Part 2635 Practice of Medicine

Part 2635: Chapter 1 Surgery/Post-Operative Care

Rule 1.1 Scope. The following regulation sets forth the policies of the Mississippi State Board of Medical Licensure regarding post-operative surgical care rendered by individuals licensed to practice medicine, osteopathic medicine and podiatric medicine in the state of Mississippi.


Rule 1.2 Definitions. For the purpose of Part 2635, Chapter 1 only, the following terms have the meanings indicated:

A. “Auxiliary” or “Auxiliaries” shall include, but is not limited to, registered nurses, licensed practical nurses, certified nursing assistants, physical therapists, nurse practitioners and optometrists.

B. “Under the supervision” means to critically watch, direct, advise and oversee, and to inspect and examine the actions of another health care practitioner.

C. “Physician” means any person licensed to practice medicine, osteopathic medicine or podiatric medicine in the state of Mississippi.

D. “Surgery” is defined as any operative procedure, including the use of lasers, performed upon the body of a living human being for the purposes of preserving health, diagnosing or curing disease, repairing injury, correcting deformity or defects, prolonging life, relieving suffering or any elective procedure for aesthetic, reconstructive or cosmetic purposes, to include, but not be limited to: incision or curettage of tissue or organ; suture or other repair of tissue or organ, including a closed as well as an open reduction of a fracture; extraction of tissue including premature extraction of the products of conception from the uterus; insertion of natural or artificial implants; or an endoscopic procedure.


Rule 1.3 Informed Consent. The responsibility for medical and surgical diagnoses is that of the licensed physician. In addition, it is the responsibility of the operating physician to explain the procedure and to obtain informed consent of the patient. It is not necessary, however, that the operating physician obtain or witness the signature of a patient on a written form evidencing informed consent.


Rule 1.4 Post-Surgical Care. The management of post-surgical care is the responsibility of the operating physician. The operating physician should provide those aspects of post-surgical care which are within the unique competence of the physician. Patients are best served by having post-surgical care conducted by the physician who best knows their condition—the operating physician.

Where the operating physician cannot personally provide post-surgical care, the physician must arrange before surgery for post-surgical care to be performed by another qualified physician who is acceptable to the patient. In this case, the operating physician may delegate discretionary post-operative activities to a qualified licensed physician. Like the operating physician, the physician
to whom a patient has been referred for post-surgical care should provide, at a minimum, those aspects of post-surgical care that are not delegable.

Unless otherwise provided by law, delegation of post-surgical activities to an auxiliary is permitted only if the auxiliary is under the supervision of the operating physician or the physician to whom the operating physician has referred a patient for post-surgical care. While an auxiliary may be authorized by law to provide certain aspects of post-surgical care, this does not relieve the operating physician of his or her responsibility to provide post-surgical care or arrange for the delegation of post-surgical care, when appropriate, as required by this rule.

Those aspects of post-surgical care which may be delegated to an auxiliary must be determined on a case-by-case basis, but shall be limited to those procedures which the auxiliary is authorized by law to perform and within the unique competence and training of the auxiliary.


Rule 1.5 Effective Date of Rules. The rules pertaining to Surgery/Post-Operative Care shall become effective October 23, 1994. Amended March 16, 2017.


Part 2635: Chapter 2 Office Based Surgery

Rule 2.1 Scope. This regulation sets forth the policies of the Mississippi State Board of Medical Licensure regarding office based surgery rendered by individuals licensed to practice medicine, osteopathic medicine and podiatric medicine in the state of Mississippi.


Rule 2.2 Definitions. For the purpose of Part 2635, Chapter 2 only, the following terms have the meanings indicated:

A. “Surgery” is defined as any operative procedure, including the use of lasers, performed upon the body of a living human being for the purposes of preserving health, diagnosing or curing disease, repairing injury, correcting deformity or defects, prolonging life, relieving suffering or any elective procedure for aesthetic, reconstructive or cosmetic purposes, to include, but not be limited to: incision or curettage of tissue or organ; suture or other repair of tissue or organ, including a closed as well as an open reduction of a fracture; extraction of tissue including premature extraction of the products of conception from the uterus; insertion of natural or artificial implants; or an endoscopic procedure. The use of local, general or topical anesthesia and/or intravenous sedation is the prerogative of the surgeon.

B. “Surgeon” is defined as a licensed physician performing any procedure included within the definition of surgery.

C. Implicit within the use of the term “equipment” is the requirement that the specific item named must meet current performance standards.

D. “Office surgery” is defined as surgery which is performed outside a hospital, an ambulatory surgical center, abortion clinic, or other medical facility licensed by the Mississippi State Department of Health or a successor agency. Physicians performing Level II or Level III office based surgery must register with the Mississippi State Board
of Medical Licensure. A copy of the registration form is attached hereto (Appendix A).

E. A “Surgical Event” for the purpose of this regulation is recognized as a potentially harmful or life-threatening episode related to either the anesthetic or the surgery. Any “Surgical Event” in the immediate perioperative period that must be reported are those which are life-threatening, or require special treatment, or require hospitalization, including, but not limited to the following: (1) serious cardiopulmonary or anesthetic events; (2) major anesthetic or surgical complications; (3) temporary or permanent disability; (4) coma; or (5) death.


Rule 2.3 General Requirements for Office Surgery. For all surgical procedures, the level of sterilization shall meet current OSHA requirements.

The surgeon must maintain complete records of each surgical procedure, including anesthesia records, when applicable and the records on all Level II and Level III cases shall contain written informed consent from the patient reflecting the patient’s knowledge of identified risks, consent to the procedure, type of anesthesia and anesthesia provider.

The surgeon must maintain a log of all Level II and Level III surgical procedures performed, which must include a confidential patient identifier, the type of procedure, the type of anesthesia used, the duration of the procedure, the type of post-operative care, and any surgical events. The log and all surgical records shall be provided to investigators of the Mississippi State Board of Medical Licensure upon request.

In any liposuction procedure, the surgeon is responsible for determining the appropriate amount of supernatant fat to be removed from a particular patient. Using the tumescent method of liposuction, the surgeon must fully document the anticipated amount of material to be removed in a manner consistent with recognized standards of care. Post-operatively, any deviation from the anticipated amount, and the reason for deviation, should be fully documented in the operative report. Morbidly obese patients should have liposuction performed in the hospital setting unless the surgeon can document significant advantage to an alternative setting.

A policy and procedure manual must be maintained in the office and updated annually. The policy and procedure manual must contain the following: duties and responsibilities of all personnel, cleaning and infection control, and emergency procedures.

The surgeon shall report to the Mississippi State Board of Medical Licensure any surgical events that occur within the office based surgical setting. This report shall be made within 15 days after the occurrence of a surgical event. A suggested form for reporting is attached hereto (Appendix B). The filing of a report of surgical event as required by this rule does not, in and of itself, constitute an acknowledgment or admission of malpractice, error, or omission. Upon receipt of the report, the Board may, in its discretion, obtain patient and other records pursuant to authority granted in Mississippi Code, Section 73-25-28.

The surgeon must have a written response plan for emergencies within his or her facility.
In offices where Level II and Level III office based surgery is performed, a sign must be prominently posted in the office which states that the office is a doctor’s office regulated pursuant to the rules of the Mississippi State Board of Medical Licensure. This notice must also appear prominently within the required patient informed consent.

Office surgery facilities should adhere to recognized standards such as those promulgated by the American Society of Anesthesiologists’ *Guidelines for Office-Based Anesthesia* or American Association of Nurse Anesthetists’ *Standards for Office Based Anesthesia*.

**Source:** Miss. Code Ann. §73-43-11 (1972, as amended).

**Rule 2.4 Level I Office Surgery.**

A. **Scope**

1. Level I office surgery includes, but not limited to, the following:
   i. Minor procedures such as excision of skin lesions, moles, warts, cysts, lipomas, Loop Electrosurgical Excision Procedures (LEEP), laser cone of cervix, laser/cautery ablation of warts or other lesions, and repair of lacerations or surgery limited to the skin and subcutaneous tissue performed under topical or local anesthesia not involving drug-induced alteration of consciousness.
   ii. Incision and drainage of superficial abscesses, limited endoscopies such as proctoscopies, flexible sigmoidoscopies, hysteroscopies, skin biopsies, arthrocentesis, paracentesis, dilation of urethra, cystoscopy procedures, and closed reduction of simple fractures or small joint dislocations (i.e., finger and toe joints).
   iii. Procedures requiring only topical, local or no anesthesia. Only minimal or no preoperative sedation should be required or used. No drug-induced alteration of respiratory effort or consciousness other than minimal preoperative tranquilization of the patient is permitted in Level I Office Surgery.
   iv. Chances of complication requiring hospitalization are remote.

