aids <u>must</u> shall
be made in a testing room that has been certified by the

Page 53 of 77

1326 department, or by an agent approved by the department, not to 1327 exceed the following sound pressure levels at the specified 1328 frequencies: 250Hz-40dB, 500Hz-40dB, 750Hz-40dB, 1000Hz-40dB, 1329 1500Hz-42dB, 2000Hz-47dB, 3000Hz-52dB, 4000Hz-57dB, 6000Hz-62dB, 1330 and 8000Hz-67dB. An exception to this requirement shall be made 1331 in the case of a client who, after being provided written notice 1332 of the benefits and advantages of having the test conducted in a 1333 certified testing room, requests that the test be conducted in a 1334 place other than the licensee's certified testing room. Such 1335 request must shall be documented by a waiver that which includes 1336 the written notice and is signed by the licensee and the client 1337 before prior to the testing. The waiver must shall be executed 1338 on a form provided by the department. The executed waiver must 1339 shall be attached to the client's copy of the contract, and a copy of the executed waiver must shall be retained in the 1340 1341 licensee's file.

(7) The board <u>may</u> shall have the power to prescribe the minimum procedures and equipment used in the conducting of hearing assessments and for the fitting and selling of <u>prescription</u> hearing aids. The board shall adopt and enforce rules necessary to <u>implement</u> carry out the provisions of this subsection and subsection (6).

(8) Any duly authorized officer or employee of the
department <u>may</u> shall have the right to make such inspections and
investigations as are necessary in order to determine the state

Page 54 of 77

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of compliance with the provisions of this section and the applicable rules and may enter the premises of a licensee and inspect the records of same upon reasonable belief that a violation of this law is being or has been committed or that the licensee has failed or is failing to comply with the provisions of this part.

1357Section 19. Section 468.1245, Florida Statutes, is amended1358to read:

1359 468.1245 Itemized listing of prices; delivery of 1360 prescription hearing aid; receipt; guarantee; packaging; 1361 disclaimer.-

Before Prior to delivery of services or products to a 1362 (1)1363 prospective purchaser, a licensee must shall disclose, upon 1364 request by the prospective purchaser, an itemized listing of prices, which must listing shall include separate price 1365 1366 estimates for each service component and each product. Provision of such itemized listing of prices may shall not be predicated 1367 1368 on the prospective purchaser's payment of any charge or 1369 agreement to purchase any service or product.

(2) Any licensee who fits and sells a <u>prescription</u> hearing aid shall, at the time of delivery, provide the purchaser with a receipt containing the seller's signature, the address of his or her regular place of business, and his or her license or certification number, if applicable, together with the brand, model, manufacturer or manufacturer's identification code, and

Page 55 of 77

1376 serial number of the prescription hearing aid furnished and the 1377 amount charged for the prescription hearing aid. The receipt 1378 must also shall specify whether the prescription hearing aid is new, used, or rebuilt, and shall specify the length of time and 1379 other terms of the guarantee, and by whom the prescription 1380 1381 hearing aid is guaranteed. When the client has requested an 1382 itemized list of prices, the receipt must shall also provide an 1383 itemization of the total purchase price, including, but not 1384 limited to, the cost of the aid, ear mold, batteries, and other 1385 accessories, and the cost of any services. Notice of the 1386 availability of this service must be displayed in a conspicuous 1387 manner in the office. The receipt must also shall state that any 1388 complaint concerning the prescription hearing aid and its 1389 guarantee, if not reconciled with the licensee from whom the 1390 prescription hearing aid was purchased, should be directed by 1391 the purchaser to the department. The address and telephone 1392 number of such office must shall be stated on the receipt.

(3) <u>A prescription</u> No hearing aid may <u>not</u> be sold to any person unless both the packaging containing the <u>prescription</u> hearing aid and the contract provided pursuant to subsection (2) carry the following disclaimer in 10-point or larger type: "A hearing aid will not restore normal hearing, nor will it prevent further hearing loss."

