MISSISSIPPI LEGISLATURE

By: Representative Ford

To: Public Health and Welfare

## HOUSE BILL NO. 416

AN ACT TO AMEND SECTION 73-21-69, MISSISSIPPI CODE OF 1972, 1 TO EXTEND THE DATE OF REPEAL ON THE PROVISIONS OF LAW THAT CREATE 2 3 THE STATE BOARD OF PHARMACY AND PRESCRIBE ITS DUTIES AND POWERS; TO REENACT SECTIONS 73-21-71 THROUGH 73-21-123, MISSISSIPPI CODE OF 1972, WHICH CREATE THE STATE BOARD OF PHARMACY AND PRESCRIBE 4 5 ITS DUTIES AND POWERS; AND FOR RELATED PURPOSES. 6 7 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MISSISSIPPI: SECTION 1. Section 73-21-69, Mississippi Code of 1972, is 8 amended as follows: 9 10 73-21-69. Sections 73-21-71 through 73-21-123, which create the State Board of Pharmacy and prescribe its duties and powers, 11 shall stand repealed on July 1, 2003. 12 SECTION 2. Section 73-21-71, Mississippi Code of 1972, is 13 reenacted as follows: 14 73-21-71. This chapter shall be known as the "Mississippi 15 Pharmacy Practice Act." 16 17 SECTION 3. Section 73-21-73, Mississippi Code of 1972, is reenacted as follows: 18 73-21-73. As used in this chapter, unless the context 19 requires otherwise: 20 "Administer" shall mean the direct application of a 21 (a) prescription drug pursuant to a lawful order of a practitioner to 22 the body of a patient by injection, inhalation, ingestion or any 23 other means. 24 "Board of Pharmacy," "Pharmacy Board," "MSBP" or 25 (b) "board" shall mean the State Board of Pharmacy. 26 27 (C) "Compounding" means (i) the production, preparation, propagation, conversion or processing of a sterile or 28 nonsterile drug or device either directly or indirectly by 29 H. B. No. 416 G3/5 02/HR07/R924 PAGE 1 (RF\HS)

extraction from substances of natural origin or independently by 30 31 means of chemical or biological synthesis or from bulk chemicals or the preparation, mixing, measuring, assembling, packaging or 32 33 labeling of a drug or device as a result of a practitioner's 34 prescription drug order or initiative based on the 35 practitioner/patient/pharmacist relationship in the course of professional practice, or (ii) for the purpose of, as an incident 36 to, research, teaching or chemical analysis and not for sale or 37 dispensing. Compounding also includes the preparation of drugs or 38 devices in anticipation of prescription drug orders based on 39 40 routine regularly observed prescribing patterns.

(d) "Continuing education unit" shall mean ten (10)
clock hours of study or other such activity as may be approved by
the board, including, but not limited to, all programs which have
been approved by the American Council on Pharmaceutical Education.

45 (e) "Deliver" or "delivery" shall mean the actual,
46 constructive or attempted transfer of a drug or device from one
47 person to another, whether or not for a consideration.

(f) "Device" shall mean an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent or other similar or related article, including any component part or accessory which is required under federal or state law to be prescribed by a practitioner and dispensed by a pharmacist.

(g) "Dispense" or "dispensing" shall mean the interpretation of a valid prescription, order of a practitioner by a pharmacist and the subsequent preparation of the drug or device for administration to or use by a patient or other individual entitled to receive the drug.

(h) "Distribute" shall mean the delivery of a drug or
device other than by administering or dispensing to persons other
than the ultimate consumer.

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(i) "Drug" shall mean:

H. B. No. 416 02/HR07/R924 PAGE 2 (RF\HS) (i) Articles recognized as drugs in the official
United States Pharmacopeia, official National Formulary, official
Homeopathic Pharmacopeia, other drug compendium or any supplement
to any of them;

(ii) Articles intended for use in the diagnosis,
cure, mitigation, treatment or prevention of disease in man or
other animals;

(iii) Articles other than food intended to affect
the structure or any function of the body of man or other animals;
and

(iv) Articles intended for use as a component of
any articles specified in subparagraph (i), (ii) or (iii) of this
paragraph.

(j) "Drugroom" shall mean a business, which does not require the services of a pharmacist, where prescription drugs or prescription devices are bought, sold, maintained or provided to consumers.

(k) "Extern" shall mean a student in the professional program of a school of pharmacy accredited by the American Council on Pharmaceutical Education who is making normal progress toward completion of a professional degree in pharmacy.

83 (1)"Foreign pharmacy graduate" shall mean a person whose undergraduate pharmacy degree was conferred by a recognized 84 school of pharmacy outside of the United States, the District of 85 86 Columbia and Puerto Rico. Recognized schools of pharmacy are those colleges and universities listed in the World Health 87 Organization's World Directory of Schools of Pharmacy, or 88 otherwise approved by the Foreign Pharmacy Graduate Examination 89 Committee (FPGEC) certification program as established by the 90 National Association of Boards of Pharmacy. 91

92 (m) "Generic equivalent drug product" shall mean a drug 93 product which (i) contains the identical active chemical 94 ingredient of the same strength, quantity and dosage form; (ii) is

H. B. No. 416 02/HR07/R924 PAGE 3 (RF\HS) 95 of the same generic drug name as determined by the United States 96 Adoptive Names and accepted by the United States Food and Drug 97 Administration; and (iii) conforms to such rules and regulations 98 as may be adopted by the board for the protection of the public to 99 assure that such drug product is therapeutically equivalent.

(n) "Interested directly" shall mean being employed by,
having full or partial ownership of, or control of, any facility
permitted or licensed by the Mississippi State Board of Pharmacy.

(o) "Interested indirectly" shall mean having a spouse
who is employed by any facility permitted or licensed by the
Mississippi State Board of Pharmacy.

106 (p) "Intern" shall mean a person who has graduated from 107 a school of pharmacy but has not yet become licensed as a 108 pharmacist.

(q) "Manufacturer" shall mean a person, business or other entity engaged in the production, preparation, propagation, conversion or processing of a prescription drug or device, if such actions are associated with promotion and marketing of such drugs or devices.

(r) "Manufacturer's distributor" shall mean any person or business who is not an employee of a manufacturer, but who distributes sample drugs or devices, as defined under subsection (i) of this section, under contract or business arrangement for a manufacturer to practitioners.

119 (s) "Manufacturing" of prescription products shall mean the production, preparation, propagation, conversion or processing 120 121 of a drug or device, either directly or indirectly, by extraction from substances from natural origin or independently by means of 122 chemical or biological synthesis, or from bulk chemicals and 123 includes any packaging or repackaging of the substance(s) or 124 labeling or relabeling of its container, if such actions are 125 126 associated with promotion and marketing of such drug or devices.

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(t) "Misappropriation of a prescription drug" shall mean to illegally or unlawfully convert a drug, as defined in subsection (i) of this section, to one's own use or to the use of another.

(u) "Nonprescription drugs" shall mean nonnarcotic medicines or drugs that may be sold without a prescription and are prepackaged and labeled for use by the consumer in accordance with the requirements of the statutes and regulations of this state and the federal government.