2. **Standards for Level I Office Surgery**
   i. **Training Required**
      The surgeon’s continuing medical education should include management of toxicity or hypersensitivity to local anesthetic drugs. The surgeon’s continuing medical education shall include Basic Life Support Certification.
   ii. **Equipment and Supplies Required**
      Oral airway, positive pressure ventilation device, epinephrine (or other vasopressor), corticosteroids, antihistamines and atropine, if any anesthesia is used. The equipment and skills to establish intravenous access must be available if any other medications are administered. The equipment and supplies should reflect the patient population, i.e., pediatrics, etc.
   iii. **Assistance of Other Personnel Required**
      No other assistance is required, unless the specific surgical procedure being performed requires an assistant.

**Source:** Miss. Code Ann. §73-43-11 (1972, as amended).

**Rule 2.5 Level II Office Surgery.**

A. **Scope**
1. Level II Office Surgery is that in which perioperative medication and sedation are used orally, intravenously, intramuscularly, or rectally. If perioperative or intraoperative medication is administered, intraoperative and postoperative monitoring is required. Such procedures include, but are not limited to: hernia repair, hemorrhoidectomy, reduction of simple fractures, large joint dislocations, breast biopsies, dilatation and curettage, thoracentesis, and colonoscopy.

2. Level II Office surgery also includes any surgery in which the patient is sufficiently sedated to allow the patient to tolerate unpleasant procedures while maintaining adequate cardiorespiratory function and the ability to respond purposefully to verbal command and/or tactile stimulation. Patients whose only response is reflex withdrawal from a painful stimulus are sedated to a greater degree than encompassed by this definition.

3. Any procedures that may yield an excessive loss of blood should be covered under Level II.

B. Transfer Agreement Required
The surgeon must have a written transfer agreement from a licensed hospital within reasonable proximity. The transfer agreement should also include physician coverage of transferred patients if the physician does not have privileges at the hospital.

C. Level of Anesthetic
Local or peripheral nerve block, including Bier Block, plus intravenous or intramuscular sedation, but with preservation of vital reflexes.

D. Training Required
To perform office based surgery, the physician must be able to document satisfactory completion of surgical training such as Board certification or Board eligibility by a Board approved by the American Board of Medical Specialties or American Board of Osteopathic Specialties. The certification should include training in the procedures performed in the office setting. Alternative credentialing for procedures outside the physician’s core curriculum must be applied for through the Mississippi State Board of Medical Licensure and reviewed by a multi-specialty board appointed by the Director. In addition to the surgeon, there must be at least one assistant certified in Basic Life Support present during any Level II or III procedure. There should be at least one person certified in Advanced Cardiac Life Support present during any Level II or III procedure unless there is an anesthesiologist or certified registered nurse anesthetist to manage the anesthetic.

E. Equipment and Supplies Required
1. Full and current crash cart at the location the anesthetizing is being carried out.

   The crash cart must include, at a minimum, the following resuscitative medications, or other resuscitative medication subsequently marketed and available after initial adoption of this regulation, provided said medication has the same FDA approved indications and usage as the medications specified below:
   i. Adrenalin (epinephrine) Abboject 1mg-1:10,000; 10ml
   ii. Adrenalin (epinephrine) ampules 1mg-1:1000; 1ml
   iii. Atropine Abboject 0.1mg/ml; 5ml
   iv. Benadryl (diphenhydramine) syringe 50mg/ml; 1ml
   v. Calcium chloride Abboject 10%; 100mg/ml; 10ml
   vi. Dextrose Abboject 50%; 25g/50ml
   vii. Dilantin (phenytoin) syringe 250mg/5ml
viii. Dopamine 400mg/250ml pre-mixed
ix. Heparin 10,000 units/ml; 1 ml vial
x. Inderal (propranolol) 1mg/ml; 1 ml ampule
xi. Isuprel (isoproterenol) 1mg/5ml; 1:5000 ampule
xii. Lanoxin (digoxin) 0.5 mg/2ml ampule
xiii. Lasix (furosemide) 40 mg/4ml vial
xiv. Lidocaine Abboject 2%; 100mg/5ml
xv. Lidocaine 2 grams/500ml pre-mixed
xvi. Magnesium sulfate 50%; 20ml vial (1g/2ml)
xvii. Narcan (naloxone) 0.4mg/ml; 1ml ampule
xviii. Pronestyl (procainamide) 100mg/ml; 10ml vial
xix. Romazicon 5ml or 10 ml (0.1mg/ml)
xx. Sodium bicarbonate Abboject 50mEq/50ml
xxi. Solu-medrol (methylprednisolone) 125mg/2ml vial
xxii. Verapamil syringe 5mg/2ml

The above dosage levels may be adjusted, depending on ages of the patient population.

2. Suction devices, endotracheal tubes, laryngoscopes, etc.
3. Positive pressure ventilation device (e.g., Ambu) plus oxygen supply.
4. Double tourniquet for the Bier Block procedure.
5. Monitors for blood pressure/EKG/Oxygen saturation and portable approved defibrillator.
6. Emergency intubation equipment.
7. Adequate operating room lighting with onsite backup sufficient to supply required equipment perioperative equipment and monitors for a minimum of two (2) hours.
8. Sterilization equipment or facilities meeting Joint Commission requirements.
9. IV solution and IV equipment.

F. Assistance of Other Personnel Required
In addition to the surgeon there must be at least one assistant certified in Basic Life Support present during any Level II or III procedure. There should be at least one person certified in Advanced Cardiac Life Support present during any Level II or III procedure unless there is an anesthesiologist or certified registered nurse anesthetist to manage the anesthetic.

A registered nurse may only administer analgesic doses of medications on the direct order of a physician. An assisting anesthesia provider, including nurse providing sedation, may not function in any other capacity during the procedure. If additional assistance is required by the specific procedure or patient circumstances, such assistance must be provided by a physician, registered nurse, licensed practical nurse, or operating room technician.


Rule 2.6 Level III Office Surgery.
A. Scope
1. Level III Office Surgery is that surgery which involves, or might foreseeably require, the use of a general anesthesia or major conduction anesthesia and perioperative sedation. This includes the use of:
   i. Intravenous sedation beyond that defined for Level II office surgery;
ii. General Anesthesia: loss of consciousness and loss of vital reflexes with probable requirement of external support of pulmonary or cardiac functions; or
iii. Major Conduction anesthesia.

2. Only patients classified under the American Society of Anesthesiologist’s (ASA) risk classification criteria as Class I, II, or III are appropriate candidates for Level III office surgery. For ASA Class III patients, the surgeon must document in the patient’s record the justification for an office procedure rather than other surgical venues. The record must also document precautions taken that make the office a preferred venue for the particular procedure to be performed.

B. Transfer Agreement Required
   The surgeon must have a written transfer agreement from a licensed hospital within reasonable proximity. The transfer agreement must include physician coverage of transferred patients if the physician does not have privileges at the hospital. Level of Anesthetic
   1. General Anesthetic: loss of consciousness and loss of vital reflexes with probable requirement of external support of pulmonary or cardiac functions.
   2. Major Conduction: epidural, spinal, caudal or any block of a nerve or plexus more proximal than the hip or shoulder joint including visceral nerve blocks.

C. Training Required
   1. To perform office based surgery, the physician must be able to document satisfactory completion of surgical training such as board certification or board eligibility by a board approved by the American Board of Medical Specialties or American Board of Osteopathic Specialties. The certification should include training in the procedures performed in the office setting. Alternative credentialing for procedures outside the physician’s core curriculum must be applied for through the Mississippi State Board of Medical Licensure and reviewed by a multi-specialty board appointed by the Executive Director.
   2. In addition to the surgeon there must be at least one assistant certified in Basic Life Support present during any Level II or III procedure. There should be at least one person certified in Advanced Cardiac Life Support present during any Level II or III procedure unless there is an anesthesiologist or certified registered nurse anesthetist to manage the anesthetic.
   3. Emergency procedures related to serious anesthesia complications should be formulated, periodically reviewed, practiced, updated, and posted in a conspicuous location.

D. Equipment and Supplies Required
   1. Equipment, medication and monitored post-anesthesia recovery must be available in the office. If anesthetic agents include inhaled agents, other than nitrous oxide, medications must include a stock of no less than 12 vials of Dantrolene.
   2. The facility, in terms of general preparation, equipment, and supplies, must be comparable to a free standing ambulatory surgical center, including, but not limited to, recovery capability, and must have provisions for proper record keeping.
   3. Blood pressure monitoring equipment; EKG; end tidal CO2 monitor; pulse oximeter, precordial or esophageal stethoscope, emergency intubation equipment and a temperature monitoring device must be available for all phases of perioperative care.
4. Table capable of Trendelenburg and other positions necessary to facilitate the surgical procedure.
5. IV solutions and IV equipment.
6. All equipment and supplies listed under Part 2635, Rule 2.5, Level II.
E. Assistance of Other Personnel Required
   An anesthesiologist or certified registered nurse anesthetist must administer the general or regional anesthesia and a physician, registered nurse, licensed practical nurse, or operating room technician must assist with the surgery. The anesthesia provider may not function in any other capacity during the procedure. A licensed physician or a licensed registered nurse with post-anesthesia care unit experience or the equivalent, and credentialed in Advanced Cardiac Life Support, or in the case of pediatric patients, Pediatric Advanced Life Support, must be available to monitor the patient in the recovery room until the patient has recovered from anesthesia.


