

MARYLAND Department of Health

Wes Moore, Governor ● Aruna Miller, Lt. Governor ● Laura Herrera Scott, M.D., M.P.H., Secretary

MARYLAND BOARD OF PHARMACY

4201 Patterson Avenue, Baltimore, Maryland 21215-2299 Neil Leikach, Board President ● Deena Speights-Napata, Executive Director

COMPREHENSIVE CARE PHARMACY INSPECTION FORM

1. PERMITS AND LICENSES Corporate Pharmacy Name Pharmacy Name-Doing Business as (d/b/a) or Trade Name Street Address **Business Telephone Number Business Fax Number** Inspection Date: Arrival Time: Departure Time: Type of Inspection: OAnnual OFollow-up Previous Date: Name of Inspector: No The pharmacy department provides service 24 hours. COMAR 10.34.05. The pharmacy hours of operation and after hour procedures are provided to the Comprehensive Care Facility. COMAR 10.34.05.03B Sat: _ Pharmacy Hours M-F: Sun:_ All permits, licenses, and registrations are posted conspicuously. HO §12-311, HO §12-408(b) Yes No and HO §12-6B-08 Maryland Pharmacy Permit Number _____ Expiration _____ CDS Registration Number _____ Expiration _____ DEA Registration Number Expiration The pharmacy performs sterile compounding. (If yes, complete Sterile Compounding Inspection Form) COMAR 10.34.19 The pharmacy wholesale distributes to another pharmacy. COMAR 10.34.37 The pharmacy wholesale distributes to a wholesale distributor. COMAR 10.34.37 No N/A The wholesale distribution business exceeds 5% of the pharmacy annual sales. COMAR 10.34.37 **Comments:**

2. PERSONNEL

Name of Pharmacist/Manager who is charged with ensuring compliance with all applicable laws

Pharmacist Employees	License #	Exp Date
Registered Technicians	Registration #	Exp Date
Unlicensed Personnel (non-registered)	Title	Duties

3. PERSONNEL TRAINING

All personnel h	ave received training in: (check all that apply) COMAR 10.34.21.03
Yes	No N/A Maintaining records
Yes	No N/A Patient confidentiality
Yes	No N/A Sanitation, hygiene, infection control
Yes	No N/A Biohazard precautions
Yes	No N/A Patient safety and medication errors COMAR 10.34.26.03
Yes No	There are written policies and procedures to specify duties that may be performed by unlicensed personnel under the supervision of a licensed pharmacist. COMAR 10.34.21.03
Yes No	N/A All unlicensed personnel who perform tasks in the pharmacy receive documented training for the tasks they perform. COMAR 10.34.21.03
Yes No	There is a written ongoing quality assurance program that documents the competency and accuracy of all assigned tasks. COMAR 10.34.21.03
Comments:	
4. POLICIES	S AND PROCEDURES COMAR 10.34.23.03
Yes No	Personnel access to the pharmacy COMAR 10.34.23.03C
Yes No	Scope and method of pharmacy service COMAR 10.34.23.03B
Yes No	Labeling requirements and distribution methods for medications provided in a single container COMAR 10.34.23.03E (1)
Yes No	Procedures for interim boxes. COMAR 10.34.23.03E (2)
Yes No	Documentation of policy and procedure manual provided to personnel of the pharmacy and comprehensive care facility. COMAR 10.34.23.03F
Yes No	Reporting adverse drug reactions. COMAR 10.34.23.09K
Yes No	Written policies related to re-use of returned medications.
Yes No	Documented contingency plans for continuing operations in an emergency and for disaster recovery of required records.
Comments:	