136 (v) "Person" shall mean an individual, corporation,137 partnership, association or any other legal entity.

(w) "Pharmacist" shall mean an individual health care provider licensed by this state to engage in the practice of pharmacy. This recognizes a pharmacist as a learned professional who is authorized to provide patient services.

142 (x) "Pharmacy" shall mean any location for which a 143 pharmacy permit is required and in which prescription drugs are 144 maintained, compounded and dispensed for patients by a pharmacist. 145 This definition includes any location where pharmacy-related 146 services are provided by a pharmacist.

(y) "Prepackaging" shall mean the act of placing small precounted quantities of drug products in containers suitable for dispensing or administering in anticipation of prescriptions or orders.

(z) Unlawful or unauthorized "possession" shall mean
physical holding or control by a pharmacist of a controlled
substance outside the usual and lawful course of employment.

(aa) "Practice of pharmacy" shall mean a health care
service that includes, but is not limited to, the compounding,
dispensing, and labeling of drugs or devices; interpreting and
evaluating prescriptions; administering and distributing drugs and
devices; the compounding, dispensing and labeling of drugs and
devices; maintaining prescription drug records; advising and

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consulting concerning therapeutic values, content, hazards and 160 161 uses of drugs and devices; initiating or modifying of drug therapy 162 in accordance with written guidelines or protocols previously 163 established and approved by the board; selecting drugs; 164 participating in drug utilization reviews; storing prescription 165 drugs and devices; ordering lab work in accordance with written guidelines or protocols as defined by paragraph (jj) of this 166 section; providing pharmacotherapeutic consultations; supervising 167 168 supportive personnel and such other acts, services, operations or transactions necessary or incidental to the conduct of the 169 170 foregoing.

(bb) "Practitioner" shall mean a physician, dentist,
veterinarian, or other health care provider authorized by law to
diagnose and prescribe drugs.

174 (cc) "Prescription" shall mean a written, verbal or 175 electronically transmitted order issued by a practitioner for a 176 drug or device to be dispensed for a patient by a pharmacist.

(dd) "Prescription drug" or "legend drug" shall mean a drug which is required under federal law to be labeled with either of the following statements prior to being dispensed or delivered:

180 (i) "Caution: Federal law prohibits dispensing181 without prescription," or

(ii) "Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian"; or a drug which is required by any applicable federal or state law or regulation to be dispensed on prescription only or is restricted to use by practitioners only.

(ee) "Product selection" shall mean the dispensing of a
generic equivalent drug product in lieu of the drug product
ordered by the prescriber.

(ff) "Provider" or "primary health care provider" shall include a pharmacist who provides health care services within his or her scope of practice pursuant to state law and regulation.

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"Registrant" shall mean a pharmacy or other entity 193 (gg) 194 which is registered with the Mississippi State Board of Pharmacy 195 to buy, sell or maintain controlled substances.

196 (hh) "Repackager" means a person registered by the 197 Federal Food and Drug Administration as a repackager who removes a 198 prescription drug product from its marketed container and places it into another, usually of smaller size, to be distributed to 199 persons other than the consumer. 200

201 (ii) "Supportive personnel" or "pharmacist technician" shall mean those individuals utilized in pharmacies whose 202 203 responsibilities are to provide nonjudgmental technical services concerned with the preparation and distribution of drugs under the 204 205 direct supervision and responsibility of a pharmacist.

206 "Written guideline or protocol" shall mean an (jj) 207 agreement in which any practitioner authorized to prescribe drugs 208 delegates to a pharmacist authority to conduct specific prescribing functions in an institutional setting, or with 209 210 individual patients, provided that a specific protocol agreement is signed on each patient and is filed as required by law or by 211 212 rule or regulation of the board.

(kk) "Wholesaler" shall mean a person who buys or 213 214 otherwise acquires prescription drugs or prescription devices for resale or distribution, or for repackaging for resale or 215 distribution, to persons other than consumers. 216

217 SECTION 4. Section 73-21-75, Mississippi Code of 1972, is reenacted as follows: 218

219 73-21-75. (1) The State Board of Pharmacy created by former Section 73-21-9 is hereby continued and reconstituted as follows: 220 The board shall consist of seven (7) appointed members. At least 221 one (1) appointment shall be made from each congressional 222 district. Each appointed member of the board shall be appointed 223 224 by the Governor, with the advice and consent of the Senate, from a list of five (5) names submitted by the Mississippi State 225

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Pharmaceutical Association/Mississippi Pharmacists Association. 226 Of the members appointed, one (1) shall, at the time of 227 appointment, have had five (5) years' experience as a pharmacist 228 229 at a facility holding an institutional permit, and one (1) shall, 230 at the time of appointment, have had five (5) years' experience as a pharmacist at a facility holding a retail permit. Any person 231 appointed to the board shall be limited to two (2) full terms of 232 office during any fifteen-year period, including any member 233 serving on May 14, 1992. 234

(2) The members of the board appointed and serving prior to 235 236 July 1, 1983, whose terms have not expired by July 1, 1983, shall serve the balance of their terms as members of the reconstituted 237 board, and they shall be considered to be from the same 238 congressional districts from which they were originally appointed 239 if they still reside therein, even if the district boundaries have 240 241 changed subsequent to their original appointments. The Governor shall appoint the remaining members of the reconstituted board in 242 243 the manner prescribed in subsection (1) of this section on July 1, The initial members of the reconstituted board shall serve 244 1983. 245 terms of office as follows:

(a) The term of the member from the First Congressional
District shall expire on July 1, 1984; and from and after July 1,
1996, this appointment shall be designated as Post 1.

(b) The term of the member from the Second
Congressional District shall expire on July 1, 1988; and from and
after July 1, 1996, this appointment shall be designated as Post
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(c) The term of the member from the Third Congressional
District shall expire on July 1, 1986; and from and after July 1,
1996, this appointment shall be designated as Post 3.

(d) The term of the member from the FourthCongressional District shall expire on July 1, 1985; and from and

H. B. No. 416 02/HR07/R924 PAGE 8 (RF\HS) 258 after July 1, 1996, this appointment shall be designated as Post 259 4.

(e) The term of the member from the Fifth Congressional
District shall expire on July 1, 1987; and from and after July 1,
1996, this appointment shall be designated as Post 5.

(f) The term of one (1) of the members from the state at large shall expire on July 1, 1985; and from and after July 1, 1996, this appointment shall be designated as Post 6.

(g) The term of the other member from the state at
large shall expire on July 1, 1988; and from and after July 1,
1996, this appointment shall be designated as Post 7.