1399 Section 20. Section 468.1246, Florida Statutes, is amended 1400 to read:

Page 56 of 77

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1401 468.1246 Thirty-day trial period; purchaser's right to 1402 cancel; notice; refund; cancellation fee.-1403 A person selling a prescription hearing aid in this (1) state must provide the buyer with written notice of a 30-day 1404 1405 trial period and money-back guarantee. The guarantee must permit 1406 the purchaser to cancel the purchase for a valid reason as 1407 defined by rule of the board within 30 days after receiving the 1408 prescription hearing aid, by returning the prescription hearing 1409 aid or mailing written notice of cancellation to the seller. If the prescription hearing aid must be repaired, remade, or 1410 adjusted during the 30-day trial period, the running of the 30-1411 day trial period is suspended 1 day for each 24-hour period that 1412 1413 the prescription hearing aid is not in the purchaser's 1414 possession. A repaired, remade, or adjusted prescription hearing aid must be claimed by the purchaser within 3 working days after 1415 1416 notification of availability. The running of the 30-day trial period resumes on the day the purchaser reclaims a repaired, 1417 1418 remade, or adjusted prescription hearing aid or on the 4th day after notification of availability. 1419 1420 The board, in consultation with the Board of Hearing (2)1421 Aid Specialists, shall prescribe by rule the terms and 1422 conditions to be contained in the money-back guarantee and any 1423 exceptions thereto. Such rule must shall provide, at a minimum,

1424 1425

Page 57 of 77

that the charges for earmolds and service provided to fit the

prescription hearing aid may be retained by the licensee. The

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1426 rules must shall also set forth any reasonable charges to be 1427 held by the licensee as a cancellation fee. Such rule shall be 1428 effective on or before December 1, 1994. Should the board fail 1429 to adopt such rule, a licensee may not charge a cancellation fee 1430 which exceeds 5 percent of the total charge for a hearing aid 1431 alone. The terms and conditions of the guarantee, including the 1432 total amount available for refund, must shall be provided in 1433 writing to the purchaser before prior to the signing of the 1434 contract.

1435 Section 21. Section 468.1255, Florida Statutes, is amended 1436 to read:

1437 468.1255 Cancellation by medical authorization; 1438 purchaser's right to return.-

1439 In addition to any other rights and remedies the (1)1440 purchaser of a prescription hearing aid may have, the purchaser 1441 has shall have the right to rescind the transaction if the 1442 purchaser for whatever reason consults a licensed physician with 1443 specialty board certification in otolaryngology or internal 1444 medicine or a licensed family practice physician, subsequent to 1445 purchasing a prescription hearing aid, and the physician 1446 certifies in writing that the purchaser has a hearing impairment 1447 for which a prescription hearing aid will not provide a benefit 1448 or that the purchaser has a medical condition which 1449 contraindicates the use of a prescription hearing aid. 1450 (2) The purchaser of a prescription hearing aid has shall

Page 58 of 77

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1451 have the right to rescind as provided in subsection (1) only if 1452 the purchaser gives a written notice of the intent to rescind 1453 the transaction to the seller at the seller's place of business 1454 by certified mail, return receipt requested, which notice shall be posted not later than 60 days following the date of delivery 1455 1456 of the prescription hearing aid to the purchaser, and the 1457 purchaser returns the prescription hearing aid to the seller in 1458 the original condition less normal wear and tear.

(3) If the conditions of subsections (1) and (2) are met, the seller <u>must shall</u>, without request, refund to the purchaser, within 10 days <u>after of</u> the receipt of notice to rescind, a full and complete refund of all moneys received, less 5 percent. The purchaser <u>does not shall</u> incur <u>any no</u> additional liability for rescinding the transaction.