Rule 2.7 Effective Date of Rules. The above rules pertaining to office based surgery shall become effective September 1, 2001.


Part 2635 Chapter 3: Laser Devices

Rule 3.1 Laser Devices. The use of laser, pulsed light or similar devices, either for invasive or cosmetic procedures, is considered to be the practice of medicine in the state of Mississippi and therefore such use shall be limited to physicians and those directly supervised by physicians, such that a physician is on the premises and would be directly involved in the treatment if required. These rules shall not apply to any person licensed to practice dentistry if the laser, pulsed light, or similar device is used exclusively for the practice of dentistry.


Part 2635 Chapter 4: Chelation Therapy

Rule 4.1 Chelation Therapy. The use of EDTA (ethylenediaminetetraacetic acid) outside of FDA approved clinical indications or an approved research protocol (see below) is not permitted. Other off-label uses may be permissible if there is substantial, high-quality research to support such use. The research should be peer-reviewed and published in recognized journals such as those cited in PubMed or in the National Library of Medicine. Specific reference should be made to the publications and research in the medical record. Informed consent for off-label use should be
obtained. Use of EDTA in any other manner may be considered to be violation of Mississippi Code, Section 73-25-29(8)(d).

However, EDTA may be used when a licensee experienced in clinical investigations has applied for and received from the Board written approval for off-label use in a clinical investigation. The licensee applying for approval must be the principal investigator for the protocol or subject to the direction of the principal investigator.

Advertising EDTA’s administration for off-label use, except for approved research protocols, is prohibited. Such advertising may be considered to be violation of Mississippi Code, Section 73-25-29(8)(d) and/or the rules promulgated pursuant thereto.

**Adopted July 18, 2002. Amended March 16, 2017.**

*Source: Miss. Code Ann. §73-43-11 (1972, as amended).*

**Part 2635 Chapter 5: Practice of Telemedicine**

**Rule 5.1 | Definitions**

For the purpose of Part 2635, Chapter 5 only, the following terms have the meanings indicated:

A. “**Provider**” means any physician or physician assistant who holds an unrestricted license to practice medicine in the state of Mississippi.

B. “**Telemedicine**” is the practice of medicine by a licensed healthcare provider using HIPAA-compliant telecommunication systems, including information, electronic, and communication technologies, remote monitoring technologies and store-and-forward transfer technology. These technologies may be used to facilitate, but are not limited to, provider to patient or provider to provider interactions. The technology must be capable of replicating the interaction of a traditional in-person encounter between a provider and a patient. This definition does not include the practice of medicine through postal or courier services.

C. “**Emergency Telemedicine**” is a unique combination of telemedicine used in a consultative interaction between a physician board certified, or board eligible, in emergency medicine, and an appropriate skilled health professional (nurse practitioner or physician assistant).

D. “**Primary Center**” is any facility providing telemedicine services to Satellite Centers, as defined in definition ‘G’.

E. “**Remote Monitoring**” is defined as the use of technology to remotely track health care data for a patient released to his or her home or a care facility, usually for the intended purpose of reducing readmission rates.

F. “**Real-Time Telemedicine**” is defined as real-time communication using interactive audio and visual equipment, such as a video conference with a specialist, also known as ‘synchronous communication.’
G. “Satellite Center” is any facility receiving telemedicine services from a Primary Center, as defined in definition ‘D’.
H. “Store-and-Forward Transfer Technology” is defined as technology which facilitates the gathering of data from the patient, via secure email or messaging service, which is then used for formulation of a diagnosis and treatment plan, also known as ‘asynchronous communication.’


Rule 5.2 | Licensure

The practice of medicine is deemed to occur in the location of the patient. Therefore, only providers holding a valid Mississippi license are allowed to practice any form of telemedicine, as defined in R.5.1, in Mississippi. The interpretation of clinical laboratory studies as well as pathology and histopathology studies performed by physicians without Mississippi licensure is not the practice of telemedicine provided a Mississippi licensed provider is responsible for accepting, rejecting, or modifying the interpretation. The Mississippi licensed provider must maintain exclusive control over any subsequent therapy or additional diagnostics.


Rule 5.3 | Informed Consent

The provider using any form of telemedicine, as defined in R.5.1, should obtain the patient’s informed consent before providing care via telemedicine technology. In addition to information relative to treatment, the patient should be informed of the risk and benefits of being treated via a telemedicine network including how to receive follow-up care or assistance in the event of an adverse reaction to treatment or if there is a telemedicine equipment failure.


Rule 5.4 | Physician Patient Relationship

In order to practice any form of telemedicine, as defined in R.5.1, a valid “physician patient relationship” must be established. The elements of this valid relationship are:

A. verify that the person requesting the medical treatment is in fact who they claim to be;
B. conducting an appropriate history and physical examination of the patient that meets the applicable standard of care;
C. establishing a diagnosis through the use of accepted medical practices, i.e., a patient history, mental status exam, physical exam and appropriate diagnostic and laboratory testing;
D. discussing with the patient the diagnosis, risks and benefits of various treatment options to obtain informed consent;
E. insuring the availability of appropriate follow-up care; and
F. maintaining a complete medical record available to patient and other treating health care providers.
Rule 5.5 | Examination

Providers using telemedicine technologies to provide medical care to patients located in Mississippi must provide an appropriate examination prior to diagnosis and treatment of the patient. However, this exam need not be in person if the technology is sufficient to provide the same information to the physician as if the exam had been performed face to face.

Store-and-Forward Transfer Technology may be used to enhance, but never replace, real-time provider-patient interaction. Provider-patient interaction may be audio-visual or audio only where medically appropriate.

Other exams may be appropriate if a licensed health care provider is on site with the patient and is able to provide various physical findings that the physician needs to complete an adequate assessment. However, a simple questionnaire without an appropriate exam is in violation of this policy and may subject the physician to discipline by the Board.

Rule 5.6 | Medical Records

The provider treating a patient through a telemedicine network must maintain a complete record of the patient’s care. The provider must maintain the record’s confidentiality and disclose the record to the patient consistent with state and federal laws. If the patient has a primary treating physician and a telemedicine provider for the same medical condition, then the primary physician’s medical record and the telemedicine provider’s record constitute one complete patient record.

Rule 5.7 | Consultative Physician Limited

A duly licensed physician may remotely consult with a duly licensed and qualified Advanced Practice Registered Nurse (“APRN”) or Physician’s Assistant (“PA”), who is in a hospital setting, using telemedicine. The physician providing Emergency Telemedicine must be either board certified or board eligible in emergency medicine. The Board may waive this requirement under extraordinary circumstances.

For the purposes of Emergency Telemedicine services, licensees will only be authorized to provide the aforementioned services to those emergency departments of licensed hospitals who have an average daily census of fifty (50) or fewer acute care/medical surgical occupied beds as defined by their Medicare Cost Report. Exceptions may be considered by the Board for physicians affiliated with facilities maintaining greater than fifty (50) beds, but not more than one hundred (100) beds.
Satellite Centers who receive telemedicine services/assistance from a Primary Center must have a transfer agreement with a facility that offers a higher level of care, in order to send any patients who require transfer for a higher level of care.


**Rule 5.8 | Reporting Requirements**

Annual reports detailing quality assurance activities, adverse or sentinel events shall be submitted for review to the Mississippi State Board of Medical Licensure by all institutions and/or hospitals operating teleemergency programs.

**Rule 5.9 | Automated Dispensaries**

Recognizing the emergence of sophisticated technology which allows certain levels of automation to the usual and customary process of seeing a provider, to include obtaining a prescription and then filling that prescription at a pharmacy, automated dispensary systems which provide the patient’s medications pursuant to a valid telemedicine visit with a licensee of the Board will not be considered in violation of Part 2640, Rule 1.9 Requirements for Dispensing Physicians. Any physician utilizing the automated dispensary will be responsible for the proper maintenance and inventory/accountability requirements as if the physician were personally dispensing the medications to the patient from his or her stock in their personal practice, as required in Rule 1.9 of Part 2640. An automated dispensary may not dispense controlled substances, and refills of medications may not be issued without a follow-up visit with the physician.

