- (e) Improve methods for preventing, producting, understanding and dealing with the misuse and abuse of controlled dangerous substances; and,
- (3) Enter into contracts with public agencies, institutions of higher education, and private organizations or individuals for the purpose of conducting research, demonstrations, or special projects which bear directly on misuce and abuse of controlled dangerous substances.
- (e) The Authority may enter into contracts for educational and research activities.

300. Prescription drugs:

- (a) "Prescription drugs" shall mean and include any drug intended for use by man which, because of its toxicity or other potentiality for harmful effect, or the method of its use, or the collateral measures necessary for its use, is designated by the Department as not safe for use except under the supervision of a practitioner licensed by law to administer such drugs. Provided that this term shall not mean any controlled dangerous substance as defined in this subheading.
- (b) Dispensed only on prescription: Any drug which bears a cautionary label warning against dispensing without a prescription under Federal law shall be dispensed only:
- (i) Upon the written prescription of a practitioner licensed by law to administer such drugs, or
- (ii) Upon the oral prescription of such practitioner, which shall be reduced to writing and filed by the pharmacist, or
- (iii) By refilling any such written or oral prescription if such refilling is authorized by the prescriber, either in the original prescription or by oral direction. Such authorization must be reduced to writing and filed by the pharmacist.
- (c) Label required: Any prescription drug dispensed by filling or refilling a written or oral prescription of a practitioner licensed by law to administer such drugs shall in addition to any requirements of the Department or Federal law bear a label containing the name and address of the dispenser, the serial number and date of the prescription, the name of the prescriber, and, if stated in the prescription, the name and address of the patient, and the directions for use as contained in the written or oral prescription.
- (d) Exempt sales: The provisions of this subheading shall not apply to sales of dangerous drugs made to registered practitioners of pharmacy, medicine, dentistry, or veterinary medicine, or to sales made by any manufacturer, wholesale druggist, or licensed pharmacist to another manufacturer, wholesale druggist, or licensed pharmacist or to a hospital or institution operating a dispensary in which a practitioner licensed by law to administer prescription drugs is in charge, providing records of such sales are maintained, and available for inspection, showing date of sale, name and address of purchaser, and quantity purchased.
 - (e) Applicability of subheading advertising drugs: