of the United States, or official National Formulary or any supplement of them; (ii) articles intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in man or animals; (iii) articles (other than food) intended to affect the structure or any function of the body of man or animals; (iv) articles intended for use as a component of any article specified in (i), (ii) or (iii) of this subsection. The term "drug" however, does not include devices or their components, parts or accessories.

(4) The term "counterfeit drug" means a drug, or container or labeling of a drug which without authorization bears the trademark, trade name, or other identifying mark, imprint or device (or any likeness thereof), of a drug manufacturer, processor, packer, or distributor other than the person or persons who in fact manufactured, processed, packed or distributed such drug, and which thereby is falsely represented or purported to be the product of, or to have been packed or distributed by, such other drug manufacturer, processor, packer or distributor.

(5) The term "depressant or stimulant drug" means (i) any drug containing any quantity of barbituric acid, any salt of barbituric acid or any derivation of barbituric acid which has been designated under the provisions of the Federal Drug Act as habit forming; (ii) any drug containing any quantity of amphetamine or any of its optical isomers, any salt of amphetamine or any salt of an optical isomer or amphetamine; (iii) any substance designated under the provisions of the Federal Drug Act as habit forming because of its stimulant effect on the central nervous system; (iv) any drug containing any quantity of any substance designated under the provisions of the Federal Drug Act as having potential for abuse due to its hallucinogenic effect or its depressant or stimulant effect on the central nervous system. ; BUT SHALL NOT MEAN ANY NARCOTIC DRUG AS DEFINED BY SECTION 276 OF ARTICLE 27 OF THIS CODE.

(6) The term "manufacture, compound or process" includes repackaging or otherwise changing the container, wrapper, or labeling of any drug package, in the furtherance of the distribution of the drug from the original place of manufacture to the person making final delivery or sale to the ultimate consumer. "Manufacturers, compounders, and processors" refer to any persons engaged in activities defined under this subsection.

(7) The term "practitioner" means physician, dentist, veterinarian or other person licensed in this State to prescribe or administer drugs subject to this section.

(8) The term "Federal Drug Act" means the Federal Food, Drug and Cosmetic Act 52 Stat. 1040 (1938), 21 U.S.C. Sections 301-392, as amended from time to time.

(b) No one but the following persons shall manufacture, compound or process any depressant or stimulant drug, except those exempted under Section 511(f) of the Federal Drug Act, in this State:

(1) Manufacturers, compounders, and processors regularly engaged in preparing pharmaceutical chemicals or prescription drugs for distribution to pharmacies, hospitals, clinics, public health agencies, physicians, laboratories, research or educational institu-