1465Section 22.Section 468.1265, Florida Statutes, is amended1466to read:

1467 468.1265 Sale or distribution of <u>prescription</u> hearing aids 1468 through mail; penalty.—It is unlawful for any person to sell or 1469 distribute <u>prescription</u> hearing aids through the mail to the 1470 ultimate consumer. Any person who violates this section commits 1471 a misdemeanor of the second degree, punishable as provided in s. 1472 775.082 or s. 775.083.

1473 Section 23. Section 468.1275, Florida Statutes, is amended 1474 to read:

1475

468.1275 Place of business; display of license.-Each

Page 59 of 77

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1476 licensee who fits and sells a <u>prescription</u> hearing aid shall 1477 declare and establish a regular place of business, at which his 1478 or her license shall be conspicuously displayed.

1479 Section 24. Section 484.0401, Florida Statutes, is amended 1480 to read:

1481 484.0401 Purpose.-The Legislature recognizes that the 1482 dispensing of prescription hearing aids requires particularized 1483 knowledge and skill to ensure that the interests of the hearing-1484 impaired public will be adequately served and safely protected. 1485 It recognizes that a poorly selected or fitted prescription 1486 hearing aid not only will give little satisfaction but may interfere with hearing ability and, therefore, deems it 1487 1488 necessary in the interest of the public health, safety, and welfare to regulate the dispensing of prescription hearing aids 1489 in this state. Restrictions on the fitting and selling of 1490 1491 prescription hearing aids shall be imposed only to the extent necessary to protect the public from physical and economic harm, 1492 1493 and restrictions shall not be imposed in a manner which will 1494 unreasonably affect the competitive market.

1495Section 25.Section 484.041, Florida Statutes, is1496reordered and amended to read:

1497 484.041 Definitions.—As used in this part, the term:
1498 (1) "Board" means the Board of Hearing Aid Specialists.
1499 (2) "Department" means the Department of Health.
1500 (3) "Dispensing prescription hearing aids" means and

Page 60 of 77

includes:

1501

2023

1502 Conducting and interpreting hearing tests for purposes (a) 1503 of selecting suitable prescription hearing aids, making earmolds or ear impressions, and providing appropriate counseling. 1504 1505 (b) All acts pertaining to the selling, renting, leasing, 1506 pricing, delivery, and warranty of prescription hearing aids. (6) (4) "Hearing aid specialist" means a person duly 1507 1508 licensed in this state to practice the dispensing of 1509 prescription hearing aids. 1510 (4) (5) "Hearing aid" means any wearable an amplifying device designed for, offered for the purpose of, or represented 1511 1512 as aiding persons with, or compensating for, impaired hearing to be worn by a hearing-impaired person to improve hearing. 1513 1514 (10) (6) "Trainee" means a person studying prescription

1514 <u>(10)(0)</u> flamee means a person studying <u>prescription</u> 1515 hearing aid dispensing under the direct supervision of an active 1516 licensed hearing aid specialist for the purpose of qualifying 1517 for certification to sit for the licensure examination.

1518 <u>(5)</u> (7) "Hearing aid establishment" means any establishment 1519 in <u>this</u> the state which <u>employs a licensed hearing aid</u> 1520 <u>specialist who</u> offers, advertises, and performs hearing aid 1521 services for the general public.

1522(7)"Over-the-counter hearing aid" means an air-conduction1523hearing aid that does not require implantation or other surgical1524intervention and is intended for use by a person 18 years of age1525or older to compensate for perceived mild to moderate hearing

Page 61 of 77

1526 impairment.

1527 (8) "Prescription hearing aid" means a hearing aid that
 1528 satisfies the requirements of this part and is not an over-the 1529 counter hearing aid.

1530 <u>(9)(8)</u> "Sponsor" means an active, licensed hearing aid 1531 specialist under whose direct supervision one or more trainees 1532 are studying <u>prescription</u> hearing aid dispensing for the purpose 1533 of qualifying for certification to sit for the licensure 1534 examination.