(3) At the expiration of a term, members of the board shall 269 be appointed in the manner prescribed in subsection (1) of this 270 section for terms of five (5) years from the expiration date of 271 272 the previous terms. Any vacancy on the board prior to the expiration of a term for any reason, including resignation, 273 removal, disqualification, death or disability, shall be filled by 274 275 appointment of the Governor in the manner prescribed in subsection 276 (1) of this section for the balance of the unexpired term. The 277 Mississippi State Pharmaceutical Association/Mississippi 278 Pharmacists Association shall submit a list of nominees no more 279 than thirty (30) days after a vacancy occurs, and the Governor 280 shall fill such vacancies within ninety (90) days after each such vacancy occurs. 281

(4) To be qualified to be a member of the board, a personshall:

(a) Be an adult citizen of Mississippi for a period of
at least five (5) years preceding his appointment to the board;
(b) Be a pharmacist licensed and in good standing to
practice pharmacy in the State of Mississippi;

(c) Have at least five (5) years' experience as apharmacist; and

H. B. No. 416 02/HR07/R924 PAGE 9 (RF\HS) (d) Be actively engaged full time in the practice ofpharmacy in Mississippi.

(5) The Governor may remove any or all members of the board 292 293 on proof of unprofessional conduct, continued absence from the 294 state, or for failure to perform the duties of his office. Any member who shall not attend two (2) consecutive meetings of the 295 board for any reason other than illness of such member shall be 296 subject to removal by the Governor. The president of the board 297 shall notify the Governor in writing when any such member has 298 failed to attend two (2) consecutive regular meetings. 299 No removal 300 shall be made without first giving the accused an opportunity to 301 be heard in refutation of the charges made against him, and he 302 shall be entitled to receive a copy of the charges at the time of 303 filing.

304 **SECTION 5.** Section 73-21-77, Mississippi Code of 1972, is 305 reenacted as follows:

306 73-21-77. (1) Each person appointed as a member of the 307 board shall qualify by taking the oath prescribed by the 308 Constitution for the state officers, and shall file certificate 309 thereof in the Office of the Secretary of State within fifteen 310 (15) days after his appointment.

311 (2) There shall be a president of the board and such other 312 officers as deemed necessary by the board elected by and from its 313 membership.

(3) The board shall meet at least once each quarter to
transact business, and may meet at such additional times as it may
deem necessary. Such additional meetings may be called by the
president of the board or a majority of the members of the board.
(4) The place for each meeting shall be determined prior to

319 giving notice of such meeting and shall not be changed after such 320 notice is given without adequate subsequent notice.

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321 (5) A majority of the members of the board shall constitute
322 a quorum for the conduct of the meeting and all actions of the
323 board shall be by a majority.

(6) Each member of the board shall receive a per diem as
provided in Section 25-3-69, not to exceed thirty (30) days in any
one (1) period of twelve (12) months, for each day actually
engaged in meetings of the board, together with necessary
traveling and other expenses as provided in Section 25-3-41.

329 **SECTION 6.** Section 73-21-79, Mississippi Code of 1972, is 330 reenacted as follows:

331 73-21-79. (1) The board shall employ an executive director 332 of the board. The executive director shall be a citizen of 333 Mississippi and a pharmacist licensed and in good standing to 334 practice pharmacy in the State of Mississippi, who has had five 335 (5) years' experience as a pharmacist.

The executive director shall receive a salary to be set 336 (2) by the board, subject to the approval of the State Personnel 337 338 Board, and shall be entitled to necessary expenses incurred in the performance of his official duties. He shall devote full time to 339 340 the duties of his office and shall not be interested directly or 341 indirectly as defined in Section 73-21-73 in the operation of a 342 pharmacy in Mississippi or any other facility permitted by the 343 board or engaged in any other business that will interfere with the duties of his office. 344

345 (3) The duties and responsibilities of the executive
346 director shall be defined by rules and regulations prescribed by
347 the board.

(4) The board may, in its discretion, employ persons in addition to the executive director in such other positions or capacities as it deems necessary to the proper conduct of board business. Any pharmacist-investigator employed by the board may have other part-time employment, provided that he shall not accept any employment that would cause a conflict of interest in his

H. B. No. 416 02/HR07/R924 PAGE 11 (RF\HS) 354 pharmacist-investigator duties. The board may employ legal 355 counsel to assist in the conduct of its business.

356 **SECTION 7.** Section 73-21-81, Mississippi Code of 1972, is 357 reenacted as follows:

358 73-21-81. The responsibility for the enforcement of the provisions of this chapter shall be vested in the board. 359 The 360 board shall have all of the duties, powers and authority specifically granted by and necessary to the enforcement of this 361 362 chapter. The board may make, adopt, amend and repeal such rules and regulations as may be deemed necessary by the board from time 363 364 to time for the proper administration and enforcement of this chapter, in accordance with the provisions of the Mississippi 365 366 Administrative Procedures Law (Section 25-43-1 et seq.).

367 SECTION 8. Section 73-21-83, Mississippi Code of 1972, is
368 reenacted as follows:

(1) The board shall be responsible for the 369 73-21-83. control and regulation of the practice of pharmacy, to include the 370 371 regulation of pharmacy externs or interns and pharmacist technicians, in this state, the regulation of the wholesaler 372 373 distribution of drugs and devices as defined in Section 73-21-73, 374 and the distribution of sample drugs or devices by manufacturer's 375 distributors as defined in Section 73-21-73 by persons other than the original manufacturer or distributor in this state. 376

A license for the practice of pharmacy shall be obtained 377 (2) 378 by all persons prior to their engaging in the practice of pharmacy. However, the provisions of this chapter shall not apply 379 380 to physicians, dentists, veterinarians, osteopaths or other practitioners of the healing arts who are licensed under the laws 381 of the State of Mississippi and are authorized to dispense and 382 383 administer prescription drugs in the course of their professional 384 practice.

385 (3) The initial licensure fee shall be set by the board but386 shall not exceed Two Hundred Dollars (\$200.00).

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All students actively enrolled in a professional school 387 (4) 388 of pharmacy accredited by the American Council on Pharmaceutical Education who are making satisfactory progress toward graduation 389 390 and who act as an extern or intern under the direct supervision of 391 a pharmacist in a location permitted by the Board of Pharmacy must 392 obtain a pharmacy student registration prior to engaging in such activity. The student registration fee shall be set by the board 393 but shall not exceed One Hundred Dollars (\$100.00). 394

395 (5) All persons licensed to practice pharmacy prior to July
396 1, 1991, by the State Board of Pharmacy under Section 73-21-89
397 shall continue to be licensed under the provisions of Section
398 73-21-91.

399 SECTION 9. Section 73-21-85, Mississippi Code of 1972, is
400 reenacted as follows:

401 73-21-85. (1) To obtain a license to engage in the practice 402 of pharmacy by examination, or by score transfer, the applicant 403 shall:

404 (a) Have submitted a written application on the form405 prescribed by the board;

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(b) Be of good moral character;

407 (c) Have graduated from a school or college of pharmacy
408 accredited by the American Council of Pharmaceutical Education and
409 have been granted a pharmacy degree therefrom;

(d) Have successfully passed an examination approved bythe board;

(e) Have paid all fees specified by the board for
examination, not to exceed the cost to the board of administering
the examination;

(f) Have paid all fees specified by the board for licensure; and

417 (g) Have submitted evidence of externship and/or418 internship as specified by the board.

