

Enforcement v250402

Unassigned

Name	Test Pharmacy	Case #		Permit	
Address	726 S Scales	Person		Inspection Date	04/08/2025
	Street	Providing Info		Inspection User	Brashears, Krysta
	Reidsville, NC	Person In	Jack Campbell	Inspection	DISTRICT3
	27320	Charge		District	
# RPhs		Rx Volume/Date			
# Techs		Hours			
Follow-Up CAP	No	Office	No		
CAP Requested	No	Commercial Use	No		
CAP	No	Ship to Other	No		
Documentation		States			
Received		States Shipped			
Additional	No	То			
Documents		Commercial Use			
Office Use		Documented			
Office Use		Clinical			
Comments		Indication			

Comments

None

General DME			
	Answer	Question	
1)	Unanswered	GS 90-85.23 - Permit posted conspicuously in the facility.	
Permi	ts		
2)	Unanswered	46.1608(a)(1) - Adequate qualified personnel to properly render services in a manner prescribed by law.	
3)	Unanswered	46.1608(a)(3) - If disp. Medical O2, must ensure (A) 24hr back up; (B) O2 analyzer @ facility, if concentrations are disp.	
4)	Unanswered	46.1608(a)(4) - Suitable facility for inventory, fabrication work space, and record retention.	
5)	Unanswered	46.1608(a)(5) - Copy of the pharmacy laws of NC, Pharmacy Practice Act, and current copy of the rules and regulations in the pharmacy.	
6)	Unanswered	46.1608(a)(6) - Functioning lavatory with hot/cold running water or hand washing appliances or waterless hand cleaner.	
7)	Unanswered	46.1608(a)(7) - Clean, orderly, and sanitary.	
8)	Unanswered	46.1608(a)(8) - Services applicable to its customer base.	
9)	Unanswered	46.1608(a)(11) - Essential services available 24 hours 7 days a week.	
10)	Unanswered	46.1608(a)(12) - Maintain written procedure at location for handling complaints.	
11)	Unanswered	46.1608(a)(13) - Compliance with local and state fire and building laws.	
12)	Unanswered	46.1608(a)(14) - Compliance with OSHA requirements, including Universal Precautions.	
13)	Unanswered	46.1608(b) - Permit must be countersigned by the person in charge.	
14)	Unanswered	46.1804(a) - Device dispensing and medical equipment delivery occurs only by bona fide employees.	
15)	Unanswered	46.1806 (c) - Records retained for three (3) years.	
16)	Unanswered	46.1806(i) - Transfer of orig. rx info. for refill is permissible b/t device & med equip holders as transferring permit	

Gene	eral DME	
	Answer	Question
		holder provides all records necessary for disp & does not interfere w/ the svc. & claims process of receiving permit holder.
17)	Unanswered	46.1809(4) - Record of emergency dispensing/delivery of devices/ medical equipment and prescriber notified.
18)	Unanswered	46.2504(a)&(c) - Pt. Counseling: name, description, purpose of device/ equipment, route dosage, administration, continuity of service, special directions, side effects, interactions, self monitoring techniques, proper storage, refill information, patient name, address, telephone number, DOB, gender, and medical history.
19)	Unanswered	46.2504(b) - Toll free telephone service is required when a registrant's primary population is not accessible through a local exchange. Proficiency in explaining and demonstrating the safe and proper use of devices and equipment and for documenting the demonstration of such proficiency.
20)	Unanswered	46.2504(d) - Information for counseling kept current.
21)	Unanswered	46.2504(g) - Records retained for three (3) years
22)	Unanswered	46.2504(h) - Documentation that written notice of warranty was given to the patient concerning service after the sale.
Devic		
23)	Unanswered	46.2602 - Devices shall be dispensed to outpatients only pursuant to an order from a practitioner.
24)	Unanswered	46.2603 - Educated and trained sufficiently in safe and proper delivery of medical equipment.
25)	Unanswered	46.2604(a) - Device and medical equipment order must comply with Board Rules 2301-2305 and shall be maintained at the dispensing site. Serial numbers shall be maintained as part of the records.
26)	Unanswered	46.2604(b) - Records retained for three (3) years.
27)	Unanswered	46.2604(c) - File copy of every item sold or rented with serial number or tracking number or code in compliance with FDA Medical Device Tracking requirements.
28)	Unanswered	46.2606 - Convey warnings issued by government agencies and manufacturers.
Rehab	oilitation Equipme	nt
29)	Unanswered	46.2609(b)(1-3) - Solicitation of results of assessments and evaluation. Choice of commercial vs. custom equipment. Measurements of Patients and Clients. Documenting goals and objectives.
30)	Unanswered	46.2609(b)(5) - Instruct patient and family in safe and proper use and care of equipment.
31)	Unanswered	46.2609(b)(6) - Provide service and support and within 72 hours provide a response to a patient request.
32)	Unanswered	46.2609(b)(7) -Documentation that written notice of warranty was given to the patient concerning service after the sale. Provide specific written statement of warranty on the equipment provided, including commercial warranties and those for adapted or custom fabricated items.
33)	Unanswered	46.2609(b)(8) - Liability insurance of at least 1,000,000.
34)	Unanswered	46.2609(b)(9)(A) - Compatibility and safety of interfacing techniques of custom fabricated parts with commercially available equipment.
35)	Unanswered	46.2609(b)(9)(B) - Understanding the properties of the materials being used in custom designed and modified equipment to assure long term durability.
36)	Unanswered	46.2609(b)(9)(C) - Documenting goals and objectives of the referring medical or education personnel, as well as short and long term effectiveness of the equipment in meeting those goals and objectives.
37)	Unanswered	46.2609(b)(9)(D) - Complaints and problems, including complaint file.
Medic	al Gas/Oxygen/Re	espiratory Equipment
38)	Unanswered	46.2610(a)(2) - Comply with all DOT regulations if transporting medical gases.
39)	Unanswered	46.2610(a)(3) - Comply with FDA and all state agency requirements regarding transfilling and repackaging if transfilling.
40)	Unanswered	46.2610(a)(4) - Demonstrate that O2 provided in cylinder or liquid form meets minimal purity standards for medical grade O2.

