ASSESSMENT OF RISK OF CONVENTIONALLY MANUFACTURED HD PRODUCTS

DRUG NAME: _____

TYPE OF ____ TABLE 1 ____ TABLE 2 ____ TABLE 3

HAZARDOUS DRUGS: ANTINEOPLASTICS NON-ANTINEOPLASTICS

NON-ANTINEOPLASTICS ADVERSE REPRODUCTION

RISK OF EXPOSURE

MANUFACTURER'S PACKAGING: ____ YES ____ NO ____ N/A

FORM: ____ TABLET ____ CAPSULE ____ LIQUID ____ OTHER

MANIPULATION REQUIRED? ____ NO ____ YES
IF YES, MUST BE STORED IN A C-PEC

CONTAINMENT STRATEGIES AND/OR WORK PRACTICES COMPLY WITH:

HD'S KEPT IN A SEGREGATED & MARKED AREA WITHIN THE PHARMACY AWAY FROM PHARMACY STOCK

HD'S COUNTED WITH HD DESIGNATED COUNTING EQUIPMENT (TRAYS, SPATULA, ETC.)

HD'S NEVER USED WITH A MECHANICAL PILL COUNTER

IF MANIPULATION IS REQUIRED OR IF ALL 3 OF THE CONTAINMENT STRATEGIES ARE **NOT MET**, THIS HD DOES NOT MEET THE DEFINITION FOR A **CONVENTIONALLY MANUFACTURED HD PRODUCT** AND THE HD MUST BE KEPT IN A **C-PEC**

ASSESSMENT PERFORMED BY: _____ DATE: _____

ASSESSMENT PERFORMED BY: _____ RENEWAL DATE: _____

ASSESSMENT PERFORMED BY: _____ RENEWAL DATE: _____

ASSESSMENT PERFORMED BY: _____ RENEWAL DATE: _____

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