5. SECURITY COMAR 10.34.05 The pharmacy is designed to prevent unauthorized entry when the prescription area is closed during any period that the rest of the establishment is open. (Briefly describe how access is restricted.) COMAR 10.34.05.02A (5) The pharmacy and/or pharmacy department has a security system. COMAR 10.34.05.02A(2) The permit holder prevents individuals from being in the prescription area when a pharmacist is not immediately available on the premises to provide pharmacy service. COMAR 10.34.23.05C Comments: 6. PHYSICAL REQUIREMENTS AND EQUIPMENT COMAR 10.34.23.05 Pharmacy area is clean, neat, and organized. HO §12-403(b)(11)(ii)2 and No COMAR 10.34.10.01A(3). The pharmacy provides a compounding service (non-sterile procedures). If yes, the pharmacy maintains equipment that enables it to prepare and Yes dispense prescriptions properly within its scope of practice. COMAR 10.34.07.02 The pharmacy has a Class A prescription balance and weights, or a prescription balance with equivalent or superior sensitivity. COMAR 10.34.07.01-1A No The pharmacy has hot and cold running water. The medication refrigerator(s) contain only prescription items. COMAR No 10.34.07.01-1B The medication refrigerator(s) have a thermometer. COMAR 10.34.07.01-1B The current temperature of the medication refrigerator(s) is between (36F-46F). No **USP Temperature** The current temperature of the pharmacy department is between [59 to 86 No degrees F]. COMAR 10.34.05.02A (1) (a) **Temperature** If the pharmacy stocks medications requiring freezing, the freezer is maintained at No temperatures required by the medication stored within it. **Temperature** The pharmacy maintains a library of current reference sources consistent with No scope of practice that is accessible to all appropriate personnel. COMAR

10.34.07.03

Yes	No	The	pharmacy 10.34.0	has online resources. HO §12-403(b)(15) and COMAR 7.03
Yes	No	The 1		possesses the current edition of <i>The Maryland Pharmacy Laws</i> gulations. HO §12-403(b) (10) (ii)
Yes	No	Medi	the ma	d supplies within the pharmacy are properly stored according to anufacturer's specifications and State and federal laws and ons with respect to:
	Yes	No		Sanitation
	Yes	No T	1	Temperature
	Yes	No 🔚	1	Light
	Yes	No 🔚	1	Ventilation
	Yes	No		Segregation
	Yes	No]	Security
Comm	onts.			
	ents			
5 DDF	a CDID	TION	LADELI	NG FH FG AND GEODAGE
				NG, FILES, AND STORAGE
Yes	No	N/A	403(b)	py prescription files are maintained chronologically for 5 years. HO §12-
		The f	` ,	label requirements are met if a drug is dispensed pursuant to a
		THE		otion. COMAR 10.34.23.08:
	Yes 🗌	No	N/A	The name and address of the pharmacy;
	Yes 🔲	No	N/A	The serial number of the prescription;
	Yes	No	N/A	The date the prescription was dispensed;
	Yes 🗍	No E	N/A	The name of the prescriber;
	Yes	No	N/A	The name of the patient;
	Yes	No	N/A	The name and strength of the drug or devices;
	Yes	No	N/A	The quantity of the drug or device;
	Yes	No	N/A	The required precautionary information regarding controlled substances;
	Yes	No	N/A	The required cautionary statements or auxiliary labels;
	Yes	No	N/A	The name of generic manufacturer;
	Yes	No	N/A	The expiration date is indicated;
	Yes	No	N/A	(Medications in Parenteral Admixtures) The name and amount of drug(s) added;
	Ш			(
	Yes	No	N/A	(Medications in Parenteral Admixtures) The name of the pharmacist responsible
				for the admixture;
	Yes	No	N/A	(Medications in Parenteral Admixtures) The rate of infusion; and (Medications in Parenteral Admixtures) The frequency of infusion
Yes	No 🔲	N/A	Medica	ation provided per dosing period in a single container, slot, blister package,
				any other method of delivering an entire single dosing unit, or as part of
	🗀		, c	a multi-dose dispensing package, are labeled with at least the following:
	Yes	No	N/A	(1) Drug name;
	Yes	No	N/A	(2) Drug strength;
	Yes	No	N/A	(3) Name of manufacturer;
	Yes	No	N/A	(4) Name of the patient;