Of paramount importance to any automated dispensary process is the continued emphasis on a patient’s freedom of choice, as it pertains to selecting a pharmacy to fill any prescriptions authorized. The failure of any system utilizing an automated dispensary to appropriately advise the patient of his or her right to choose where their medications are filled will constitute a violation of Part 2640, Rule 1.12 Freedom of Choice.

Any telemedicine service devices or systems which contain automated dispensaries, containing medications ordered and maintained by physician licensees, shall be subject to the oversight of the Board and the Mississippi Board of Pharmacy, as stated in Part 2640, Rule 1.9, and may not operate in this state until approved by both Boards.


**Part 2635 Chapter 6: Electrodiagnostic Testing**

**Rule 6.1 General.** Electrodiagnostic testing includes two primary categories: needle electromyography testing and nerve conduction testing.
The purpose of both categories of electrodiagnostic testing is to detect abnormalities of the peripheral neuromuscular system or to determine the extent and degree of recovery of neuromuscular abnormalities.


Rule 6.2 Delegation of Electrodiagnostic Testing Procedures. Electrodiagnostic testing is a clinical diagnostic study that must be considered only in the light of the clinical finding. The person performing electrodiagnostic testing must be able to elicit the pertinent history and perform the necessary examination to define the clinical problems. Differential diagnoses must be considered, and as abnormalities unfold or fail to unfold during the course of testing, the electrodiagnostic testing may be modified until a probable diagnosis is reached.

Electrodiagnostic testing procedures may be delegated to a specifically trained non-physician or physician in a residency or fellowship training program. The responsible electrodiagnostic physician need not be physically present but must be immediately available within the same building throughout the performance of the entire procedure.


Part 2635 Chapter 7: Internet Prescribing

Rule 7.1 Internet Prescribing. Essential components of proper prescribing and legitimate medical practice require that the physician obtains a thorough medical history and conducts an appropriate physical and/or mental examination before prescribing any medication.

Prescribing drugs to individuals that the physician has never met and based solely on answers to a set of questions, as is found in Internet or toll-free telephone prescribing fails to meet an acceptable standard of care and could constitute unprofessional conduct subject to disciplinary action.


Part 2635 Chapter 8: Medical Expert Activities by Physicians

Rule 8.1 Authority and Purpose. The Mississippi State Board of Medical Licensure (hereinafter referred to as “the Board”) adopts these rules governing medical expert activities by physicians pursuant to Chapters 25 and 43 of Title 73 of the Mississippi Code. The Mississippi State Board of Medical Licensure finds it necessary to fulfill its statutory responsibilities by adopting these rules in order to protect the public, to set professional standards, to enforce the provisions of law regarding the performance of medical expert activities by physicians, and to further other legitimate government purposes in the public interest.


Rule 8.2 Scope. These rules apply to any physician who performs medical expert activities regarding any person, facility, or entity located within the state of Mississippi, or regarding an
event alleged to have occurred within the state of Mississippi, regardless of the location, type, or status of the physician’s medical expert activity, the presence or absence of the physician expert’s license to practice medicine in Mississippi, the physician expert’s presence or absence of a physician-patient relationship in Mississippi, the type of medical expert activity performed (e.g., oral testimony or a written statement), or the setting in which the medical expert activity is performed (e.g., a state or federal court or administrative agency).

No part of these rules is intended to conflict with or supercede the authority of any state or federal court or administrative agency to designate a physician as a medical expert in a legal matter then pending before the court or agency. The Board does not intend for these rules to conflict with or supercede the description or regulation of the function of a physician serving as an “expert” as that term is used in the Mississippi Rules of Evidence or in other provisions of law, rules, or decisions of any court or administrative agency.

No part of these rules is intended to conflict with or supercede the authority of a person other than a physician to serve as an expert in a legal matter. Furthermore, the Board does not intend for these rules to have any effect on physicians’ participation in legal proceedings in a capacity other than as a medical expert.


Rule 8.3 Definition of Medical Expert Activities. For the purposes of these rules only, the Mississippi State Board of Medical Licensure has determined that the definition of the term “medical expert activities” includes, but is not limited to, the use of medical knowledge and professional judgment by a physician to:

A. Suggest or recommend to a person any medical advice or other agency (whether material or not material).
B. Perform medical services (including, but not limited to, a physical or mental examination of a person).
C. Conduct a review of a person’s medical record.
D. Serve as a medical consultant.
E. Render a medical opinion concerning the diagnosis or treatment of a person.
F. Produce a written medical expert opinion report, affidavit, or declaration.
G. Give testimony under oath as a medical expert at a state or federal hearing, deposition, trial, administrative agency proceeding, alternative dispute resolution proceeding, or any other legal proceeding, regarding the medical issues in a legal matter or claim for injuries that is then pending in a court or administrative agency, or which may be filed or asserted whether or not such claim ever results in a pending legal matter and which involves a person, facility, or entity located within the state of Mississippi, or an event alleged to have occurred within the state of Mississippi.


Rule 8.4 Licensure and Qualification Requirements. Except as otherwise provided by law, rule or regulation of this state, any medical expert activity by a physician regarding a legal matter pending in a state or federal court or administrative agency in Mississippi must be performed by a physician who holds a current unrestricted medical license in Mississippi, another state or foreign jurisdiction, and who has the qualifications to serve as a medical expert on the issue(s) in question.
by virtue of knowledge, skill, experience, training, or education. This rule does not supersede the policies and rules of the Board in regards to unreferred diagnostic screening tests.

The practice of any physician not licensed in Mississippi that meets the licensure and qualification requirements stated in the above paragraph shall be deemed automatically by the Board to be authorized to include the performance of medical expert activities as an otherwise lawful practice, without any need for licensure verification or further requirement for licensure. In accordance with the provisions of law in Mississippi, any physician not licensed in Mississippi whose practice is deemed automatically by the Board to be authorized to include the performance of medical expert activities as an otherwise lawful practice shall be subject to regulation by the Board regarding the physician’s performance of such medical expert activities in the state of Mississippi.


Rule 8.5 Professional Standards. Any physician who performs medical expert activities must:
A. Comply with these rules and all applicable provisions of Mississippi law (e.g., statutes, court rules and decisions, and other administrative agency rules) with regard to the performance of medical expert activities.
B. Comply with medical ethics principles, including, but not limited to, ethics principles established by the American Medical Association and relevant medical specialty associations.
C. Be honest in all professional interactions involving his or her medical expert activities.
D. Not accept payment for medical expert activities that is contingent upon the result or content of any medical diagnosis, opinion, advice, services, report, or review; or that is contingent upon the outcome of any case, claim, or legal matter then pending or contemplated.
E. Not make or use any false, fraudulent, or forged statement or document.


Rule 8.6 Professional Accountability for Violation of Rules. Any physician who performs medical expert activities, whether or not licensed to practice medicine in Mississippi, may be disciplined or otherwise held professionally accountable by the Board, upon a finding by the Board that the physician is unqualified as evidenced by behavior including, but not limited to, incompetent professional practice, unprofessional conduct, or any other dishonorable or unethical conduct likely to deceive, defraud, or harm the public.

Any violation of Part 2635, Rule 8.5 as enumerated above shall constitute unprofessional conduct in violation of Mississippi Code, Section 73-25-29(8).


Rule 8.7 Complaint Procedure, Investigation, Due Process, and Actions Available to the Board. Any person who has reason to believe that any physician may have failed to comply with any part of these rules in the performance of medical expert activities may make a complaint to the Mississippi State Board of Medical Licensure on a complaint form that is furnished by the Board.

Any physician, whether or not licensed to practice medicine in Mississippi, who performs medical expert activities in the context of a legal matter regarding any person, facility, entity, or event located within the state of Mississippi may be subject to an investigation by the Mississippi State
Board of Medical Licensure upon the receipt of a complaint regarding the physician’s conduct or practice. Any such physician shall be afforded the due process procedures of the law and Board rules. The Board, in its sole discretion, may refer the complaint to the medical licensure authority of another state, or to any other appropriate legal authority.

Any physician may request, or may be summoned by the Board, to appear before the Board at a hearing to consider the physician’s compliance with these rules. Any physician’s failure to appear when summoned to a hearing may be deemed by the Board to be a waiver of the physician’s due process opportunity to appear before the Board and may result in a finding by the Board that the physician is out of compliance with these rules in absentia.

In disciplining a physician licensed to practice medicine in Mississippi or otherwise holding any physician professionally accountable pursuant to these rules and to the statutes, rulings, and other rules and provisions of Mississippi law, the actions that the Mississippi State Board of Medical Licensure may take include, but are not limited to, one or more of the following:

A. Denying, suspending, restricting, or revoking a Mississippi license to practice medicine.
B. Administering a public or private reprimand to a Mississippi licensed physician.
C. Assessing up to $10,000 of the reasonable investigation costs expended by the Board in investigating a Mississippi licensed physician.
D. Moving for an injunction in Chancery Court to prohibit any physician’s further performance of medical expert activities.
E. Petitioning the Chancery Court to cite any noncompliant physician for contempt of court.
F. Referring the matter to another medical licensure authority or other legal authority for action regarding any physician.
G. Any other action regarding any physician that the Board may deem proper under the circumstances (e.g., issuing an advisory letter of concern; issuing a notice of warning; issuing a cease and desist notice; or adopting a resolution of disapproval of any physician’s medical expert activities).

Any physician who is found by the Mississippi State Board of Medical Licensure to have failed to comply with any part of these rules may be reported by the Board to any person or organization appropriate under the circumstances in order to enforce or comply with the law or to protect the public, including, but not limited to, the National Practitioner Data Bank, the U.S. Department of Health and Human Services Office of the Inspector General, the Centers for Medicare and Medicaid Services, the Federation of State Medical Boards, the medical licensure authority or state medical association in any state in which the physician is licensed to practice medicine, the American Board of Medical Specialties and any of its member specialty boards, the Mississippi Attorney General or District Attorney, the United States Attorney, any state or federal court or administrative agency, any national or state professional organization or medical specialty association, and any other appropriate person, government agency, healthcare entity, or legal authority.


Rule 8.8 Compliance Policy and Exemptions. In assuring compliance with these rules, the duty shall be on the physician, not on the party who engaged the physician to perform medical expert activities and not on any other person or entity, to ensure that his or her medical expert activities
comply with these rules. Any physician who claims to be exempt from these rules shall have the burden of proving to the Board that the exemption is valid.

Amended May 20, 2010.


References.


Mississippi Rule of Evidence 702

“Rules, Laws, and Policies of the Mississippi State Board of Medical Licensure.” Published by the Mississippi State Board of Medical Licensure and available at Internet address www.msbml.ms.gov

Hall v. Hilbun, 466 So. 2d 856 (Miss. 1985)


Findings of Fact adopted by the Mississippi State Board of Medical Licensure on May 18, 2006.**

**COMMENT: Based on information presented to the Board at a public hearing on this matter on March 9, 2006, and on May 18, 2006, and on research and analysis of information obtained by Board members and its staff and attorneys, and also on comments received from numerous sources, including the Board’s Consumer Health Committee, leaders of the medical and legal professions, former judges, officials from the Federation of State Medical Boards, and members of the public, the Mississippi State Board of Medical Licensure makes the following Findings of Fact:

1. A physician’s professional practice, conducted pursuant to the privilege of possessing a medical license, historically has been subject to regulation by other members of the medical profession, by methods such as peer review, performance evaluation, quality assurance monitoring, and other methods of regulation. However, there is a problem in Mississippi with the lack of regulation of medical expert activities by physicians. This lack of regulation causes the performance of medical expert activities to be vulnerable to fraud, abuse, dishonesty, deception, incompetence, and other forms of unprofessional, dishonorable, and unethical conduct by physician experts, all of which are harmful to the public.

2. A physician’s performance of medical expert activities involves a lawful part of a physician’s practice that is historically an area of state concern and that the Board has the statutory authority and duty to regulate in order to protect the public.

3. A physician’s medical expert activities involve practices that are likely to affect the health, safety,
Part 2635 Chapter 9 Community-Based Immunization Programs

**Rule 9.1 Scope.** The administration of vaccinations constitutes the practice of medicine, as defined by Mississippi Code Section 73-43-11, and thus may only be performed by a physician licensed rights, remedies, and general welfare of persons in Mississippi.

4. In keeping with the public policy and provisions of law in Mississippi, the performance of medical expert activities, regardless of the physician expert’s location or state(s) of medical licensure, is a lawful practice that requires a qualified physician, and is therefore subject to regulation by, and professional accountability to, the Mississippi State Board of Medical Licensure.

5. Due to its physician membership and statutory authority, the Mississippi State Board of Medical Licensure is uniquely able to establish and enforce licensure requirements, qualification requirements, and Professional Standards related to the performance of medical expert activities by physicians, especially with regard to ethical conduct and competent practice.

6. Regardless of a physician’s state(s) of medical licensure, a physician who performs medical expert activities in a legal matter has an ethical duty to practice according to the standards of medical professionalism, to perform all medical expert activities in an honest and competent manner, and to strive to report to appropriate entities any physician who is deficient in character or competence or who engages in fraud or deception.

7. In keeping with the public policy and provisions of law in Mississippi and principles of medical ethics, it is unprofessional, dishonorable, and unethical for a physician to willfully state an opinion or a material fact as a medical expert in the context of a legal matter that the physician knows or should know is false, or that a reasonable person could objectively conclude was a misrepresentation or other distortion of the truth, or was intended by the physician to mislead or deceive a judge, juror, lawyer, litigant, other expert, hearing officer, administrative body, investigator, legal authority, or any finder of fact.

8. In adopting these rules, the Mississippi State Board of Medical Licensure has attempted to tailor these rules as closely as possible to the current provisions of Mississippi law, in order to regulate medical expert activities for the legitimate government purpose of protecting the public and to further other legitimate government purposes in the public interest.

9. In adopting these rules, the Mississippi State Board of Medical Licensure states that its intent is only to regulate the conduct and practice of physicians who perform medical expert activities in Mississippi. The Board does not intend for these rules to be subverted or misused by participants in legal proceedings as a procedural weapon to intimidate or harass a physician expert or to delay or otherwise complicate the administration of justice.

The Mississippi State Board of Medical Licensure shall provide a copy of these rules, with these Comments appended, to the Mississippi Supreme Court, the Mississippi Court of Appeals, the respective conferences of the Mississippi Circuit, Chancery, and County Judges, the Administrative Office of the Courts, the Mississippi Attorney General, the United States District Courts and United States attorneys located in Mississippi, the Mississippi Workers’ Compensation Commission, the Mississippi Bar Association, the Mississippi State Medical Association, the Federation of State Medical Boards, and any other appropriate person or organization at the discretion of the Board’s Executive Director, with the request that those organizations give notice to their members or other interested parties of the existence of these rules.
to practice medicine in this state, or by a licensed nurse under the direction and supervision of a licensed physician.


Rule 9.2 Position. It is the position of the Mississippi State Board of Medical Licensure that vaccinations administered pursuant to a community-based public immunization program are considered to be under the direction and supervision of a physician, and thus do not constitute the unlawful practice of medicine, when all of the following criteria are met:

A. the vaccinations are administered to the public by a licensed provider who is:
   1. authorized under Mississippi statute or regulation to provide vaccinations and is
   2. subject to the regulation of a Mississippi regulatory agency.

B. The vaccinations are carried out pursuant to state and federal public health immunization programs or other programs which:
   1. shall be approved in advance by the Board;
   2. shall be conducted under the general supervision of a physician
      a. licensed in the state of Mississippi,
      b. who actively practices medicine at least 20 hours/week, and
      c. resides in the state of Mississippi; and,
   3. a single physician assumes responsibility for the safe administration of the vaccine.


Part 2635 Chapter 10: Release of Medical Records

Rule 10.1 | Definitions

For the purpose of Part 2635, Chapter 10 only, the following terms have the meanings indicated:

A. “Licensee” means any person licensed to practice by the Mississippi State Board of Medical Licensure (the “Board”).

B. “Medical Records” means all records and/or documents relating to the treatment of a patient, including, but not limited to, family histories, medical histories, report of clinical findings and diagnosis, laboratory test results, x-rays, reports of examination and/or evaluation, billing records, and any hospital admission/discharge records which the licensee may have, or which is otherwise maintained by the group or facility wherein said licensee practices medicine.

C. “Patient” means any person who receives or should have received health care from a licensee, under a contract, express or implied, whether or not the licensee is compensated for services rendered.

D. “Legal Representative” means an attorney, guardian, custodian, or in the case of a deceased patient, the executor/administrator of the estate, surviving spouse, heirs and/or devisees.¹

E. “Authorized Requesting Party” includes patient and legal representative as defined above who holds a valid written release and authorization.

¹ See Miss. Code Ann., §41-10-3 for further authority and information.
Rule 10.2 | Medical Records - Property of Licensee

Medical records, as defined herein, are and shall remain the property of the licensee in whose facility said records are maintained, subject to reasonable access to the information by authorized individuals or entities.

In the case of employed or contracted licensees (those lacking authority to manage or maintain medical records), medical record ownership shall be determined by federal and state statutes and regulations. Licensees in such relationships shall make reasonable efforts to assure reasonable access to the information by authorized individuals or entities. Further, licensees should inform patients of procedures for release of records if the licensee is not the custodian of the records.

