

TITLE 16
CHAPTER 2
PART 19 **OCCUPATIONAL AND PROFESSIONAL LICENSING**
 ACUPUNCTURE AND ORIENTAL MEDICINE PRACTITIONERS
 EXPANDED PRACTICE CERTIFICATIONS

16.2.19.1 **ISSUING AGENCY:** New Mexico Board of Acupuncture and Oriental Medicine.
[16.2.19.1 NMAC - N, 11-28-09]

16.2.19.2 **SCOPE:** All doctors of oriental medicine who are certified for expanded practice or who are applicants for certification for expanded practice, as well as all educational programs and students enrolled in an educational program.
[16.2.19.2 NMAC - N, 11-28-09]

16.2.19.3 **STATUTORY AUTHORITY:** This part is promulgated pursuant to the Acupuncture and Oriental Medicine Practice Act, Section 6-1-14.A-8.1.
[16.2.19.3 NMAC - N, 11-28-09]

16.2.19.4 **DURATION:** Permanent.
[16.2.19.4 NMAC - N, 11-28-09]

16.2.19.5 **EFFECTIVE DATE:** November 28, 2009, unless a later date is cited at the end of a section.
[16.2.19.5 NMAC - N, 11-28-09]

16.2.19.6 **OBJECTIVE:** This part lists the certification requirements for each of the following expanded practice categories: basic injection therapy, injection therapy, intravenous therapy and bioidentical hormone therapy.
[16.2.19.6 NMAC - N, 11-28-09]

16.2.19.7 **DEFINITIONS:**

- A. The ~~definition~~ [in this section] ~~is~~ ~~are~~ in addition to those in the act and 16.2.1 NMAC.
- B. ~~[The following definition applies to the rules and the act: "educational course" is a comprehensive foundation of studies, approved by the board leading to demonstration of entry-level competence in the specified knowledge and skills required for the four respective certifications in expanded practice; an educational course is not an educational program as this term is used in the act and the rules and as defined in 16.2.1 NMAC.] [The following definitions are from 16.19.36 NMAC for clarification of regulations for doctors of oriental medicine, certified in expanded practice;~~
1. ~~_____~~ "Aseptic Technique" means proper manipulation of preparations to maintain sterility.
 2. ~~_____~~ "ASHP" American Society of Health-Systems Pharmacists.
 3. ~~_____~~ "Beyond-use date" (BUD) means the date, or an appropriate, date and time, after which a compounded preparation is not to be used and is determined from the date and time the preparation is compounded.
 4. ~~_____~~ "Certification" means independent third party documentation declaring that the specific requirements have been met.
 5. ~~_____~~ "Cleanroom" means a room in which the concentration of airborne particles is controlled to meet a specified airborne particulate cleanliness class. Microorganisms in the environment are monitored so that a microbial level for air, surface, and personnel gear are not exceeded for a specified cleanliness class.
 6. ~~_____~~ "Compounded sterile preparations" (CSP's) include, but are not limited, to the following dosage forms which must be sterile when administered to patients:
 - (a) ~~_____~~ parenteral preparations;
 - (b) ~~_____~~ aqueous bronchial and nasal inhalations;
 - (c) ~~_____~~ injections (e.g. colloidal dispersions, emulsions, solutions, suspensions);
 - (d) ~~_____~~ irrigations for wounds and body cavities;
 - (e) ~~_____~~ ophthalmic drops and ointments; and
 7. ~~_____~~ "Compounding aseptic isolator" (CAI) means an enclosed ISO Class 5 environment for compounding pharmaceutical ingredients or preparations. It is designed to maintain an aseptic compounding environment within the isolator throughout the compounding and material transfer processes. Air exchange into the isolator from the surrounding environment should not occur unless the air has first passed through a microbial retentive filter (HEPA minimum).
 8. ~~_____~~ "Critical area" means an ISO Class 5 environment.
 9. ~~_____~~ "Critical site" means a location that includes any component or fluid pathway surfaces (e.g., vial septa, injection ports, beakers) or openings (e.g., opened ampules, needle hubs) exposed and at risk of direct contact with air (e.g., ambient room or HEPA filtered), moisture (e.g., oral and mucosal secretions), or touch contamination. Risk of microbial particulate contamination of the critical site increases with the size of the openings and exposure time.

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Chapter 1

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