Yes $\bigcap_{N_0} N/A$ (5) Lot number; and
Yes N_0 $N'A$ (6) Expiration date.
Yes No N/A The pharmacist and technician initials are on prescriptions or patient dr profiles or computerized patient records. COMAR 10.34.08.01
Comments:
8. QUALITY ASSURANCE – PATIENT SAFETY / MEDICATION ERRORS
Yes No There are written procedures to follow when reporting a suspected medication error to the permit holder, pharmacist, health care facility, or other health care provider. COMAR 10.34.26.04
Yes No The pharmacy maintains a minimum of two (2) continuous years of records clearly demonstrating the content of annual educational training provided to each member of the pharmacy staff involved in the medication delivery system regarding the role and responsibility of pharmacy staff in preventing medication errors. COMAR 10.34.26.03B
Comments:
9. CONFIDENTIALITY Yes No Confidentiality is maintained in the creation, storage, access, disposal and disclosure of patient records. HO §12-403(b)(13), COMAR 10.34.10.03A and HIPAA Regulations.
Yes No Any identifiable information contained in a patient's record is not disclosed unless authorized by the patient, or an order of the court, or as authorized pursuant to HG §4-301 through §4-307.COMAR 10.34.10.03B
Comments:
10. DRUG CONTROL AND ACCOUNTABILITY COMAR 10.34.23.09
Yes No N/A The pharmacy maintains invoices as required by law for accurate control an accountability of all pharmaceuticals. COMAR 10.34.24.03
Yes No N/A The pharmacy has written policies and procedures for the safe handling of dru recalls. See www.recalls.gov
Yes No N/A The pharmacy maintains records of all recalls. See www.recalls.gov
Yes No No N/A The pharmacy has a written procedure in place for removal of all expired drug (both prescription and OTC) COMAR 10.34.12.01
Yes No No N/A The pharmacy has a process for discontinued medication and returned medication that accounts for proper storage and labeling. COMAR 10.34.23.09
Comments:

11. EMERGENCY DRUG KIT COMAR 10.34.23.09.F The emergency drug kit is secured with a tamper evident seal or electronic security system. COMAR 10.34.23.09.F.(1) The emergency drug kit meets the labeling requirements as set forth on COMAR No 10.34.23.09.F.(2). A written policy exits regarding what medications and quantities are to be contained in interim drugs box as well as procedures for restocking medications. COMAR 10.34.23.03 Comments:____ 12. CONTROLLED SUBSTANCES Hard copy prescription files are maintained chronologically for 5 years. COMAR 10.34.20.03; HO §12-403(b)(13). (CDS-Federal law requires record retention for 7 years) A perpetual inventory is maintained for Schedule II controlled substances. No (Recommended) Schedule II controlled substances are dispersed throughout the stock of non-No controlled substances, or stored in such a manner as to obstruct theft or diversion. COMAR 10.19.03.12B (2) The pharmacy has a copy of the most recent required biennial inventory of No Schedule II- V controlled substances. COMAR 10.19.03.05B Inventory date: Inventory Opening or Closing (Check one) completed at: No The inventories and records of Schedule II-V drugs are maintained and readily available, COMAR 10.19.03.05 and 21 CFR 1304.03 Records are kept of all receipts of controlled substances entered into the No pharmacy inventory (including DEA Form 222 or CSOS orders). COMAR 10.19.03.05 The prescription label for controlled drugs include the following warning: "CAUTION Federal law prohibits the transfer of this drug to any person other than the patient for whom it was prescribed", in 6-point type or the Pharmacy utilizes an auxiliary label that contains this warning, COMAR 10.19.03.08D (1) All controlled substances prescriptions comply with COMAR 10.19.03 Yes The permit holder or pharmacist designee(s) has written policies and procedures for investigating discrepancies and reporting of theft or loss. COMAR 10.19.03.12B (4) For the return of Schedule III-V drugs, the pharmacy uses a distribution system No that classifies medications as pharmacy inventory until the utilization of the medication by the patient. COMAR 10.34.23.09.C Comments:

13. AUTOMATED MEDICATION SYSTEMS Yes No (if No, go to #14)
Yes No N/A The facility uses an automated device(s) as defined in COMAR 10.34.28.02.
Written policies and procedures exist for (check all that apply): COMAR
10.34.28.05
Yes $N_0 N_{N/A}$ Control of access to the device.
Yes No N/A Accounting for medication added and removed from the system.
Yes No N/A Sufficient safeguards are in place to ensure accurate replenishment of the
automated medication system. If yes, describe safeguards. COMAR 10.34.28.06
Comments:
Yes No N/A Adequate records are maintained for at least two years addressing the following (check all that apply). COMAR 10.34.28.11
Yes No N/A Maintenance records.
Yes No N/A System failure reports.
Yes No N/A Accuracy audits.
Yes No N/A Quality Assurance Reports.
Yes No N/A Reports on system access and changes in access.
Yes No N/A Training records.
Yes No No N/A The pharmacy has records, documents, or other evidence of a quality assurance program regarding the automated medication system in accordance with
the requirements of COMAR 10.34.28
Comments:
14. OUTSOURCING Yes No (if No, go to #13)
Yes No No N/A The facility outsources the preparation of medications or performs outsourcing functions for other pharmacies. COMAR 10.34.04.02
Yes No N/A The facility serves as a primary pharmacy outsourcer to other pharmacies. COMAR 10.34.04.02
Yes No N/A The facility serves as a secondary pharmacy. COMAR 10.34.04.02
Yes No N/A Written policies exist for maintenance of documentation regarding transfer of prescription records. COMAR 10.34.04.06
Yes No No N/A Documentation is maintained, including the names and locations of the
pharmacies, names of pharmacists, and a record of the preparations made. COMAR 10.34.04.03 and .05

