By: Representative Read

To: Public Health and Welfare

HOUSE BILL NO. 10

AN ACT TO REENACT SECTION 73-21-108, MISSISSIPPI CODE OF 1 1972, WHICH REQUIRES ANY PERSON OR BUSINESS THAT SELLS HOME 2 3 MEDICAL EQUIPMENT TO OBTAIN A MEDICAL EQUIPMENT SUPPLIER PERMIT FROM THE STATE BOARD OF PHARMACY, AND WHICH WAS REPEALED BY OPERATION OF LAW ON JULY 1, 2001; TO RATIFY THE ISSUANCE OF 4 5 PERMITS UNDER SECTION 73-21-108 AND MEDICAID REIMBURSEMENT 6 PAYMENTS MADE TO HOLDERS OF THOSE PERMITS DURING THE PERIOD FROM 7 JULY 1, 2001, UNTIL THE EFFECTIVE DATE OF THIS ACT; AND FOR RELATED PURPOSES. 8 9 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MISSISSIPPI: 10 11 SECTION 1. Section 73-21-108, Mississippi Code of 1972, is reenacted as follows: 12 73-21-108. (1) Definitions. For the purposes of this 13 14 section: "Home medical equipment" means technologically 15 (a) 16 sophisticated medical equipment and devices usable in a home care setting, including, but not limited to: 17 18 (i) Oxygen for human consumption, oxygen concentrators and/or oxygen delivery systems and equipment; 19 (ii) Ventilators; 20 21 (iii) Respiratory disease management devices; (iv) Electronic and computer driven wheelchairs 22 and seating systems; 23 24 (v) Apnea monitors; 25 (vi) Transcutaneous electrical nerve stimulator (TENS) units; 26 (vii) Low air loss cutaneous pressure management 27 devices; 28 29 (viii) Sequential compression devices; 30 (ix) Neonatal home phototherapy devices; N3/5 H. B. No. 10 023E/HR40/R25.1

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(x) Feeding pumps; and

32 (xi) Other similar equipment as defined in33 regulations adopted by the board.

The term "home medical equipment" does not include medical 34 35 equipment used in the normal course of treating patients by 36 hospitals, hospices, long-term care facilities or home health agencies, or medical equipment used or dispensed by health care 37 professionals licensed by the State of Mississippi if the 38 professional is practicing within the scope of his or her 39 professional practice. In addition, the term does not include 40 41 items such as upper and lower extremity prosthetics, canes, crutches, walkers, bathtub grab bars, standard wheelchairs, 42 43 commode chairs and bath benches.

(b) "Home medical equipment services" means the
delivery, installation, maintenance, replacement, and/or
instruction in the use of home medical equipment, used by a sick
or disabled individual, to allow the individual to be cared for
and maintained in a home or noninstitutional environment.

49 (c) "Medical gas" means those gases and liquid oxygen50 intended for human consumption.

51 (d) "Order" means an order issued by a licensed 52 practitioner legally authorized to order home medical equipment 53 and/or medical gases.

Permit required. (a) No person, business or entity 54 (2) 55 located in this state or outside of this state that is subject to this section shall sell, rent or provide or offer to sell, rent or 56 57 provide directly to patients in this state any home medical equipment, legend devices, and/or medical gas unless such person, 58 business or entity first obtains a Medical Equipment Supplier 59 Permit from the board. 60

(b) The permitting requirements of this section apply
to all persons, companies, agencies and other business entities
that are in the business of supplying home medical equipment to

H. B. No. 10 023E/HR40/R25.1 PAGE 2 (RF\BD) 64 patients in their places of residence and that bill the patient or 65 the patient's insurance, Medicare, Medicaid or other third party 66 payor for the rent or sale of that equipment.

(c) The board shall require a separate permit for each
facility location directly or indirectly owned or operated in this
state.

(d) The application for a permit shall be made to the board on a form supplied by the board and shall be accompanied by a fee of not more than Three Hundred Dollars (\$300.00), as prescribed by the board. Once issued, every permit must be renewed annually, and the renewal fee shall be not more than One Hundred Seventy-five Dollars (\$175.00), as prescribed by the board.

77 (e) All permits issued under this section shall expire annually on June 30 of each year. Applications for renewal must 78 be made to the board on or before June 30 and must be accompanied 79 by the fee as prescribed by the board. A late renewal fee of One 80 Hundred Dollars (\$100.00) shall be added to all renewal 81 applications received by the board after June 30 of each renewal 82 83 period. The permit shall become void if the renewal application, renewal fee and the late renewal fee are not received by the board 84 85 by September 30 of each year.

86 (3) Exemptions. (a) The permitting requirements of this
87 section do not apply to the following entities or practitioners
88 unless they have a separate business entity, company, corporation
89 or division that is in the business of providing home medical
90 equipment for sale or rent to patients at their places of
91 residence:

Home health agencies; 92 (i) (ii) Hospitals; 93 Wholesalers and/or manufacturers; 94 (iii) 95 (iv) Medical doctors, physical therapists, respiratory therapists, occupational therapists, speech 96 H. B. No. 10 023E/HR40/R25.1 PAGE 3 (RF\BD)

97 pathologists, optometrists, chiropractors and podiatrists who use 98 home medical equipment and/or legend devices in their individual 99 practices;

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(v) Pharmacies;

101 (vi) Hospice programs;

102 (vii) Nursing homes and/or long-term care 103 facilities;

104 (viii) Veterinarians; dentists; and emergency105 medical services.

