(12) If it is, or purports to be, or is represented as a drug composed wholly or partly of insulin, unless (a) it is from a batch with respect to which a certificate or release has been issued pursuant to Section ~~506~~ 356 of the Federal Act, and (b) such certificate or release is in effect with respect to such drug.

(13) If it is, or purports to be, or is represented as a drug composed wholly or partly of any kind of penicillin, streptomycin, chlortetracycline, chloramphenicol, bacitracin, or any other antibiotic drug, or any derivative thereof, unless (a) it is from a batch with respect to which a certificate or release has been issued pursuant to Section ~~507~~ 357 of the Federal Act, and (b) the certificate or release is in effect with respect to the drug; provided, that this subsection shall not apply to any drug or class of drugs exempted by regulations promulgated under Section ~~507~~ 357 (c) or (d) of the Federal Act. For the purpose of this subsection the term "antibiotic drug" means any drug intended for use by man containing any quantity of any chemical substance which is produced by microorganisms and which has the capacity to inhibit or destroy microorganisms in dilute solution (including, the chemically synthesized equivalent of the substance).

(14) If it is a color additive, the intended use of which in or on drugs is for the purpose of coloring only, unless its packaging and labeling are in conformity with such packaging and labeling requirements applicable to such color additive, prescribed under the provisions of Section 188F (b) of this subtitle or under the provisions of the Federal Act.

(15) In the case of any prescription drug distributed or offered for sale in this State, unless the manufacturer, packer, or distributor thereof includes in all advertisements and other descriptive printed matter issued or caused to be issued by the manufacturer, packer or distributor with respect to that drug a true statement of (a) the established name, as defined in paragraph (6) of this section, printed prominently and in type at least half as large as that used for any trade or brand name thereof, (b) the formula showing quantitatively each ingredient of the drug to the extent required for labels under Section ~~502~~ 352 (e) of the Federal Act, and (c) such other information in brief summary relating to side effects, contraindications, and effectiveness as shall be required in regulations issued under the Federal Act.

(16) If a trademark, trade name or other identifying mark, imprint or device of another or any likeness of the foregoing has been placed thereon or upon its container with intent to defraud.

(17) Drugs and devices which are, in accordance with the practice of the trade, to be processed, labeled or repacked in substantial quantities at establishments other than those where originally processed or packed shall be exempt from any labeling or packaging requirements of this subheading; provided, that such drugs and devices are being delivered, manufactured, processed, labeled, repacked or otherwise held in compliance with regulations issued by the Secretary, or under the Federal Act.

189C.

(a) A drug intended for use by man which: (i) is a habit-forming drug to which Section 189B (4) applies; or (ii) because