Rule 10.3 | Regulatory and Legal Requests

The Board has the authority to investigate licensees as part of its mission to protect the public. Further, continued licensure by the Board requires the production of medical records when requested. When provided an administrative (i.e., legal) request for in-person inspection or production of copies for removal by the Board, licensees shall comply and provide all records as requested.

Rule 10.4 | Transfer of Patient Records to Another Licensee

A licensee shall not refuse for any reason to make the information contained in the medical records available upon valid request by authorized requesting party to another licensee presently treating the patient. The licensee has a right to request a written release from the patient or legal representative of the patient, authorizing the transfer prior to transfer of said documents. Upon receipt of the written release and authorization, the licensee must tender a copy of said documents to the other licensee within a reasonable period of time. Transfer of said documents shall not be withheld because of an unpaid bill for medical services, but the licensee is entitled to reasonable compensation paid in advance for any copy expenses as provided in Part 2635, Rule 10.6.

Rule 10.5 | Release of Patient Records to Patient

A licensee shall, upon request of authorized requesting party holding a written release and authorization, provide a copy of a patient's medical record to the authorized requesting party within a reasonable period of time.

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2 Miss. Code Ann., §73-43-11

3 30 Miss. Admin. Code Pt.2640, R.1.4 Patient Record
In those cases where release of psychiatric/psychological records directly to a patient would be deemed harmful to the patient's mental health or well-being, the licensee shall not be obligated to release the records directly to the patient, but shall, upon request, release the records to the patient's legal representative. The licensee has a right to request a written authorization prior to release of the records to any party other than the patient. Upon receipt of the written release and authorization, the licensee must tender a copy of the records to the authorized requesting party within a reasonable period of time. Transfer of the records shall not be withheld because of an unpaid bill for medical services, but the licensee is entitled to reasonable compensation paid in advance for any copy expenses as provided in Part 2635, Rule 10.6.


Rule 10.6 | Narrative Summary of Medical Record

In some cases, a requesting party may wish to obtain a narrative summary of the medical record, in lieu of, or in addition to a copy of the medical record. Upon such a request, the licensee may provide the narrative summary. The licensee may charge a reasonable fee for the time devoted to preparation of the medical record narrative summary.


Rule 10.7 | Duplication and Administrative Fees

A. Licensees have a right to be reimbursed for duplication and other expenses relating to requests for medical records. The copying charge is set by Mississippi Code, Section 11-1-52 as follows:

1. Any medical provider or hospital or nursing home or other medical facility shall charge no more than the following amounts to an authorized requesting party for photocopying any patient's records:
   i. Twenty Dollars ($20.00) for pages one (1) through twenty (20);
   ii. One Dollar ($1.00) per page for the next eighty (80) pages;
   iii. Fifty Cents (50¢) per page for all pages thereafter.
   iv. Ten percent (10%) of the total charge may be added for postage and handling.
   v. Fifteen Dollars ($15.00) may be recovered by the medical provider or hospital or nursing home or other medical facility for retrieving medical records in archives at a location off the premises where the facility/office is located.
   vi. In addition, the actual costs of reproducing x-rays or other special records may be included.
   vii. The duplication and administrative fees authorized herein are not intended to include or restrict any fees charged in relation to expert testimony.


Rule 10.8 | Exclusion

Federal or state agencies providing benefit programs as well as contractual third-party payers and administrators are excluded from the above stated fees. Records that are requested by state or federal agencies as well as contracted payers and administrators may be billed at rates established
by those payers and contracts. The release of records as requested by state or federal agencies or third-party payers and administrators may not be refused for failure to pay required fees.


Rule 10.9 Violation of Rules

A refusal by a licensee to release patient records shall constitute unprofessional conduct, dishonorable or unethical conduct likely to deceive, defraud or harm the public in violation of Mississippi Code, Section 73-25-29(8)(d).


Part 2635 Chapter 11: Withdrawn March 16, 2017

Part 2635 Chapter 12: Physician Advertising

Rule 12.1 Scope. The following rule on physician advertising applies to all individuals licensed to practice medicine, osteopathic medicine or podiatric medicine in the state of Mississippi.


Rule 12.2 Definitions. For the purpose of Part 2635, Chapter 12 only, the following terms have the meanings indicated:

A. “Board” means the Mississippi State Board of Medical Licensure.
B. “Physician” means any individual licensed to practice medicine, osteopathic medicine or podiatric medicine in the state of Mississippi.
C. “Advertisement” or “Advertising” means any form of public communication, such as office signage, newspaper, magazine, telephone directory, medical directory, radio, television, direct mail, billboard, sign, computer, business card, billing statement, letterhead or any other means by which physicians may communicate with the public or patients.


Rule 12.3 Requirements.

A. Subject to the requirements set forth herein below, any advertisement by a physician may include:
   1. The educational background or specialty of the physician.
   2. The basis on which fees are determined, including charges for specific services.
   3. Available credit or other methods of payment.
   4. Any other non-deceptive information.
B. A physician may publicize himself or herself as a physician through any form of advertisement, provided the communication, (i) shall not be misleading because of the
omission of necessary information, (ii) shall not contain any false or misleading statement, or (iii) shall not otherwise operate to deceive.

C. Because the public may be deceived by the use of medical terms or illustrations that are difficult to understand, physicians should design the advertisement to communicate the information contained therein to the public in a readily comprehensible manner.

D. It is unethical to advertise in such a manner as to create unjustified medical expectations by the public. The key issue is whether advertising or publicity is true and not materially misleading.

E. In addition to the above general requirements, any advertisement or other form of public communication shall comply with the following specific requirements:

1. All advertisements and written communications pursuant to these rules shall include the name of at least one (1) physician responsible for its content. In the case of office signage at least one sign in reasonable proximity to the main entrance must bear the name of the responsible physician.

2. Whenever a physician is identified in an advertisement or other written communication, the physician should not be identified solely as “Doctor” or “Dr.” but shall be identified as M.D. for medical doctors, D.O. for osteopathic physicians and D.P.M. for podiatric physicians.

3. A physician who advertises a specific fee for a particular service or procedure shall honor the advertised fee for at least ninety (90) days unless the advertisement specifies a longer period; provided that for advertisements in the yellow pages of a telephone directory or other media not published more frequently than annually, the advertised fee shall be honored for no less than one (1) year following publication.

4. A physician shall not make statements which are merely self-laudatory or statements describing or characterizing the quality of the physician’s services.

5. No physician shall advertise or otherwise hold himself or herself out to the public as being “Board Certified” without, (i) a complete disclosure in the advertisement of the specialty board by which the physician was certified, and (ii) can submit proof of current certification by a specialty board recognized by the American Board of Medical Specialties or the American Osteopathic Association. The term “Board Certified” frequently appears in conjunction with a list of services that the physician or clinic provides. The general public could easily be misled into thinking that the physician is certified in all of those services.

6. No physician shall hold himself or herself out as a specialist in a particular field unless that physician has either, (i) completed a residency program recognized by the Accreditation Council for Graduate Medical Education, by the American Osteopathic Association or by the American Podiatric Medical Association and can submit proof that such training was completed, or (ii) can submit proof that the licensee was “grandfathered” into a specialty by board certification by a recognized specialty board of the American Board of Medical Specialties or the American Osteopathic Association.

7. No physician shall compare his or her service with other physicians' services, unless the comparison can be factually substantiated; this precludes the use of terms such as “the best,” “one of the best,” or “one of the most experienced” or the like.

8. Where an advertisement includes a consumer-endorser's experience (i.e., patient testimonials), the advertisement must contain clear and prominent disclosure of (a)
what the generally expected outcome would be in the depicted circumstances, and (b) the limited applicability of the endorser's experience. Although testimonials and endorsements are authorized under this rule, compliance will be strictly monitored as endorsements and testimonials are inherently misleading to the lay public and to those untrained in medicine.

9. Any claims of success, efficacy or result (i.e., cure) must have scientific evidence in substantiation of such claims.

10. Any claims that purport to represent “typical” results (results that consumers will generally achieve) must be based on a study of a sample of all patients who entered the program, or, if the claim refers to a subset of those patients, a sample of that subset.

11. Any claim made regarding the safety of a medical procedure or drug must also disclose the risk of adverse medical complications.

12. No physician shall claim to have any drug or medication or use of a drug or medication for a specific ailment or condition unless such drug or medication has an F.D.A. approved indication for such purpose.

13. Any claim that improvements can be achieved through surgery in a specified time period must also include disclosure of the typical recovery time.

F. Consistent with federal regulatory standards which apply to commercial advertising, a physician who is considering the placement of an advertisement or publicity release, whether in print, radio or television, should determine in advance that the communication or message is explicitly and implicitly truthful and not misleading. These standards require the advertiser to have a reasonable basis for claims before they are used in advertising. The reasonable basis must be established by those facts known to the advertiser, and those which a reasonable, prudent advertiser should have discovered.