(b) Although community pharmacies are exempt from the permitting requirements of this section, they shall be subject to the same regulations that are applicable to permitted businesses or entities for the sale or rental of home medical equipment covered by this section.

(c) Nothing in this section shall prohibit trained individuals from using oxygen, liquid oxygen and/or legend devices in emergencies.

(d) Nothing in this section shall prohibit the prehospital emergency administration of oxygen by licensed health care providers, emergency medical technicians, first responders, fire fighters, law enforcement officers and other emergency personnel trained in the proper use of emergency oxygen.

(4) Order required. Home medical equipment suppliers shall
not provide any home medical equipment to a patient without a
valid order from an authorized licensed practitioner.

(5) Regulations. The board shall adopt regulations for the distribution and sale or rental of home medical equipment, legend devices and medical gases that promote the public health and welfare and comply with at least the minimum standards, terms and conditions of federal laws and regulations. The regulations shall include, without limitation:

H. B. No. 10 023E/HR40/R25.1 PAGE 4 (RF\BD) (a) Minimum information from each home medical
equipment, legend device and medical gas supplier required for
permitting and renewal permits;

(b) Minimum qualifications of persons who engage in thedistribution of home medical equipment;

(c) Appropriate education, training or experience ofpersons employed by home medical equipment suppliers;

135 (d) Minimum standards for storage of home medical136 equipment;

(e) Minimum requirements for the establishment and
maintenance of all records for the sale, rental and servicing of
home medical equipment; and

140 (f) Minimum standards of operation and professional141 conduct.

142 (6) Medical Equipment Advisory Committee to the board.

(a) A Medical Equipment Advisory Committee (MEAC),
composed of three (3) members selected by the Mississippi
Association of Medical Equipment Suppliers and approved by the
board, shall review and make recommendations to the board
regarding all regulations dealing with home medical equipment,
legend devices and medical gases that are proposed by the board
and before they are adopted by the board.

(b) All MEAC members must have been actively involved in the home medical equipment business for a minimum of five (5) years before the selection to the committee and shall hold and maintain, in good standing, a permit issued by the board under this section.

(c) The MEAC members shall meet at least quarterly and review all home medical equipment suppliers' inspection reports. All complaints and reports of investigations of violations of law or regulations regarding home medical equipment, legend devices and medical gases shall first be reviewed by the MEAC. After review, the MEAC may make recommendations to the board's

H. B. No. 10 023E/HR40/R25.1 PAGE 5 (RF\BD) 161 Investigations Review Committee regarding further administrative 162 action by the board.

(d) The MEAC shall keep and maintain minutes of all
meetings of the MEAC and shall provide copies of the minutes to
the board on a quarterly basis.

166 (7) Revocation, Suspension or Restriction of Permit and
 167 Penalties.

(a) The board may revoke, suspend, restrict or refuse
to issue or renew a permit or impose a monetary penalty, in
accordance with Section 73-21-103 except that the monetary penalty
shall not exceed Ten Thousand Dollars (\$10,000.00) per violation,
if the business or holder of a permit or applicant for a permit
issued under this section has committed or is found guilty by the
board of any of the following:

(i) Violation of any federal, state or local law
or regulations relating to home medical equipment, legend devices
or medical gases.

178 (ii) Violation of any of the provisions of this179 section or regulations adopted under this section.

(iii) Commission of an act or engaging in a course
of conduct that constitutes a clear and present danger to the
public health and safety.

(iv) Filing a claim or assisting in the filing of a claim for reimbursement for home medical equipment or home medical equipment services that were not provided or that were not authorized to be provided.

187 (v) Failure to comply with any lawful order of the188 board.

(b) Disciplinary action by the board against a business
or any person holding a permit under this section shall be in
accordance with Section 73-21-99.

192 SECTION 2. (1) The issuance of any permits by the State193 Board of Pharmacy in accordance with the provisions of Section

H. B. No. 10 023E/HR40/R25.1 PAGE 6 (RF\BD) 194 73-21-108 as it existed on June 30, 2001, during the period from 195 July 1, 2001, until the effective date of House Bill No. 10, Third 196 Extraordinary Session of 2002, are ratified, approved and 197 confirmed.

198 (2) Any reimbursement payments for durable medical equipment services or medical supplies made by the Division of Medicaid to 199 200 holders of permits issued by the State Board of Pharmacy in accordance with the provisions of Section 73-21-108 as it existed 201 on June 30, 2001, during the period from July 1, 2001, until the 202 effective date of House Bill No. 10, Third Extraordinary Session 203 204 of 2002, are ratified, approved and confirmed; however, this subsection does not prevent or restrict the Division of Medicaid 205 from exercising any of the authority granted under Section 206 207 43-13-121 with respect to any of those reimbursement payments.

208 **SECTION 3.** This act shall take effect and be in force from 209 and after its passage.