This subsection shall not apply to officers and employees of this State, or a political subdivision of this State or of the United States while acting in the course of their official duties; or to drug manufacturers, or their agents or employees authorized to possess stimulants or depressant drugs under the provisions of this subtitle while they are actually engaged in investigative activities directed toward safeguarding said drug manufacturer's trademark.

(f) No person shall make, sell, dispose of, or keep in possession, control or custody, or conceal any punch, die, plate, stone, or other thing designed to print, imprint, or reproduce the trademark, trade name, or other identifying mark, imprint, or device of another or any likeness of any of the foregoing upon any drug or container or labeling thereof so as to render such drug a counterfeit drug.

(g) (1) From and after July 1, 1968, every person engaged in manufacturing, compounding, processing, selling, delivering or otherwise disposing of any depressant or stimulant drug shall prepare a complete record of all stocks of each drug on hand and keep such record for a period of three (3) years. If such record has already been prepared in accordance with Section 511(d) of the Federal Drug Act, and has been retained and is available to the Department upon request, no additional record shall be required. Every person manufacturing, compounding or processing such drugs shall include in the record prepared the kind and quantity of each drug manufactured, compounded or processed and the date of such manufacture, compounding or processing. Every person selling, delivering, or otherwise disposing of such drugs shall include in the record prepared, the kind and quantity of each such drug received, sold, delivered, or otherwise disposed of, the name and address of the person from whom it was received, and to whom it was sold, delivered, or otherwise disposed of, along with the dates of such transactions.

(2) Persons required to prepare and keep records as provided in (g) (1) above, shall, upon request, permit officers and/or employees authorized by the Department to have access to and copy such records. Officers and/or employees properly authorized by the Department, may enter at reasonable times any factory, warehouse, establishment or vehicle in which any depressant or stimulant drug is held, manufactured, compounded, processed, sold, delivered or otherwise disposed of, to inspect, within reasonable limits in a reasonable manner, such factory, warehouse, establishment or vehicle, and all pertinent equipment, finished and unfinished material, containers and labeling therein, including records, files, papers, processes, controls and facilities, with the right to inventory any stock of any such drug and obtain samples of any such drug; provided, however, that the right of inspection shall not apply to financial data, sales data (other than shipment data), pricing data, personnel data or research data.

(3) The provisions of subsections (g) (1) and (2) shall not apply to a licensed practitioner as defined in (b) (5) with respect to depressant or stimulant drugs received, prepared, processed, administered or dispensed by him in the course of his professional practice, unless he regularly engages in dispensing such drugs to his patients for which they are charged, either separately or together with charges for other professional services.