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To obtain a license to engage in the practice of (2) 419 420 pharmacy, a foreign pharmacy graduate applicant shall obtain the National Association of Boards of Pharmacy's Foreign Pharmacy 421 422 Graduate Examination Committee's certification, which shall 423 include, but not be limited to, successfully passing the Foreign Pharmacy Graduate Equivalency Examination and attaining a total 424 score of at least five hundred fifty (550) on the Test of English 425 as a Foreign Language (TOEFL), and shall: 426

427 (a) Have submitted a written application on the form428 prescribed by the board;

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(b) Be of good moral character;

430 (c) Have graduated and been granted a pharmacy degree
431 from a college or school of pharmacy recognized and approved by
432 the National Association of Boards of Pharmacy's Foreign Pharmacy
433 Graduate Examination Committee;

(d) Have paid all fees specified by the board for
examination, not to exceed the cost to the board of administering
the examination;

437 (e) Have successfully passed an examination approved by438 the board;

439 (f) Have completed the number of internship hours as440 set forth by regulations of the board; and

(g) Have paid all fees specified by the board forlicensure.

(3) Each application or filing made under this section shall
include the Social Security number(s) of the applicant in
accordance with Section 93-11-64, Mississippi Code of 1972.

446 **SECTION 10.** Section 73-21-87, Mississippi Code of 1972, is 447 reenacted as follows:

448 73-21-87. (1) To obtain a license to engage in the practice 449 of pharmacy by reciprocity or license transfer, the applicant 450 shall:

H. B. No. 416 02/HR07/R924 PAGE 14 (RF\HS) (a) Have submitted a written application on the formprescribed by the board;

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(b) Be of good moral character;

454 (c) Have possessed at the time of initial licensure as
455 a pharmacist such other qualifications necessary to have been
456 eligible for licensure at that time in that state;

(d) Have presented to the board proof that any license or licenses granted to the applicant by any other states have not been suspended, revoked, cancelled or otherwise restricted for any reason except nonrenewal or the failure to obtain required continuing education credits; and

462 (e) Have paid all fees specified by the board for463 licensure.

464 (2) No applicant shall be eligible for licensure by 465 reciprocity or license transfer or unless the state in which the 466 applicant was initially licensed also grants a reciprocal license 467 or transfer license to pharmacists licensed by this state under 468 like circumstances and conditions.

469 (3) Each application or filing made under this section shall
470 include the Social Security number(s) of the applicant in
471 accordance with Section 93-11-64, Mississippi Code of 1972.

472 **SECTION 11.** Section 73-21-89, Mississippi Code of 1972, is 473 reenacted as follows:

474 73-21-89. (1) The board shall issue a license to practice
475 pharmacy to any person, if such person be otherwise qualified,
476 upon presentation to the board of:

477 (a) Satisfactory proof that the applicant has been
478 graduated from the University of Mississippi School of Pharmacy;
479 (b) Written application for licensure; and
480 (c) Payment of all fees specified by the board for

481 licensure.

482 (2) The board shall not issue any new licenses pursuant to483 this section after June 30, 1987.

H. B. No. 416 02/HR07/R924 PAGE 15 (RF\HS) 484 (3) Each application or filing made under this section shall
485 include the Social Security number(s) of the applicant in
486 accordance with Section 93-11-64, Mississippi Code of 1972.
487 SECTION 12. Section 73-21-91, Mississippi Code of 1972, is

488 reenacted as follows:

489 73-21-91. (1) Every pharmacist shall renew his license490 biennially. To renew his license, a pharmacist shall:

491 (a) Submit an application for renewal on the form492 prescribed by the board;

(b) Submit satisfactory evidence of the completion in the last licensure period of such continuing education units as shall be required by the board, but in no case less than two (2) continuing education units in the last licensure period;

497 Pay such renewal fees as required by the board, not (C) to exceed Two Hundred Dollars (\$200.00) for each biennial 498 499 licensing period, provided that the board may add a surcharge of not more than Five Dollars (\$5.00) to a license renewal fee to 500 501 fund a program to aid impaired pharmacists or pharmacy students. Any pharmacist license renewal received postmarked after December 502 503 31 of the renewal period will be returned and a Fifty Dollar 504 (\$50.00) late renewal fee will be assessed prior to renewal.

505 (2) Any pharmacist who has defaulted in license renewal may be reinstated within two (2) years upon payment of renewal fees in 506 arrears and presentation of evidence of the required continuing 507 508 education. Any pharmacist defaulting in license renewal for a period in excess of two (2) years shall be required to 509 successfully complete the examination given by the board pursuant 510 to Section 73-21-85 before being eligible for reinstatement as a 511 pharmacist in Mississippi, or shall be required to appear before 512 513 the board to be examined for his competence and knowledge of the practice of pharmacy, and may be required to submit evidence of 514 515 continuing education. If such person is found fit by the board to practice pharmacy in this state, the board may reinstate his 516

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519 (3) Each application or filing made under this section shall
520 include the Social Security number(s) of the applicant in
521 accordance with Section 93-11-64, Mississippi Code of 1972.

522 **SECTION 13.** Section 73-21-93, Mississippi Code of 1972, is 523 reenacted as follows:

524 73-21-93. (1) The examination for licensure required under 525 Section 73-21-85 shall be given by the board at least once during 526 each year. The board shall determine the content and subject 527 matter of each examination, the place, time and date of the 528 administration of the examination and those persons who have 529 successfully passed the examination.

(2) The examination shall be prepared to measure the
competence of the applicant to engage in the practice of pharmacy.
The board may employ and cooperate with any organization or
consultant in the preparation and grading of an appropriate
examination, but shall retain the sole discretion and
responsibility of determining which applicants have successfully
passed such an examination.

537 (3) The board shall have authority to use the laboratories 538 of the school of pharmacy and other facilities of the University 539 of Mississippi for the purpose of examining applicants.

540 **SECTION 14.** Section 73-21-95, Mississippi Code of 1972, is 541 reenacted as follows:

542 73-21-95. The assistant pharmacist license is hereby 543 abolished after April 30, 1984. The board shall issue a license 544 to practice pharmacy to those persons presently holding an 545 assistant pharmacist license upon their meeting the requirements 546 of Section 73-21-91.

547 **SECTION 15.** Section 73-21-97, Mississippi Code of 1972, is 548 reenacted as follows:

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73-21-97. (1) The board may refuse to issue or renew, or 549 may suspend, reprimand, revoke or restrict the license, 550 registration or permit of any person upon one or more of the 551 552 following grounds: 553 (a) Unprofessional conduct as defined by the rules and regulations of the board; 554 (b) 555 Incapacity of a nature that prevents a pharmacist 556 from engaging in the practice of pharmacy with reasonable skill, 557 confidence and safety to the public; Being found guilty by a court of competent 558 (C) jurisdiction of one or more of the following: 559 560 (i) A felony; (ii) Any act involving moral turpitude or gross 561 562 immorality; or Violation of pharmacy or drug laws of this 563 (iii) 564 state or rules or regulations pertaining thereto, or of statutes, rules or regulations of any other state or the federal government; 565 566 (d) Fraud or intentional misrepresentation by a 567 licensee or permit holder in securing the issuance or renewal of a 568 license or permit; Engaging or aiding and abetting an individual to 569 (e) 570 engage in the practice of pharmacy without a license; Violation of any of the provisions of this chapter 571 (f) or rules or regulations adopted pursuant to this chapter; 572 573 (q) Failure to comply with lawful orders of the board; 574 Negligently or willfully acting in a manner (h) inconsistent with the health or safety of the public; 575 576 Addiction to or dependence on alcohol or controlled (i) substances or the unauthorized use or possession of controlled 577 578 substances; Misappropriation of any prescription drug; 579 (j)

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(k) Being found guilty by the licensing agency in
another state of violating the statutes, rules or regulations of
that jurisdiction; or

583 (1) The unlawful or unauthorized possession of a584 controlled substance.