Yes No N/A The permit holder employs an outside agency/business entity for the provision of any pharmacy services, inclusive of staffing, remote order entry, and management.
If yes: Name of agency, state of incorporation, service contracted, and State of Maryland License/Permit Number:
COMAR 10.34.04.06E
The pharmacist from the <u>primary</u> pharmacy documents in a readily retrievable and identifiable manner (Check all that apply): COMAR 10.34.04.06
Yes No N/A The prescription order was prepared by a secondary pharmacy. Yes No N/A The prescription order was prepared by a secondary pharmacy.
Yes No N/A The name of the secondary pharmacy. The name of the pharmacist who transmitted the prescription order to the secondary pharmacy.
Yes No N/A The name of the pharmacist at the secondary pharmacy to whom the prescription order was transmitted if the transmission occurred in an oral manner.
Yes No No N/A The date on which the prescription order was transmitted to the secondary pharmacy.
Yes No N/A The date on which the preparation was sent to the primary pharmacy.
Yes No N/A The primary and secondary pharmacies are both licensed in the State of Maryland, or operated by the federal government. COMAR 10.34.04.06F
Yes No N/A The primary pharmacy maintains, in a readily retrievable and identifiable manner, a record of preparations received from the secondary pharmacy. COMAR 10.34.04.06G
The permit holder at the <u>secondary</u> pharmacy maintains documentation in a readily retrievable and identifiable manner, which includes (check all that apply): COMAR 10.34.04.07
Yes No No N/A Records of the prescription orders transmitted from another pharmacy.
Yes No N/A The date on which the prescription order was transmitted from the primary pharmacy.
Yes No N/A The name and information identifying the specific location of the primary pharmacy.
Yes No N/A The name of the pharmacist who transmitted the prescription to the secondary pharmacy if the transmission occurred in an oral manner.
Yes No No N/A The name of the pharmacist at the secondary pharmacy who accepted the transmitted prescription order.
Yes No N/A The name of the pharmacist at the secondary pharmacy who verified/performed the final check of the prescription order.
Comments:

15. DISTRIBUTION N/A Sales of prescription drugs other than by patient specific prescription orders exceeds 5% of the pharmacy's annual sales. If yes, Maryland distributors license #_____COMAR 10.34.22.01. Comments: 16. MEDICATION ORDERS (COMAR 10.34.23.09.H) No Medications are dispensed from the pharmacy only in response to medication orders issued by authorized prescribers or by prescriber per institution approved protocols. 17. MEDICATION PACKAGING (COMAR 10.34.23.07) No No N/A The pharmacy prepares packaged medications. (If yes complete questions below) Packaged from the original manufacturer's container: Yes $N_0 \setminus N_A$ The pharmacy uses a lot number and expiration date assigned by the pharmacy instead of the distributor or manufacturer information in a master log if kept with respect to drugs that are packaged within the pharmacy facility from the original manufacturer's container which includes the: Yes No N/A Name of drug; Strength; N/A Yes No N/A Manufacturer; No Yes No N/A Lot Number assigned by the pharmacy; No N/A Lot number assigned by the distributor or manufacturer; Yes N/A Quantity packaged; No Yes Manufacturer's expiration date; Yes No N/A Lot number assigned by the distributor or manufacturer; Yes No N/A Date of packaging; Yes No N/A Name of the pharmacy technician who performed packaging functions; No N/A Name and initials of verifying licensed pharmacist. Packaged from Another Pharmacy: No N/A The licensed pharmacist packages medication received from another pharmacy licensed in Maryland or operated by the government of the United States provided that: Yes No N/A (1) The licensed pharmacist determines that the medication has been handled in a manner which preserves the strength, quality, purity, and identity of the drug or device during an interim period

between the time it was dispensed by the original pharmacy and to directly send medication to the packaging pharmacy;