G. The above rules do not prohibit physicians or clinics from authorizing the use of the physician's name or clinic name in medical directories, HMO directories, preferred provider agreements or other communications intended primarily for referral purposes.


Rule 12.4 Violation of Rules. The above rules on physician advertising shall not be interpreted to alter or amend that which is otherwise provided by Mississippi statutory law or the rules on advertising adopted by the Federal Trade Commission.

If any physician subject to this rule advertises or enters into any communication in violation of the above rules, such act shall constitute unprofessional conduct, which includes dishonorable or unethical conduct likely to deceive, defraud or harm the public, in violation of Mississippi Code, Sections 73-25-29(8)(d) and 73-27-13(h)(iv).


Part 2635: Chapter 13: Complementary and Alternative Therapies

Rule 13.1 | Scope and Purpose
Rule 13.2 | Definitions

For the purpose of Part 2635, Chapter 13 only, the following terms have the meanings indicated:

A. “Board” means the Mississippi State Board of Medical Licensure.

B. “Complementary”, “Alternative”, and “Regenerative Medicine/Therapy” means those health care methods of diagnosis, treatment, or interventions that are not acknowledged to be conventional but that may be offered by some licensed physicians in addition to, or as an alternative to, conventional medicine. Examples of these therapies include, but are not limited to: IV infusion/hydration therapy, oriental medicine techniques and practices other than Licensed Acupuncture, utilization of Artificial Intelligence, and stem cell therapy.

C. “Conventional Medical Practices” means those medical interventions that are taught extensively at U.S. medical schools, generally provided at U.S. hospitals, or meet the requirements of the generally accepted standard of care.

D. “Informed and Shared Decision Making” means the process by which a physician discusses, in the context of the use of complementary, alternative, and/or regenerative therapies, the risks and benefits of such treatment with the patient. The patient is given an opportunity to express preferences and values before collaboratively evaluating and arriving at treatment decisions.

E. “Informed Consent” means evidence documenting appropriate patient consent to a therapy or procedure.

F. “Unproven Intervention” means any therapy that lacks compelling evidence, based upon scientific studies, to validate its treatment efficacy.

Rule 13.3 | Alternative Medicine Practices

The Board is aware that a growing number of licensees and patients are both implementing and seeking complementary and alternative medicine in their health care. Further, the Board recognizes that innovative practices that could benefit patients and improve care should be given reasonable and responsible degrees of latitude.

In reviewing this subject, the Board is also aware of the fact that consumer fraud occurs across the country, and, unfortunately, not infrequently in the practice of medicine. If consumer protection means anything, it should protect people weakened by illness from the dangers attendant to

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4 Regulations regarding Licensed Acupuncture can be found at Title 30, Part 2625 The Practice of Acupuncture
unsound, invalidated, and/or otherwise unsubstantiated practices. Licensees should never agree to perform invalidated or unsound treatments or therapies.

The Board feels that licensees may incorporate alternative therapies if research results are promising, and only if the methods utilized are reasonably likely to benefit patients without undue risk. A full and frank discussion of the risks and benefits of all medical practices is expected, and is in the patient’s best interest.

Licensees should practice pursuant to informed and shared decision making when determining the utilization of complementary therapies. This style of process is conducive to openly weighing the risks and benefits of the therapies under consideration. While this process is ideal, the licensee is ultimately responsible for the decision-making process.

Where evidence is unavailable for a particular treatment in the form of clinical trials or case studies, licensees must only proceed with an appropriate rationale for the proposed treatment, and justification of its use, in relation to the patient’s symptoms or condition. Novel, experimental, and unproven interventions should only be proposed when traditional or accepted proven treatment modalities have been exhausted. In such instances, there must still be a basis in theory or peer-acknowledged practice. The burden rests solely on the licensee in regard to the substantiation supporting the use of a particular therapy. Licensees should be prepared to support any claims made about benefits of treatments or devices with documented evidence, for example with studies published in peer-reviewed publications.

Licensees must refrain from charging excessive fees for treatments provided. Further, licensees should not recommend, provide, or charge for unnecessary medical services, nor should they make intentional misrepresentations to increase the level of payment they receive.\(^5\)

*Source: Miss. Code Ann. §73-43-11 (1972, as amended).*

**Rule 13.4 | Informed Consent**

Licensees who choose to utilize alternative therapies must obtain written informed consent from the patient prior to the utilization of said therapies. Said informed consent consists of the following elements:

1. The patient, the licensee, and the credentials of the licensee are all identified;
2. The types of transmissions regarding the therapy are identified (e.g., prescription refills, appointment scheduling, patient education, etc.);
3. Overt agreement from the patient with the licensee’s determination about whether or not the condition being diagnosed and/or treated is appropriate for alternative therapy;
4. Express patient consent to forward patient-identifiable information to a third party, if necessary;
5. An accurate description of the benefits and risks of treatment or intervention, based on scientific evidence, as well as an explanation of alternatives to treatment or an intervention, and the right to withdraw from treatment or an intervention without denial of standard of care to patients.

\(^5\) American Medical Association, *Code of Medical Ethics*, Opinion 11.3.1.
Rule 13.5 | Evaluation

Parity of evaluation standards should be established for patients, whether the licensee is using conventional medical practices or alternative therapy.

Prior to offering any recommendations for conventional and/or alternative treatments, the physician shall conduct an appropriate medical history and physical examination of the patient, as well as an appropriate review of the patient’s medical records. This evaluation shall include, but is not limited to, conventional methods of diagnosis, and may include other methods of diagnosis as long as the methodology utilized for diagnosis is based upon the same standards of safety and reliability as conventional methods, and shall be documented in the patient’s medical record. The record should also document the following:

1. What medical options have been discussed, offered or tried, and if so, to what effect, or a statement as to whether or not certain options have been refused by the patient or guardian;
2. That proper referral has been offered for appropriate treatment;
3. That the risks and benefits of the use of the recommended treatment, to the extent known, have been appropriately discussed with the patient or guardian; and
4. That the licensee has determined the extent to which the treatment could interfere with any other recommended or ongoing treatment.

Rule 13.6 | Treatment Plan

A documented treatment plan tailored to the individual needs of the patient by which treatment progress or success can be evaluated with stated objectives, such as pain relief and/or improved physical and/or psychosocial function. Said treatment plan must consider pertinent medical history, previous medical records and physical examination, as well as the need for further testing, consultations, referrals or the use of other treatment modalities.

The treatment offered shall meet the following criteria:

1. A favorable risk/benefit ratio compared to other treatments for the same condition;
2. Be based upon a reasonable expectation that it will result in a favorable patient outcome, including preventive practices;
3. Be based upon the expectation that a greater benefit will be achieved than that which can be expected with no treatment.

Rule 13.7 | Medical Records

Any licensee who provides alternative therapy as a component of practice must, as with all other forms of practice, maintain a complete record which substantiates the care provided. Said record shall, at a minimum, include the following:

1. The medical history and physical examination(s);
2. Diagnostic, therapeutic and laboratory results;
3. Results of evaluations, consultations and referrals;
4. Treatment objectives;
5. Discussion of risks and benefits;
6. Appropriate informed consent;
7. Treatments;
8. Medications (including date, type, dosage and quantity prescribed);
9. Instructions and agreements; and
10. Periodic reviews

Records shall be current and maintained in an accessible manner, and readily available for review and inspection.


Rule 13.8 | Education

All licensees who offer alternative therapies must be able to demonstrate knowledge and understanding of the medical and scientific knowledge connected with any method they are offering or using in their medical practices as a result of related education and training. In order to implement best practices for alternative therapies, licensees must understand the relevant clinical issues and shall obtain sufficient targeted continuing medical education and training.


Rule 13.9 | Advertising

As to the advertising of alternative therapies, data purportedly supporting unproven interventions commonly undermines information about risks and overemphasizes information about benefits. Information presented in advertising, including but not limited to clinic websites and social media, shall be represented accurately and come from reputable peer-reviewed publications or respected external organizations.

Even where an appropriate informed consent process seems to be in place, deceptive or fraudulent information contained within practice advertising, websites, and other marketing materials could mislead patients into consenting to treatment, thereby invalidating the informed consent process.

Treatment options described and accompanied by supporting information in the form of journal articles, patient testimonials, claims of partnerships with academic institutions, mentions of affiliations with professional societies or networks, statements regarding receipt of FDA approval or explicit mention of exemption from FDA oversight, listings of patents granted, statements that clinical trials of investigational interventions are being conducted, and accolades related either to the practice itself or its affiliated physicians and researchers, which serve to exaggerate, inflate, or misrepresent information derived from legitimate or questionable sources, shall be deemed a
violation of the Board’s advertising regulations and unprofessional conduct likely to deceive, defraud, or harm the public.

