IREDELL HEALTH SYSTEM

Labeling, Packaging, Dispensing and Transporting HAZARDOUS DRUGS				
Approved by:	Last Revised/Reviewed Date:			
Laura Rollings, PharmD, BCPS, BCGP	04/2021			
Randi Raynor, PharmD, MBA, BCPS				
P&T Committee	Date: 04/2020			

Purpose

Hazardous Drugs (HDs) may pose serious health risks to employees that handle them regularly. The following policy details appropriate measures for labeling, packaging, dispensing and transporting of HDs.

Policy

HDs identified as requiring special handling precautions shall be clearly labeled at all times.

This entity shall use packaging containers and materials that will maintain physical integrity, stability, and sterility of the HDs during transport. Packaging materials shall protect the HD from damage, leakage, contamination and degradation, while protecting healthcare workers who transport HDs. Dedicated equipment for packaging shall be made available for HDs. Automated packaging machines shall be used only as specified in *Assessment of Risk* (AOR).

HDs must be transported in containers that minimize the risk of breakage or leakage. Individuals responsible for transporting antineoplastic injectable agents and all other HDs, as specified by AOR, shall be aware of what is being transported and have received training concerning handling spills and exposure events. Designated transport containers shall be utilized as specified in this policy and shall not contain products for more than one patient at a time.

Procedure

Labeling

Identified HDs shall display the labels, as specified in the chart below, upon completed preparation by pharmacy.

• Packaging

Personnel packaging unit-dose undamaged oral antineoplastic agents and all other HDs, as specified in AOR, shall don chemotherapy gloves. Personnel packaging unit-dose damaged (i.e. containers with broken tablets or extraneous powders) agents shall don chemotherapy gloves and an impervious gown. Eye and respiratory protection shall be applied if risk for spills is present.

Manual packaging method shall take place in the designated area, with designated counting trays and spatulas.

• Dispensing

Finished parenteral antineoplastic products shall be stored in the designated transport container in the main pharmacy while waiting for final dispensing. All other finished HDs shall be stored as described in the AOR while awaiting final dispensing.

Transporting

Personnel transporting antineoplastic injectable agents and all other HDs, as specified in AOR, shall don single pair of chemotherapy gloves. Double-bagged items as specified in AOR shall be transported while wearing a single pair of chemotherapy gloves.

	Labeling	Packaging	Dispensing	Transporting
NIOSH Group 1 injectable HDs	CAUTION CANCER CHEMOTHERAPY DISPOSE OF PROPERLY	Double sealable bagged (Procedure described in Compounding HAZARDOUS DRUGS policy) PPE requirement	Designated transport container, within a double sealable bag	Performed by trained personnel Chemotherapy gloves required
NIOSH Group 1 oral HDs	ANTINEOPLASTIC HAZARDOUS DRUG OBSERVE SPECIAL HANDLING, ADMINISTRATION AND DISPOSAL REQUIREMENTS	Manual packagaing method PPE requirement	Plastic sealable bag sized to approriate dosage form	Specifications in AOR
NIOSH Group 2 & 3 HDs (refer to individual AOR documentation)	NON-ANTINEOPLASTIC HAZARDOUS DRUG OBSERVE SPECIAL HANDLING, ADMINISTRATION AND DISPOSAL REQUIREMENTS	Manual Packaging Method OR Automated Packaging Machine - Specified in AOR PPE requirement as specified in AOR	Specifications in AOR	Specifications in AOR

INITIAL EFFECTIVE DATE: 05/2020 DATES REVISIONS EFFECTIVE: DATES REVIEWED (no changes): 04/2021