Yes N	e	icensed pharmacist packages and dispenses all at one time the ntire quantity of the prescription medications received from nother pharmacy for packaging;
Yes N	$N_0 \square N/A \square$ (3) The r	manufacturer's name is present on the container received from he other pharmacy; and
Yes N	$N_0 \square N/A \square$ (4) The 1	licensed pharmacist maintains a master log that includes the following information:
Y	Yes No No N/A	(a) Name of the drug;
Y		(b) Lot number assigned by the packaging pharmacy;
Y	Yes No N/A	(c) Strength;
Y	Yes No No N/A	(d) Manufacturer;
Y	Yes No No N/A	(e) Name, address, and telephone number of the original dispensing pharmacy;
Y	Yes No No N/A	(f) Prescription number for the original dispensing pharmacy;
Y	Yes No No N/A	(g) Quantity packaged;
Y	Yes No N/A	(h) Expiration date as assigned by the original dispensing pharmacy;
Y	Yes No N/A	(i) Date of packaging;
Y	Yes No N/A	(j) Name of pharmacy technician who performed packing function;
Y	Yes No N/A	(k) Name and initials of verifying licensed pharmacist; and
Y	Yes No No N/A	(l) Name of the patient.
Comments:		
18. CONSULTA	ANT SERVICES	
Yes No	There are written policies consulting pharma	for and documentation of timely medication review by acists at all sites.
19. DELIVERY		
Yes No	There are policies for and sites.	documentation of timely delivery of medications to all
Yes No		and documentation of timely delivery of controlled notes to all sites.

nactor Signatura	
pector Signature	
armacist Name:	Date:
nt)	
eived a copy of the inspection report on _	Date and Signature of the Pharmacist

FINAL03/06/2023

CONTROLLED DANGEROUS SUBSTANCES WORKSHEET

Clear Form

Permit#:					
Pharmacist Signature:					
Rx#:	. 1.				
Date Fill	ed:				
	NDO	C Number		ON HAND	PEI
	NDC			INVENTORY	INV
					<u> </u>
					<u> </u>
COMMENTS:					
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					-
	SCHEDULE II				- -
	SCHEDULE II	AUDIT			_
		AUDIT			-
	SCHEDULE II Drug Date of last Inspection	AUDIT			
Amount at last inspection/biennia	SCHEDULE II Drug Date of last Inspection	AUDIT	(A)		_
Amount at last inspection/biennia Purchased since inspection/bienn	SCHEDULE II Drug Date of last Inspection	AUDIT	(A) (B)		_
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial	(A)		_
Amount at last inspection/biennia Purchased since inspection/bienn	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial	(A) (B)		
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial	(A) (B) (C) = A (D)	A + B	_
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial 0	(A) (B) (C) = A (D) (E) = C	A + B	
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory Quantity on Hand	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial 0	(A) (B) (C) = A (D) (E) = C (F)	A + B C - D	
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory	SCHEDULE II Drug Date of last Inspection	AUDIT n/Biennial 0	(A) (B) (C) = A (D) (E) = C (F)	A + B	ge
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory Quantity on Hand	SCHEDULE II Drug Date of last Inspection	AUDIT on/Biennial 0	(A) (B) (C) = A (D) (E) = C (F)	A + B C - D F-E) or (E-F)	ge
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory Quantity on Hand Discrepancy	SCHEDULE II Drug Date of last Inspectional al nial	AUDIT on/Biennial 0	(A) (B) (C) = A (D) (E) = C (F)	A + B C - D F-E) or (E-F)	ge
Amount at last inspection/biennia Purchased since inspection/bienn Total inventory Quantity dispensed Expected inventory Quantity on Hand	SCHEDULE II Drug Date of last Inspectional al nial	AUDIT on/Biennial 0	(A) (B) (C) = A (D) (E) = C (F)	A + B C - D F-E) or (E-F)	ge

PRESCRIPTION REVIEW

CII# DATE

COMMENTS:		
	CIII - CV # DATE	
COMMENTS:		