Although not all-encompassing, the following represents instances of improper or misleading advertising practices which the Board would consider unprofessional and deceptive in nature:

1. Asserting certification of products or practices by international standards organizations or claiming training certification, in order to legitimize alternative therapies;
2. Convening scientific or medical advisory boards featuring prominent business leaders and academic faculty members in order to legitimize alternative therapies;
3. Registering trials whose apparent purpose is solely to attract patients willing to pay to participate in them;
4. Using the statement or impression of “ethics review” to convey a sense of legitimacy to products or procedures;
5. Renting of laboratory or business space within a legitimate scientific or government institution in order to legitimize alternative therapies;
6. Using membership in established academic or professional societies to suggest legitimacy by association;
7. Publication of open-ended voluntary monitoring data sets rather than undertaking controlled clinical trials;
8. Suggesting that patent applications or grants indicate clinical utility rather than initiation of an application process or recognition of novelty and inventiveness;
9. Publishing research and commentary in journals with limited anonymous peer review;
10. Citing preclinical and other research findings to justify clinical application without sufficient efficacy testing in humans;
11. Forming organizations to self-regulate in ways that support premature commercialization; and
12. Providing expert opinions or celebrity comments on unsupported clinical uses or standing of the provider.


Rule 13.10 | Violation of Rules

The use of alternative, complementary, and/or regenerative therapies outside the requirements and regulations stated herein constitutes unprofessional conduct, dishonorable or unethical conduct likely to deceive, defraud or harm the public, in violation of Miss. Code Ann., § 73-25-29(8)(d).


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6 Title 30, Part 2635 Chapter 12: Physician Advertising

7 Miss. Code Ann., §73-25-29(8)(d)
Part 2635: Chapter 14: Temporary Practice by an Athletic Team Physician

Rule 14.1 | Scope and Purpose

The purpose of this regulation is to set forth certain exemptions and stipulations as to the practice of medicine within Mississippi by physicians travelling from out of state with sports teams for sporting events conducted within the state. Further, it is the intent of this regulation to sort forth the requirements of those physicians to practice medicine in Mississippi, temporarily, without obtaining Mississippi licensure.


Rule 14.2 | Definitions

For the purpose of Part 2635, Chapter 14 only, the following terms have the meanings indicated:

A. “Athletic Team” or “Team” means a group of people representing a specific organization engaged in sporting activities, such as baseball or football, which require medical personnel to treat or evaluate injuries sustained pursuant to the activity.

B. “Staff Members” means those individuals directly affiliated with the sporting program or entity whose purpose is to support the players or members of the team during the event. This includes, but is not necessarily limited to: trainers, coaches, equipment personnel, communications staff, band members, cheerleaders, and the team mascot. This would not include parents, boosters, or other individuals simply present or attending the activity or sporting event.

C. “Team Physician” means those health care professionals, holding an unrestricted medical license in their athletic team’s state of origin, who travel with their team to away games/events for the purposes of providing medical treatment and evaluation for players and staff members of said team.


Rule 14.3 | Athletic Team Physicians

As part of any sport, teams require the presence of trained medical personnel, to include physicians, in order to treat injuries incurred during the course of the activity. As such, when athletic teams travel to away games or events outside their respective state, said medical personnel routinely travel with the team to provide said care.

Understanding these principles of athletics, a physician licensed in another state, territory or jurisdiction of the United States is exempt from the licensure requirements in Mississippi under the following conditions related to athletic team based practice:

i) The physician is employed or formally designated as the team physician by an athletic team visiting Mississippi for a specific sporting event;

ii) The physician limits the practice of medicine in Mississippi to medical treatment of the members, coaches and staff, as defined herein, of the sports entity that employs or has designated the physician and;

iii) Said physician is licensed in the state the sports entity or organization is based or housed.
Additionally, physicians authorized to practice under this rule may also treat members from the home team in Mississippi if said physician has specialized training or experience beyond that of the home team physician.

The extent of the medical practice allowed under this rule is limited to the following aspects of the game or event:

- Pre-game warm-up and any postgame activities;
- During the actual game or event;
- Travel to and from the sporting event within Mississippi; and
- In-state lodging of the team and other covered staff.

Further, it is the responsibility of the team or organization employing the physician to verify said physician is licensed and in good standing in the appropriate jurisdiction as required under this rule.


Rule 14.4 | Violation of Rules

The practice of medicine outside of the requirements and regulations stated herein constitutes the illegal practice of medicine, in violation of Miss. Code Ann., §97-23-43, and violators shall be subject to all fines and penalties described therein.


Part 2635 Chapter 15: Hospice Practice

Rule 15.1 In-Home Hospice Good Faith

Recognizing the unique team-based approach utilized when treating in-home hospice patients, the following represents four factors required to establish a proper physician-patient relationship:

i) The medical director must receive an order from the treating/referring physician requesting the patient be admitted for hospice care. Self-referral by the physician medical director may be necessary, and on those occasions, a second physician must be consulted to affirm the decision for hospice admission. Physician Medical Directors who self-refer a patient to their hospice, or to any hospice with whom the director has a contractual relationship, must obtain informed consent from the patient. Additionally, Physician Medical Directors must disclose to the primary care provider for the patient, in writing, that the patient has been admitted to hospice;

ii) That the treating hospice physician or medical director has thoroughly reviewed the medical records of the patient, as provided by the referring physician, has documented the review, and has determined just cause exists for hospice admission (expected death
in six months or less), with documented follow-up review at every certification period thereafter;

iii) That the actions of the physician are deemed within the course of legitimate professional practice, as defined by the Centers for Medicare and Medicaid Services (CMS); and

iv) That an evaluation of the patient occurs no later than thirty (30) days after the admission of the patient to hospice. The evaluation shall consist of either a face to face with the physician, face to face with a mid-level provider (PA or APRN), or a telemedicine visit by the medical director with nursing support in the home. Regardless of how the evaluation is accomplished, the author of any controlled substance prescriptions must have evaluated the patient within the thirty (30) day time-period.

It shall be considered unprofessional conduct for a medical director to participate in active recruitment for patient admission to hospice. For the purposes of this regulation, the term “active recruitment” shall mean any unsolicited interaction with a patient for the purposes of convincing a patient to enroll in hospice. As an example: having hospice staff or affiliates visit nursing home patients, with whom the physician has no prior relationship, for the ultimate purpose of soliciting their enrollment in hospice.

It shall be considered unprofessional conduct for physicians to document participation at Inter-Disciplinary Group (IDG) meetings when they did not attend the meeting(s).

Nothing in this section shall preclude a hospice physician from fulfilling their duties to provide physician services as needed to hospice patients.


Part 2635: Chapter 16: Medical Examiners

Rule 16.1 | Scope and Purpose

The purpose of this regulation is to set forth certain exemptions, stipulations, and expectations as to the practice of medicine within Mississippi by physicians who serve as the State Medical Examiner or a Deputy Medical Examiner. Further, it is the intent of this regulation to set forth the requirements of those physicians to practice medicine in Mississippi, temporarily, without obtaining an unrestricted Mississippi medical license. The Board defers to state statute on any duties or requirements not specifically mentioned within this regulation.


Rule 16.2 | Definitions

8 As defined in The Social Security Act, Title 18, §1861 (dd)(2)(B), as amended.
For the purpose of Part 2635, Chapter 15 only, the following terms have the meanings indicated:

A. “Medical Examiner” means the person appointed by the Commissioner of Public Safety pursuant to Miss. Code Ann., §41-61-55 to investigate and certify deaths that affect the public interest.

B. “Deputy Medical Examiner” means those professional individuals employed by The Department of Public Safety who serve under the direction of the Medical Examiner, and who perform autopsies and post-mortem examinations to determine cause of death via medical processes, such as pathology, and who may testify as an expert regarding their findings.


Rule 16.3 | Temporary Practice

Recognizing the unique challenges in hiring and retaining Deputy Medical Examiners, along with the need to expeditiously conduct autopsies in order to avoid evidentiary spoilage, applicants for licensure to serve in the role of Deputy Medical Examiner may practice within Mississippi temporarily, without an unrestricted medical license, while going through the licensure process. Said physicians must first submit their application, thereby starting the licensure process, and must verify they are licensed in good standing in another state or acceptable jurisdiction. This temporary practice period shall not exceed six (6) months from the date the application is received.

Further, contract physicians who are hired on a temporary basis by The Department of Public Safety may also practice without a license, after verifying their unrestricted licensure as described above, for a period of up to one (1) month. Thereafter, said physicians must apply for a full license in Mississippi.


Rule 16.4 | Violation of Rules

The practice of medicine outside of the requirements and regulations stated herein constitutes the illegal practice of medicine, in violation of Miss. Code Ann., §97-23-43, and violators shall be subject to all fines and penalties described therein.