585 (2) In lieu of suspension, revocation or restriction of a 586 license as provided for above, the board may warn or reprimand the 587 offending pharmacist.

In addition to the grounds specified in subsection (1) 588 (3) of this section, the board shall be authorized to suspend the 589 590 license, registration or permit of any person for being out of compliance with an order for support, as defined in Section 591 93-11-153. The procedure for suspension of a license, 592 593 registration or permit for being out of compliance with an order for support, and the procedure for the reissuance or reinstatement 594 of a license, registration or permit suspended for that purpose, 595 and the payment of any fees for the reissuance or reinstatement of 596 597 a license, registration or permit suspended for that purpose, shall be governed by Section 93-11-157 or 93-11-163, as the case 598 599 may be. If there is any conflict between any provision of Section 600 93-11-157 or 93-11-163 and any provision of this chapter, the 601 provisions of Section 93-11-157 or 93-11-163, as the case may be, shall control. 602

603 **SECTION 16.** Section 73-21-99, Mississippi Code of 1972, is 604 reenacted as follows:

73-21-99. (1) Disciplinary action by the board against a
licensee, registrant or permit holder, or license, registration or
permit shall require the following:

(a) A sworn affidavit filed with the board charging a
licensee or permit holder with an act which is grounds for
disciplinary action as provided in Section 73-21-97; and

(b) An order of the Investigations Review Committee ofthe board which shall cause the executive director of the board to

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fix a time and place for a hearing by the board. The executive 613 director shall cause a written notice specifying the offense or 614 offenses for which the licensee or permit holder is charged and 615 616 notice of the time and place of the hearing to be served upon the 617 licensee or permit holder at least thirty (30) days prior to the hearing date. Such notice may be served by mailing a copy thereof 618 by certified mail, postage prepaid, to the last known residence or 619 620 business address of the licensee or permit holder.

The board shall designate two (2) of its members to 621 (2) serve on a rotating no longer than three-consecutive-month basis 622 623 with the executive director and legal counsel for the board as an 624 Investigations Review Committee, and the board's investigators shall provide status reports solely to the Investigations Review 625 Committee during monthly meetings of the board. 626 Such reports 627 shall be made on all on-going investigations, and shall apply to any routine inspections which may give rise to the filing of a 628 complaint. In the event any complaint on a licensee comes before 629 630 the board for possible disciplinary action, the members of the board serving on the Investigations Review Committee which 631 632 reviewed the investigation of such complaint shall recuse themselves and not participate in the disciplinary proceeding. 633

(3) The board acting by and through its Investigation Review
Committee may, if deemed necessary, issue a letter of reprimand to
any licensee, registrant or permit holder in lieu of formal action
by the board.

(4) The board, acting by and through its executive director,
is hereby authorized and empowered to issue subpoenas for the
attendance of witnesses and the production of books and papers at
such hearing. Process issued by the board shall extend to all
parts of the state and shall be served by any person designated by
the board for such service.

644 (5) The accused shall have the right to appear either645 personally or by counsel or both to produce witnesses or evidence

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646 in his behalf, to cross-examine witnesses and to have subpoenas 647 issued by the board.

(6) At the hearing, the board shall administer oaths as may be necessary for the proper conduct of the hearing. All hearings shall be conducted by the board, which shall not be bound by strict rules of procedure or by the laws of evidence in the conduct of its proceedings, but the determination shall be based upon sufficient evidence to sustain it.

Where, in any proceeding before the board, any witness 654 (7) fails or refuses to attend upon a subpoena issued by the board, 655 656 refuses to testify, or refuses to produce any books and papers the production of which is called for by a subpoena, the attendance of 657 such witness, the giving of his testimony or the production of the 658 659 books and papers shall be enforced by any court of competent 660 jurisdiction of this state in the manner provided for the enforcement of attendance and testimony of witnesses in civil 661 cases in the courts of this state. 662

(8) The board shall, within thirty (30) days after
conclusion of the hearing, reduce its decision to writing and
forward an attested true copy thereof to the last known residence
or business address of such licensee or permit holder by way of
United States first class, certified mail, postage prepaid.

668 **SECTION 17.** Section 73-21-101, Mississippi Code of 1972, is 669 reenacted as follows:

670 73-21-101. (1) The right to appeal from the action of the board in denying, revoking, suspending or refusing to renew any 671 license, registration or permit issued by the board, or fining or 672 otherwise disciplining any person is hereby granted. Such appeal 673 shall be to the chancery court of the county of the residence of 674 675 the licensee or permit holder on the record made, including a verbatim transcript of the testimony at the hearing. The appeal 676 677 shall be taken within thirty (30) days after notice of the action 678 of the board in denying, revoking, suspending or refusing to renew

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the license or permit, or fining or otherwise disciplining the 679 The appeal shall be perfected upon filing notice of the 680 person. appeal and by the prepayment of all costs, including the cost of 681 682 the preparation of the record of the proceedings by the board, and 683 the filing of a bond in the sum of Two Hundred Dollars (\$200.00), conditioned that if the action of the board in denying, revoking, 684 685 suspending or refusing to renew the license or permit, or fining or otherwise disciplining the person, be affirmed by the chancery 686 court, the licensee or permit holder will pay the costs of the 687 appeal and the action in the chancery court. 688

689 (2) If there is an appeal, such appeal shall act as a 690 The chancery court shall dispose of the appeal and supersedeas. 691 enter its decision promptly. The hearing on the appeal may, in 692 the discretion of the chancellor, be tried in vacation. The scope of review of the chancery court shall be limited to a review of 693 the record made before the board to determine if the action of the 694 board is unlawful for the reason that it was (a) not supported by 695 696 substantial evidence, (b) arbitrary or capricious, (c) beyond the 697 power of the board to make, or (d) in violation of some statutory 698 or constitutional right of the appellant. The decision of the chancery court may be appealed to the Supreme Court in the manner 699 700 provided by law.

Actions taken by the board in suspending a license, 701 (3) registration or permit when required by Section 93-11-157 or 702 703 93-11-163 are not actions from which an appeal may be taken under 704 this section. Any appeal of a suspension of a license, registration or permit that is required by Section 93-11-157 or 705 706 93-11-163 shall be taken in accordance with the appeal procedure specified in Section 93-11-157 or 93-11-163, as the case may be, 707 708 rather than the procedure specified in this section.