1535 Section 26. Subsection (2) of section 484.042, Florida 1536 Statutes, is amended to read:

1537 484.042 Board of Hearing Aid Specialists; membership,1538 appointment, terms.-

1539 (2) Five members of the board shall be hearing aid 1540 specialists who have been licensed and practicing the dispensing 1541 of prescription hearing aids in this state for at least the 1542 preceding 4 years. The remaining four members, none of whom 1543 shall derive economic benefit from the fitting or dispensing of 1544 hearing aids, shall be appointed from the resident lay public of 1545 this state. One of the lay members shall be a prescription 1546 hearing aid user but may not neither be nor have been a hearing 1547 aid specialist or a licensee of a closely related profession. 1548 One lay member shall be an individual age 65 or over. One lay 1549 member shall be an otolaryngologist licensed pursuant to chapter 458 or chapter 459. 1550

Page 62 of 77

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1551Section 27.Subsection (2) of section 484.044, Florida1552Statutes, is amended to read:

1553

484.044 Authority to make rules.-

1554 (2)The board shall adopt rules requiring that each 1555 prospective purchaser of a prescription hearing aid be notified 1556 by the attending hearing aid specialist, at the time of the 1557 initial examination for fitting and sale of a hearing aid, of 1558 telecoil, "t" coil, or "t" switch technology. The rules shall 1559 further require that hearing aid specialists make available to 1560 prospective purchasers or clients information regarding 1561 telecoils, "t" coils, or "t" switches. These rules shall be 1562 effective on or before October 1, 1994.

Section 28. Subsection (2) of section 484.0445, Florida Statutes, is amended to read:

1565

484.0445 Training program.-

1566 A trainee shall perform the functions of a hearing aid (2) 1567 specialist in accordance with board rules only under the direct 1568 supervision of a licensed hearing aid specialist. The term 1569 "direct supervision" means that the sponsor is responsible for 1570 all work being performed by the trainee. The sponsor or a 1571 hearing aid specialist designated by the sponsor shall give 1572 final approval to work performed by the trainee and shall be 1573 physically present at the time the prescription hearing aid is 1574 delivered to the client.

1575

Section 29. Subsection (2) of section 484.045, Florida

Page 63 of 77

FLORIDA HOUSE OF REPRESEN	ITATIVES
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1576	Statutes, is amended to read:				
1577	484.045 Licensure by examination				
1578	(2) The department shall license each applicant who the				
1579	board certifies meets all of the following criteria:				
1580	(a) Has completed the application form and remitted the				
1581	required fees.+				
1582	(b) Is of good moral character <u>.</u> ;				
1583	(c) Is 18 years of age or older <u>.</u> +				
1584	(d) Is a graduate of an accredited high school or its				
1585	equivalent <u>.</u>				
1586	(e)1. Has met the requirements of the training program; or				
1587	2.a. Has a valid, current license as a hearing aid				
1588	specialist or its equivalent from another state and has been				
1589	actively practicing in such capacity for at least 12 months; or				
1590	b. Is currently certified by the National Board for				
1591	Certification in Hearing Instrument Sciences and has been				
1592	actively practicing for at least 12 months $\underline{\cdot} \dot{\tau}$				
1593	(f) Has passed an examination, as prescribed by board				
1594	rule <u>.</u> ; and				
1595	(g) Has demonstrated, in a manner designated by rule of				
1596	the board, knowledge of state laws and rules relating to the				
1597	fitting and dispensing of prescription hearing aids.				
1598	Section 30. Section 484.0501, Florida Statutes, is amended				
1599	to read:				
1600	484.0501 Minimal procedures and equipment				
Page 64 of 77					

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1601 The following minimal procedures shall be used in the (1)1602 fitting and selling of prescription hearing aids: 1603 Pure tone audiometric testing by air and bone to (a) 1604 determine the type and degree of hearing deficiency. 1605 Effective masking when indicated. (b) 1606 Appropriate testing to determine speech reception (C) 1607 thresholds, speech discrimination scores, the most comfortable 1608 listening levels, uncomfortable loudness levels, and the 1609 selection of the best fitting arrangement for maximum hearing 1610 aid benefit. 1611 (2)The following equipment shall be used: A wide range audiometer that which meets the 1612 (a) specifications of the American National Standards Institute for 1613 1614 diagnostic audiometers. 1615 A speech audiometer or a master hearing aid in order (b) 1616 to determine the most comfortable listening level and speech 1617 discrimination. 1618 (3) A final fitting ensuring physical and operational comfort of the prescription hearing aid shall be made. 1619 1620 The following medical clearance shall be obtained: If, (4)1621 upon inspection of the ear canal with an otoscope in the common procedure of a prescription hearing aid fitter and upon 1622 1623 interrogation of the client, there is any recent history of 1624 infection or any observable anomaly, the client must shall be instructed to see a physician, and a prescription hearing aid 1625 Page 65 of 77

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1626 <u>may shall</u> not be fitted until medical clearance is obtained for 1627 the condition noted. If, upon return, the condition noted is no 1628 longer observable and the client signs a medical waiver, a 1629 <u>prescription</u> hearing aid may be fitted. Any person with a 1630 significant difference between bone conduction hearing and air 1631 conduction hearing must be informed of the possibility of 1632 medical correction.

(5) (a) A prescription hearing aid establishment office must have available, or have access to, a selection of prescription hearing aid models, hearing aid supplies, and services complete enough to accommodate the various needs of the prescription hearing aid wearers.

(b) At the time of the initial examination for fitting and sale of a <u>prescription</u> hearing aid, the attending hearing aid specialist <u>shall</u> <u>must</u> notify the prospective purchaser or client of the benefits of telecoil, "t" coil, or "t" switch technology, including increased access to telephones and noninvasive access to assistive listening systems required under the Americans with Disabilities Act of 1990.

(6) Each audiometric test conducted by a licensee or
authorized trainee in the fitting and selling of prescription
hearing aids <u>must shall</u> be made in a testing room that has been
certified by the department, or by an agent approved by the
department, not to exceed the following sound pressure levels at
the specified frequencies: 250Hz-40dB, 500Hz-40dB, 750Hz-40dB,

Page 66 of 77

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1651 1000Hz-40dB, 1500Hz-42dB, 2000Hz-47dB, 3000Hz-52dB, 4000Hz-57dB, 1652 6000Hz-62dB, and 8000Hz-67dB. An exception to this requirement 1653 shall be made in the case of a client who, after being provided 1654 written notice of the benefits and advantages of having the test 1655 conducted in a certified testing room, requests that the test be 1656 conducted in a place other than the licensee's certified testing 1657 room. Such request must shall be documented by a waiver which 1658 includes the written notice and is signed by the licensee and 1659 the client before prior to the testing. The waiver must shall be 1660 executed on a form provided by the department. The executed 1661 waiver must shall be attached to the client's copy of the 1662 contract, and a copy of the executed waiver must shall be 1663 retained in the licensee's file.

1664 The board may shall have the power to prescribe the (7)minimum procedures and equipment which must shall be used in the 1665 1666 conducting of hearing assessments, and for the fitting and 1667 selling of prescription hearing aids, including equipment that will measure the prescription hearing aid's response curves to 1668 1669 ensure that they meet the manufacturer's specifications. These 1670 procedures and equipment may differ from those provided in this 1671 section in order to take full advantage of devices and equipment 1672 which may hereafter become available and which are demonstrated 1673 to be of greater efficiency and accuracy. The board shall adopt 1674 and enforce rules necessary to implement carry out the provisions of this subsection and subsection (6). 1675

