(6) IT HAS BEEN FOUND UNDER THE FEDERAL ACT OR BY THE SECRETARY TO BE A DRUG LIABLE TO DETERIORATION, AND:

(I) IT IS NOT PACKAGED IN THE FORM AND MANNER REQUIRED BY THE RULES AND REGULATIONS ADOPTED UNDER THE FEDERAL ACT OR BY THE SECRETARY; OR

(II) ITS LABEL DOES NOT BEAR A STATEMENT OF PRECAUTIONS AS REQUIRED BY THOSE RULES AND REGULATIONS;

(7) IT IS A PRESCRIPTION DRUG THAT WAS MANUFACTURED AFTER JULY 1, 1976 AND ITS LABEL DOES NOT BEAR THE NAME OF THE ACTUAL MANUFACTURER OF THE DRUG;

(8) ITS CONTAINER IS MADE, FORMED, OR FILLED IN A MANNER THAT IS MISLEADING;

(9) IT IS AN IMITATION OF ANOTHER DRUG;

(10) IT IS OFFERED FOR SALE UNDER THE NAME OF ANOTHER DRUG;

(11) IT IS OR IT IS PURPORTED TO BE A DRUG THAT IS COMPOSED IN WHOLE OR IN PART OF INSULIN AND IT IS NOT FROM A BATCH FOR WHICH A CURRENTLY UNEXPIRED CERTIFICATE OR RELEASE HAS BEEN ISSUED UNDER THE FEDERAL ACT;

(12) IT IS OR IT IS PURPORTED TO BE A DRUG COMPOSED IN WHOLE OR IN PART OF ANY KIND OF PENICILLIN, STREPTOMYCIN, CHLORTETRACYCLINE, CHLORAMPHENICOL, BACITRACIN, OR ANY OTHER ANTIBIOTIC DRUG, OR OF ANY DERIVATIVE OF THESE DRUGS AND, UNLESS THE DRUG HAS BEEN EXEMPTED BY RULES AND REGULATIONS ADOPTED UNDER THE FEDERAL ACT, IT IS NOT FROM A BATCH FOR WHICH A CURRENTLY UNEXPIRED CERTIFICATE OR RELEASE HAS BEEN ISSUED UNDER THE FEDERAL ACT;

(13) IT IS A COLOR ADDITIVE THAT IS INTENDED TO BE USED IN OR ON A DRUG FOR THE PURPOSE OF COLORING ONLY AND ITS PACKAGING OR LABELING DOES NOT CONFORM TO ANY REQUIREMENT ADOPTED UNDER § 4-239 OF THIS SUBTITLE; OR

(14) IT IS A PRESCRIPTION DRUG THAT IS DISTRIBUTED OR OFFERED FOR SALE IN THIS STATE, AND THE MANUFACTURER, PACKER, OR DISTRIBUTOR OF THE DRUG DOES NOT INCLUDE IN ANY ADVERTISEMENT, OR IN ANY OTHER DESCRIPTIVE PRINTED MATTER THAT IT ISSUES OR CAUSES TO BE ISSUED REGARDING THE DRUG, A TRUE STATEMENT OF:

(I) THE ESTABLISHED NAME OF THE DRUG, WHICH NAME IS PRINTED PROMINENTLY AND IN TYPE AT LEAST HALF AS LARGE AS THAT USED FOR ANY PRINTED TRADE OR BRAND NAME OF THE DRUG;