709 **SECTION 18.** Section 73-21-103, Mississippi Code of 1972, is 710 reenacted as follows:

H. B. No. 416 02/HR07/R924 PAGE 22 (RF\HS) 711 73-21-103. (1) Upon the finding of the existence of grounds 712 for action against any permitted facility or discipline of any 713 person holding a license, registration or permit, seeking a 714 license, registration or permit, or seeking to renew a license or 715 permit under the provisions of this chapter, the board may impose 716 one or more of the following penalties:

717 (a) Suspension of the offender's license, registration718 and/or permit for a term to be determined by the board;

719 (b) Revocation of the offender's license, registration 720 and/or permit;

(c) Restriction of the offender's license, registration and/or permit to prohibit the offender from performing certain acts or from engaging in the practice of pharmacy in a particular manner for a term to be determined by the board;

(d) Imposition of a monetary penalty as follows:
(i) For the first violation, a monetary penalty of
not less than Fifty Dollars (\$50.00) nor more than Five Hundred
Dollars (\$500.00) for each violation;

(ii) For the second violation and subsequent violations, a monetary penalty of not less than One Hundred Dollars (\$100.00) nor more than One Thousand Dollars (\$1,000.00) for each violation.

Money collected by the board under Section 73-21-103, paragraphs (1)(d)(i), (ii) and (iv) shall be deposited to the credit of the State General Fund of the State Treasury;

(iii) The board may assess a monetary penalty for those reasonable costs that are expended by the board in the investigation and conduct of a proceeding for licensure revocation, suspension or restriction, including but not limited to the cost of process service, court reporters, expert witnesses and investigators.

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Money collected by the board under Section 73-21-103, paragraph (1)(d)(iii), shall be deposited to the credit of the Special Fund of the Pharmacy Board;

(iv) The board may impose a monetary penalty for those facilities/businesses registered with the Pharmacy Board as wholesalers/manufacturers of not less than One Hundred Dollars (\$100.00) per violation and not more than Twenty-five Thousand Dollars (\$25,000.00) per violation;

750 (e) Refusal to renew offender's license, registration751 and/or permit;

(f) Placement of the offender on probation and
supervision by the board for a period to be determined by the
board;

755

(g) Public or private reprimand.

Whenever the board imposes any penalty under this subsection, the board may require rehabilitation and/or additional education as the board may deem proper under the circumstances, in addition to the penalty imposed.

760 Any person whose license, registration and/or permit has (2) 761 been suspended, revoked or restricted pursuant to this chapter, 762 whether voluntarily or by action of the board, shall have the 763 right to petition the board at reasonable intervals for reinstatement of such license, registration and/or permit. 764 Such 765 petition shall be made in writing and in the form prescribed by 766 the board. Upon investigation and hearing, the board may, in its 767 discretion, grant or deny such petition, or it may modify its 768 original finding to reflect any circumstances which have changed sufficiently to warrant such modifications. The procedure for the 769 reinstatement of a license, registration or permit that is 770 suspended for being out of compliance with an order for support, 771 as defined in Section 93-11-153, shall be governed by Section 772 773 93-11-157 or 93-11-163, as the case may be.

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(3) Nothing herein shall be construed as barring criminal
prosecutions for violation of this chapter where such violations
are deemed as criminal offenses in other statutes of this state or
of the United States.

(4) A monetary penalty assessed and levied under this section shall be paid to the board by the licensee, registrant or permit holder upon the expiration of the period allowed for appeal of such penalties under Section 73-21-101, or may be paid sooner if the licensee, registrant or permit holder elects.

783 When payment of a monetary penalty assessed and levied (5) 784 by the board against a licensee, registrant or permit holder in 785 accordance with this section is not paid by the licensee, registrant or permit holder when due under this section, the board 786 787 shall have the power to institute and maintain proceedings in its 788 name for enforcement of payment in the chancery court of the county and judicial district of residence of the licensee, 789 registrant or permit holder, or if the licensee, registrant or 790 permit holder is a nonresident of the State of Mississippi, in the 791 792 Chancery Court of the First Judicial District of Hinds County, 793 Mississippi. When such proceedings are instituted, the board 794 shall certify the record of its proceedings, together with all documents and evidence, to the chancery court and the matter shall 795 thereupon be heard in due course by the court, which shall review 796 the record and make its determination thereon. The hearing on the 797 798 matter may, in the discretion of the chancellor, be tried in 799 vacation.

(6) The board shall develop and implement a uniform penalty
policy which shall set the minimum and maximum penalty for any
given violation of board regulations and laws governing the
practice of pharmacy. The board shall adhere to its uniform
penalty policy except in such cases where the board specifically
finds, by majority vote, that a penalty in excess of, or less
than, the uniform penalty is appropriate. Such vote shall be

H. B. No. 416 02/HR07/R924 PAGE 25 (RF\HS) 807 reflected in the minutes of the board and shall not be imposed 808 unless such appears as having been adopted by the board.

809 **SECTION 19.** Section 73-21-105, Mississippi Code of 1972, is 810 reenacted as follows:

811 73-21-105. (1) Every facility/business that shall engage in the wholesale distribution of prescription drugs, to include 812 without limitation, manufacturing in this state, distribution into 813 this state, or selling or offering to sell in this state, or 814 distribution from or within this state, shall register biennially 815 with the Mississippi State Board of Pharmacy by applying for a 816 817 permit on a form supplied by the board and accompanied by a fee as set by subsection (4) of this section. The Pharmacy Board shall 818 819 by regulation determine the classification of permit(s) that shall 820 be required.

Every business/facility/pharmacy located in this state (2) 821 822 that engages in or proposes to engage in the dispensing and delivery of prescription drugs to consumers shall register with 823 824 the Mississippi State Board of Pharmacy by applying for a permit on a form supplied by the board and accompanied by a fee as set by 825 subsection (4) of this section. The Pharmacy Board shall by 826 regulation determine the classification of permit(s) that shall be 827 required. 828

(3) The board shall establish by rule or regulation the criteria which each business shall meet to qualify for a permit in each classification. The board shall issue a permit to any applicant who meets the criteria as established. The board may issue various types of permits with varying restrictions to businesses where the board deems it necessary by reason of the type of activities conducted by the business requesting a permit.

(4) The board shall specify by rule or regulation the
registration procedures to be followed, including, but not limited
to, specification of forms for use in applying for such permits
and times, places and fees for filing such applications. However,

H. B. No. 416 02/HR07/R924 PAGE 26 (RF\HS) 840 the biennial fee for an original or renewal permit shall not 841 exceed Three Hundred Dollars (\$300.00).

842 (5) Applications for permits shall include the following843 information about the proposed business:

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(a) Ownership;

845

(b) Location;

(c) Identity of the responsible person or pharmacist licensed to practice in the state, who shall be the pharmacist in charge of the pharmacy, where one is required by this chapter, and such further information as the board may deem necessary.

850 (6) Permits issued by the board pursuant to this section851 shall not be transferable or assignable.