Page 67 of 77

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1676 Any duly authorized officer or employee of the (8) 1677 department may shall have the right to make such inspections and 1678 investigations as are necessary in order to determine the state of compliance with the provisions of this section and the 1679 1680 applicable rules and may enter the premises of a licensee and 1681 inspect the records of same upon reasonable belief that a 1682 violation of this law is being or has been committed or that the 1683 licensee has failed or is failing to comply with the provisions 1684 of this part act. 1685 (9) A licensed hearing aid specialist may service, market, sell, dispense, provide customer support for, and distribute 1686 1687 prescription and over-the-counter hearing aids. Section 31. Section 484.051, Florida Statutes, is amended 1688 1689 to read: 1690 484.051 Itemization of prices; delivery of prescription 1691 hearing aid; receipt, packaging, disclaimer, guarantee.-1692 Before Prior to delivery of services or products to a (1)1693 prospective purchaser, any person who fits and sells 1694 prescription hearing aids must shall disclose on request by the 1695 prospective purchaser an itemized listing of prices, which must 1696 listing shall include separate price estimates for each service 1697 component and each product. Provision of such itemized listing 1698 of prices may shall not be predicated on the prospective 1699 purchaser's payment of any charge or agreement to purchase any service or product. 1700

Page 68 of 77

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1701 Any person who fits and sells a prescription hearing (2)1702 aid must shall, at the time of delivery, provide the purchaser 1703 with a receipt containing the seller's signature, the address of 1704 her or his regular place of business, and her or his license or 1705 trainee registration number, if applicable, together with the 1706 brand, model, manufacturer or manufacturer's identification 1707 code, and serial number of the prescription hearing aid 1708 furnished and the amount charged for the prescription hearing 1709 aid. The receipt must also shall specify whether the 1710 prescription hearing aid is new, used, or rebuilt, and shall 1711 specify the length of time and other terms of the guarantee, and by whom the prescription hearing aid is guaranteed. If When the 1712 1713 client has requested an itemized list of prices, the receipt 1714 must shall also provide an itemization of the total purchase price, including, but not limited to, the cost of the aid, 1715 1716 earmold, batteries and other accessories, and any services. Notice of the availability of this service shall be displayed in 1717 1718 a conspicuous manner in the office. The receipt must also shall 1719 state that any complaint concerning the prescription hearing aid 1720 and guarantee therefor, if not reconciled with the licensee from 1721 whom the prescription hearing aid was purchased, should be 1722 directed by the purchaser to the Department of Health. The 1723 address and telephone number of such office must shall be stated on the receipt. 1724

1725

(3) <u>A prescription</u> No hearing aid may <u>not</u> be sold to any

Page 69 of 77

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1726 person unless both the packaging containing the prescription 1727 hearing aid and the itemized receipt provided pursuant to 1728 subsection (2) carry the following disclaimer in 10-point or 1729 larger type: "A hearing aid will not restore normal hearing, nor 1730 will it prevent further hearing loss."

1731 Section 32. Section 484.0512, Florida Statutes, is amended 1732 to read:

1733484.0512Thirty-day trial period; purchaser's right to1734cancel; notice; refund; cancellation fee; criminal penalty.-

A person selling a prescription hearing aid in this 1735 (1) 1736 state must provide the buyer with written notice of a 30-day 1737 trial period and money-back guarantee. The guarantee must permit 1738 the purchaser to cancel the purchase for a valid reason, as 1739 defined by rule of the board rule, within 30 days after receiving the prescription hearing aid, by returning the 1740 1741 prescription hearing aid or mailing written notice of cancellation to the seller. If the prescription hearing aid must 1742 1743 be repaired, remade, or adjusted during the 30-day trial period, 1744 the running of the 30-day trial period is suspended 1 day for 1745 each 24-hour period that the prescription hearing aid is not in 1746 the purchaser's possession. A repaired, remade, or adjusted 1747 prescription hearing aid must be claimed by the purchaser within 1748 3 working days after notification of availability. The running 1749 of the 30-day trial period resumes on the day the purchaser reclaims the repaired, remade, or adjusted prescription hearing 1750