The board shall specify by rule or regulation minimum 852 (7) 853 standards for the responsibility in the conduct of any 854 business/facility and/or pharmacy that has been issued a permit. The board is specifically authorized to require that the portion 855 of the facility located in this state to which a pharmacy permit 856 857 applies be operated only under the direct supervision of no less 858 than one (1) pharmacist licensed to practice in this state, and to 859 provide such other special requirements as deemed necessary. Nothing in this subsection shall be construed to prevent any 860 861 person from owning a pharmacy.

862 (8) All businesses permitted by the board shall report to863 the board the occurrence of any of the following changes:

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(a) Permanent closing;

865 (b) Change of ownership, management, location or866 pharmacist in charge;

867 (c) Any and all other matters and occurrences as the 868 board may require by rule or regulation.

869 (9) Disasters, accidents and emergencies which may affect
870 the strength, purity or labeling of drugs, medications, devices or
871 other materials used in the diagnosis or the treatment of injury,
872 illness and disease shall be immediately reported to the board.

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(10) No business that is required to obtain a permit shall 873 be operated until a permit has been issued for such business by 874 the board. Any person, firm or corporation violating any of the 875 876 provisions of this section shall be quilty of a misdemeanor and, 877 upon conviction thereof, shall be punished by a fine of not less than One Hundred Dollars (\$100.00) nor more than One Thousand 878 Dollars (\$1,000.00), or imprisonment in the county jail for not 879 880 less than thirty (30) days nor more than ninety (90) days, or by 881 both such fine and imprisonment. However, the provisions of this chapter shall not apply to physicians, dentists, veterinarians, 882 883 osteopaths or other practitioners of the healing arts who are 884 licensed under the laws of the State of Mississippi and are 885 authorized to dispense and administer prescription drugs in the 886 course of their professional practice.

887 SECTION 20. Section 73-21-106, Mississippi Code of 1972, is 888 reenacted as follows:

889 73-21-106. (1) Any pharmacy located outside this state that 890 ships, mails or delivers, in any manner, controlled substances, 891 prescription or legend drugs or devices into this state shall be 892 considered a nonresident pharmacy, shall be registered with the 893 board, and shall disclose to the board all of the following:

(a) The location, names, and titles of all principal
corporate officers and all pharmacists-in-charge. A report
containing this information shall be made on a biennial basis and
within thirty (30) days after any change of office, corporate
officer or pharmacist-in-charge;

That it complies with all lawful directions and 899 (b) 900 requests for information from the regulatory or licensing agency of the state in which it is licensed as well as with all requests 901 902 for information made by the board pursuant to this section. The nonresident pharmacy shall maintain at all times a valid unexpired 903 904 license, permit or registration to conduct the pharmacy in 905 compliance with the laws of the state in which it is a resident.

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As a prerequisite to registering with the board, the nonresident pharmacy shall submit a copy of the most recent inspection report resulting from an inspection conducted by the regulatory or licensing agency of the state in which it is located; and

910 (c) That it maintains its records of controlled 911 substances, or prescription or legend drugs or devices dispensed 912 to patients in this state so that the records are readily 913 retrievable from the records of other drugs dispensed.

Any pharmacy subject to this section shall provide 914 (2) during its regular hours of operation, but not less than six (6) 915 916 days per week and for a minimum of forty (40) hours per week, a 917 toll-free telephone service to facilitate communication between patients in this state and a pharmacist at the pharmacy who has 918 access to the patient's records. This toll-free number shall be 919 920 disclosed on a label affixed to each container of drugs dispensed 921 to patients in this state.

(3) The registration fee for nonresident pharmacies shall be
the same as the fee as set by subsection (4) of Section 73-21-105.
(4) The registration requirements of this section shall
apply only to a nonresident pharmacy that only ships, mails or
delivers controlled substances, prescription or legend drugs and

928 (5) The board may deny, revoke or suspend a nonresident 929 pharmacy registration only for:

devices into this state pursuant to a prescription.

930 (a) Failure to comply with any requirement of this931 section; or

(b) Conduct that causes serious bodily or serious psychological injury to a resident of this state if the board has referred the matter to the regulatory or licensing agency in the state in which the pharmacy is located and the regulatory or licensing agency fails to initiate an investigation within forty-five (45) days of the referral.

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938 (6) It is unlawful for any nonresident pharmacy that is not 939 registered pursuant to this section to advertise its services in 940 this state, or for any person who is a resident of this state to 941 advertise the pharmacy services of a nonresident pharmacy that has 942 not registered with the board, with the knowledge that the 943 advertisement will or is likely to induce members of the public in 944 this state to use the pharmacy to fill prescriptions.

945 (7) When requested to do so by the board, each nonresident 946 pharmacy shall supply any inspection reports, controlled 947 substances dispensing records, warning notices, notice of 948 deficiency reports or any other related reports from the state in 949 which it is located concerning the operation of a nonresident 950 pharmacy for review of compliance with state and federal drug 951 laws.

952 **SECTION 21.** Section 73-21-107, Mississippi Code of 1972, is 953 reenacted as follows:

954 73-21-107. (1) The board or its representative may enter 955 and inspect, during reasonable hours, a facility which has 956 obtained or applied for a permit under Section 73-21-105 relative 957 to the following:

958

(a) Drug storage and security;

959 (b) Equipment;

960 (c) Sanitary conditions; or

961 (d) Records, reports, or other documents required to be
962 kept or made under this chapter or the Uniform Controlled
963 Substances Law (Section 41-29-101 et seq.) or rules and
964 regulations adopted under such laws.

965 (2) Prior to an entry and inspection, the board 966 representative shall state his purpose and present appropriate 967 credentials to the owner, pharmacist or agent in charge of a 968 facility.

969 (3) The board representative may:

H. B. No. 416 02/HR07/R924 PAGE 30 (RF\HS) 970 (a) Inspect and copy records, reports, and other
971 documents required to be kept or made under this chapter, the
972 Uniform Controlled Substances Law, or rules and regulations
973 adopted under such laws;

974 (b) Inspect, within reasonable limits and in a
975 reasonable manner, a facility's storage, equipment, security,
976 records, or prescription drugs or devices; or

977 (c) Inventory any stock of any prescription drugs or978 devices in the facility.

979 (4) Unless the owner, pharmacist, or agent in charge of the 980 facility consents in writing, an inspection authorized by this 981 section may not extend to:

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(a) Financial data;

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(b) Sales data other than shipment data; or

984 (c) Pricing data.

985 SECTION 22. Section 73-21-109, Mississippi Code of 1972, is 986 reenacted as follows:

987 73-21-109. No person shall make use of the terms "drugstore," "pharmacy," "apothecary" or words of similar meaning 988 which indicate that pharmaceutical services are performed in any 989 990 sign, letterhead or advertisement unless such person is a permit holder as provided in Section 73-21-105. Any person violating 991 this section shall be guilty of a misdemeanor and, upon conviction 992 thereof, shall be punished by a fine of not less than One Hundred 993 994 Dollars (\$100.00) nor more than Three Hundred Dollars (\$300.00), or by imprisonment in the county jail for not less than thirty 995 (30) days nor more than ninety (90) days, or by both. 996

997 SECTION 23. Section 73-21-111, Mississippi Code of 1972, is
998 reenacted as follows:

999 73-21-111. The board shall make, adopt, amend and repeal 1000 from time to time such rules and regulations for the regulation of 1001 supportive personnel as may be deemed necessary by the board.

H. B. No. 416 02/HR07/R924 PAGE 31 (RF\Hs) 1002 **SECTION 24.** Section 73-21-113, Mississippi Code of 1972, is 1003 reenacted as follows:

1004 73-21-113. All fees received by the board from examinations, 1005 licenses, permits and monetary penalties, and any other funds 1006 received by the board, shall be paid to the State Treasurer, who 1007 shall issue receipts therefor and deposit such funds in the State 1008 Treasury in a special fund to the credit of the board. All such 1009 funds shall be expended only pursuant to appropriation approved by 1010 the Legislature and as provided by law.

1011 SECTION 25. Section 73-21-115, Mississippi Code of 1972, is 1012 reenacted as follows:

73-21-115. (1) Every prescription written in this state by 1013 1014 a person authorized to issue such prescription shall be on prescription forms containing two (2) lines for the prescriber's 1015 signature. There shall be a signature line in the lower 1016 right-hand corner of the prescription form beneath which shall be 1017 clearly imprinted the words "substitution permissible." 1018 There 1019 shall be a signature line in the lower left-hand corner of the prescription form beneath which shall be clearly imprinted the 1020 words "dispense as written." The prescriber's signature on either 1021 signature line shall validate the prescription and shall designate 1022 1023 approval or disapproval of product selection.

1024 (2) If a prescription form which does not contain the two
1025 (2) signature lines required in subsection (1) of this section is
1026 utilized by the prescriber, he shall write in his own handwriting
1027 the words "dispense as written" thereupon to prevent product
1028 selection.

(3) A pharmacist licensed by the Mississippi State Board of
Pharmacy may dispense a one-time emergency dispensing of a
prescription of up to a seventy-two-hour supply of a prescribed
medication in the event the pharmacist is unable to contact the
prescriber to obtain refill authorization, provided that:

The prescription is not for a controlled substance;

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(a)

1034

1035 (b) In the pharmacist's professional judgment, the 1036 interruption of therapy might reasonably produce undesirable 1037 health consequences or may cause physical or mental discomfort;

1038 (c) The dispensing pharmacist notifies the prescriber
1039 or his agent of the emergency dispensing within seven (7) working
1040 days after the one-time emergency dispensing;

1041 (d) The pharmacist properly records the dispensing as a 1042 separate nonrefillable prescription. Said document shall be filed 1043 as is required of all other prescription records. This document 1044 shall be serially numbered and contain all information required of 1045 other prescriptions. In addition it shall contain the number of 1046 the prescription from which it was refilled; and

1047 (e) The pharmacist shall record on the new document the 1048 circumstances which warrant this emergency dispensing.

1049This emergency dispensing shall be done only in the permitted1050facility which contains the nonrefillable prescription.

1051 SECTION 26. Section 73-21-117, Mississippi Code of 1972, is
1052 reenacted as follows:

1053 73-21-117. (1) A pharmacist may select a generic equivalent 1054 drug product only when such selection results in lower cost to the 1055 purchaser, unless product selection is expressly prohibited by the 1056 prescriber.

1057 (2) A pharmacist shall select a generic equivalent drug1058 product when:

1059 (a) The purchaser requests the selection of a generic1060 equivalent drug product;

1061 (b) The prescriber has not expressly prohibited product1062 selection; and

1063 (c) Product selection will result in lower cost to the 1064 purchaser.

1065Before product selection is made, the pharmacist shall advise1066the purchaser of his prerogatives under this subsection.

H. B. No. 416 02/HR07/R924 PAGE 33 (RF\HS) 1067 (3) When requested by the purchaser to dispense the drug 1068 product as ordered by the prescriber, a pharmacist shall not 1069 select a generic equivalent drug product.

1070 **SECTION 27.** Section 73-21-119, Mississippi Code of 1972, is 1071 reenacted as follows:

1072 73-21-119. (1) The label of the container of any drug 1073 product which is sold within the State of Mississippi for resale 1074 at retail and which requires a prescription to be dispensed at 1075 retail shall contain at a minimum the name of the manufacturer of 1076 the final dosage unit, expiration date if applicable, batch or lot 1077 number and national drug code.

1078 (2) Whenever product selection is made, the pharmacist shall 1079 indicate on the label of the dispensed container the initials 1080 "G.E." and the proprietary name of the product dispensed or the 1081 generic name of the product dispensed and its manufacturer either 1082 written in full or appropriately abbreviated, unless the 1083 prescriber indicates that the name of the drug product shall not 1084 appear on the label.

1085 **SECTION 28.** Section 73-21-121, Mississippi Code of 1972, is 1086 reenacted as follows:

1087 73-21-121. (1) Product selection as authorized by Sections 1088 73-21-115 through 73-21-119 shall not constitute evidence of 1089 negligence by the dispensing pharmacist when such product 1090 selection is in accordance with reasonable and prudent pharmacy 1091 practice. No prescriber shall be liable for civil damages or in 1092 any criminal prosecution arising from the incorrect product 1093 selection by a pharmacist.

(2) Any person having knowledge relating to a pharmacist or to a pharmacy student which might provide grounds for disciplinary action by the board may report relevant facts to the board, and shall by reason of reporting such facts in good faith be immune from civil liability.

H. B. No. 416 02/HR07/R924 PAGE 34 (RF\HS) (3) Any person furnishing information in the form of data, reports or records to the board or to a pharmacist organization approved by the board to receive such information, where such information is furnished for the purpose of aiding a pharmacist or a pharmacy student impaired by chemical abuse or by mental or by physical illness, shall by reason of furnishing such information in good faith be immune from civil liability.

(4) The records of the board or the records of a pharmacist organization approved by the board to aid pharmacists or pharmacy students impaired by chemical abuse, where such records relate to the impairment, shall be confidential and are not considered open records; provided, however, the board may disclose this confidential information only:

1112 (a) In a disciplinary hearing before the board, or in1113 an appeal of an action or order of the board;

(b) To the pharmacist licensing or disciplinary authorities of other jurisdictions in the case of a pharmacist who lis licensed in, or seeking transfer to, another state; or

1117 (c) Pursuant to an order of a court of competent 1118 jurisdiction.

1119 SECTION 29. Section 73-21-123, Mississippi Code of 1972, is
1120 reenacted as follows:

73-21-123. Nothing in this chapter shall be construed to 1121 prevent, or in any manner interfere with, or to require a permit 1122 1123 for the sale of nonnarcotic nonprescription drugs which may be lawfully sold under the United States Food, Drug and Cosmetic Act 1124 1125 (21 USCS 301 et seq. as now or hereafter amended) without a prescription, nor shall any rule or regulation be adopted by the 1126 board under the provisions of this chapter which shall require the 1127 sale of nonprescription drugs by a licensed pharmacist of in a 1128 1129 pharmacy or otherwise apply to or interfere with the sale or 1130 distribution of such drugs.

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1131 **SECTION 30.** This act shall take effect and be in force from 1132 and after July 1, 2002.