Page 70 of 77

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2023

1751 aid or on the fourth day after notification of availability, 1752 whichever occurs earlier. 1753 (2) The board, in consultation with the Board of Speech-1754 Language Pathology and Audiology, shall prescribe by rule the 1755 terms and conditions to be contained in the money-back guarantee 1756 and any exceptions thereto. Such rules must rule shall provide, 1757 at a minimum, that the charges for earmolds and service provided 1758 to fit the prescription hearing aid may be retained by the 1759 licensee. The rules must shall also set forth any reasonable 1760 charges to be held by the licensee as a cancellation fee. Such

1761 rule shall be effective on or before December 1, 1994. Should 1762 the board fail to adopt such rule, a licensee may not charge a 1763 cancellation fee which exceeds 5 percent of the total charge for 1764 a hearing aid alone. The terms and conditions of the guarantee, 1765 including the total amount available for refund, <u>must shall</u> be 1766 provided in writing to the purchaser <u>before prior to</u> the signing 1767 of the contract.

(3) Within 30 days after the return or attempted return of the prescription hearing aid, the seller shall refund all moneys that must be refunded to a purchaser pursuant to this section. A violation of this subsection is a misdemeanor of the first degree, punishable as provided in s. 775.082 or s. 775.083.

1773 (4) For purposes of this section, the term "seller" or
1774 "person selling a <u>prescription</u> hearing aid" includes:
1775 (a) Any natural person licensed under this part or any

Page 71 of 77

1776 other natural person who signs a sales receipt required by s. 1777 484.051(2) or s. 468.1245(2) or who otherwise fits, delivers, or 1778 dispenses a prescription hearing aid. 1779 (b) Any business organization, whether a sole proprietorship, partnership, corporation, professional 1780 1781 association, joint venture, business trust, or other legal 1782 entity, that which dispenses a prescription hearing aid or 1783 enters into an agreement to dispense a prescription hearing aid. 1784 Any person who controls, manages, or operates an (C) establishment or business that dispenses a prescription hearing 1785 1786 aid or enters into an agreement to dispense a prescription 1787 hearing aid. Section 33. Section 484.0513, Florida Statutes, is amended 1788 1789 to read: 484.0513 Cancellation by medical authorization; 1790 1791 purchaser's right to return.-In addition to any other rights and remedies the 1792 (1)1793 purchaser of a prescription hearing aid may have, the purchaser 1794 has shall have the right to rescind the transaction if the 1795 purchaser for whatever reason consults a licensed physician with 1796 specialty board certification in otolaryngology or internal 1797 medicine or a licensed family practice physician, subsequent to 1798 purchasing a prescription hearing aid, and the physician 1799 certifies in writing that the purchaser has a hearing impairment for which a prescription hearing aid will not provide a benefit 1800

Page 72 of 77

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1801 or that the purchaser has a medical condition which 1802 contraindicates the use of a prescription hearing aid.

1803 The purchaser of a prescription hearing aid has shall (2) 1804 have the right to rescind as provided in subsection (1) only if the purchaser gives a written notice of the intent to rescind 1805 1806 the transaction to the seller at the seller's place of business 1807 by certified mail, return receipt requested, which must notice 1808 shall be posted within not later than 60 days after following 1809 the date of delivery of the prescription hearing aid to the purchaser, and the purchaser returns the prescription hearing 1810 1811 aid to the seller in the original condition less normal wear and 1812 tear.

(3) If the conditions of subsections (1) and (2) are met,
the seller <u>must</u> shall, without request, refund to the purchaser,
within 10 days <u>after</u> of the receipt of <u>the</u> notice to rescind, a
full and complete refund of all moneys received, less 5 percent.
The purchaser <u>does not</u> shall incur <u>any</u> no additional liability
for rescinding the transaction.

1819 Section 34. Section 484.053, Florida Statutes, is amended 1820 to read:

1821 484.053 Prohibitions; penalties.-

1822 (1) A person may not:

(a) Practice dispensing <u>prescription</u> hearing aids unless
the person is a licensed hearing aid specialist;

1825 (b) Use the name or title "hearing aid specialist" when

Page 73 of 77

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1826 the person has not been licensed under this part; 1827 Present as her or his own the license of another; (C) 1828 Give false, incomplete, or forged evidence to the (d) board or a member thereof for the purposes of obtaining a 1829 1830 license; 1831 Use or attempt to use a hearing aid specialist license (e) 1832 that is delinquent or has been suspended, revoked, or placed on 1833 inactive status; 1834 (f) Knowingly employ unlicensed persons in the practice of dispensing prescription hearing aids; or 1835 1836 (q) Knowingly conceal information relative to violations 1837 of this part. Any person who violates any provision of the 1838 (2) 1839 provisions of this section is guilty of a felony of the third 1840 degree, punishable as provided in s. 775.082 or s. 775.083. 1841 (3) If a person licensed under this part allows the sale 1842 of a prescription hearing aid by an unlicensed person not 1843 registered as a trainee or fails to comply with the requirements 1844 of s. 484.0445(2) relating to supervision of trainees, the board 1845 must shall, upon determination of that violation, order the full 1846 refund of moneys paid by the purchaser upon return of the 1847 prescription hearing aid to the seller's place of business. 1848 Section 35. Section 484.054, Florida Statutes, is amended 1849 to read: 484.054 Sale or distribution of prescription hearing aids 1850

Page 74 of 77

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1851 through mail; penalty.-It is unlawful for any person to sell or 1852 distribute <u>prescription</u> hearing aids through the mail to the 1853 ultimate consumer. Any violation of this section constitutes a 1854 misdemeanor of the second degree, punishable as provided in s. 1855 775.082 or s. 775.083.

1856 Section 36. Section 484.059, Florida Statutes, is amended 1857 to read:

1858

484.059 Exemptions.-

(1) The licensure requirements of this part do not apply to any person engaged in recommending <u>prescription</u> hearing aids as part of the academic curriculum of an accredited institution of higher education, or as part of a program conducted by a public charitable institution supported primarily by voluntary contribution, provided this organization does not dispense or sell <u>prescription</u> hearing aids or accessories.

1866 (2)The licensure requirements of this part do not apply 1867 to any person licensed to practice medicine in this the state, 1868 except that such physician must shall comply with the 1869 requirement of periodic filing of the certificate of testing and 1870 calibration of audiometric equipment as provided in this part. A 1871 No person employed by or working under the supervision of a 1872 person licensed to practice medicine may not shall perform any 1873 services or acts which would constitute the dispensing of 1874 prescription hearing aids as defined in s. 484.041 s. 484.041(3), unless such person is a licensed hearing aid 1875

Page 75 of 77

1876 specialist.

1877 (3) The licensure requirements of this part do not apply
1878 to an audiologist licensed <u>under pursuant to</u> part I of chapter
1879 468.

1880 (4) <u>Section</u> The provisions of s. 484.053(1)(a) <u>does</u> shall
1881 not apply to registered trainees operating in compliance with
1882 this part and board rules of the board.

1883 (5) The licensure requirements of this part do not apply 1884 to a person who services, markets, sells, dispenses, provides 1885 customer support for, or distributes exclusively over-the-1886 counter hearing aids, whether through in-person transactions, by 1887 mail, or online. For purposes of this subsection, over-the-1888 counter hearing aids are those that are available without the 1889 supervision, prescription, or other order, involvement, or 1890 intervention of a licensed person to consumers through in-person 1891 transactions, by mail, or online. These devices allow the user 1892 to control the device and customize it to the user's hearing 1893 needs through the use of tools, tests, or software, including, 1894 but not limited to, wireless technology or tests for self-1895 assessment of hearing loss. 1896 Section 37. The Division of Law Revision is directed to 1897 replace the phrase "the effective date of this act" wherever it 1898 occurs in this act with the date the act becomes a law. 1899 Section 38. Except as otherwise expressly provided in this

1900 act and except for this section, which shall take effect upon

Page 76 of 77

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1901 this act becoming a law, this act shall take effect July 1, 1902 2023. Page 77 of 77