Gene	ral DME	
	Answer	Question
41)	Unanswered	46.2610(a)(6)(A) - Demonstrate that each piece of equipment has been checked, is free of defect, and operates within manufacturer's specifications.
42)	Unanswered	46.2610(a)(6)(B) - Refrain from modifying equipment to the extent that the modification might reasonably cause harm.
43)	Unanswered	46.2610(a)(6)(C) - Maintain all electrical components so they do not present a fire or shock hazard.
44)	Unanswered	46.2610(a)(6)(D) - Ensure that all appropriate warning labels or labeling, including tags are present on the equipment provided.
45)	Unanswered	46.2610(b)(1) - Ensure that lot numbers and expiration dates are affixed to each cylinder delivered.
46)	Unanswered	46.2610(b)(2) - Maintain a tracking system for all medical oxygen and gas delivered.
47)	Unanswered	46.2610(b)(3) - Document all equipment serial numbers and model numbers to ensure that equipment can be retrieved if a recall is initiated.
48)	Unanswered	46.2610(b)(4) - Maintain records for equipment that requires FDA tracking.
49)	Unanswered	46.2610(c)(1) - Function and safety check prior to set up.
50)	Unanswered	46.2610(c)(2) - Protocol for cleaning and disinfecting equipment which addresses both aerobic and anaerobic pathogens.
51)	Unanswered	46.2610(c)(3) - Maintain a Material Safety Data Sheet on file for solutions and products used in cleaning and disinfecting procedures.
52)	Unanswered	46.2610(c)(4) - Maintain segregated areas for clean, dirty ,and contaminated items.
53)	Unanswered	46.2610(c)(5) - Clean and disinfect equipment according to manufacturers' specification.
54)	Unanswered	46.2610(c)(6) - Instruct the patient on proper cleaning techniques as specified by the manufacturer.
55)	Unanswered	46.2610(d)(1) - Problem reporting, tracking, recall, and resolution.
56)	Unanswered	46.2610(d)(2) - Performance of service as specified by the manufacturer and the documentation of such performance in the service records.
57)	Unanswered	46.2610(d)(3) - Routine inspection, service, and maintenance of equipment located in the patient's home according to manufacturer's specifications.
58)	Unanswered	46.2610(e)(1-7) - Equipment suppliers shall maintain repair logs to document repairs and maintenance of equipment. Documentation must include type of equipment, manufacturer, model, serial number, date of repair, type of repair, who performed the repair.
59)	Unanswered	46.2610(f) - Maintain testing equipment to ensure accurate calibration.
60)	Unanswered	46.2610(g) - Written policy for handling complaints and problems, which include complaint file.
61)	Unanswered	46.2610(h)(1)(A-E) - Counseling requirements, Orientation checklist, instructions, safety, precautions, cleaning procedures, maintenance procedures, and return demonstrations on back up O2 systems. Plan of Service.
62)	Unanswered	46.2610(h)(2) - Instruct patients about routine & emergency contact procedures.
63)	Unanswered	46.2610(h)(3) - Review written instructions to properly operate equipment.
Medica	al Equipment	
64)	Unanswered	46.2611(a)(1-3) - Solicitation of results of assessments and evaluation. Choice of commercial vs. custom equipment. Measurements of Patients and Clients. Documenting goals and objectives.
65)	Unanswered	46.2611(a)(4) - Deliver, fit, and adjust the prescribed equipment.
66)	Unanswered	46.2611(a)(5) - Instruct patient and family in safe and proper use and care of equipment.
67)	Unanswered	46.2611(a)(6) - Provide service and support and within 72 hours provide a response to a patient request.
68)	Unanswered	46.2611(a)(7) - Liability insurance of at least 1,000,000.
69)	Unanswered	46.2611(a)(8) - Demonstrate that each piece of equipment has been checked, is free of defect, and operates with in the manufacturers' specifications.

General	DME

	Answer	Question
70)	Unanswered	46.2611(a)(9) - Refrain from modifying equipment to the extent that the modification might reasonably cause harm.
71)	Unanswered	46.2611(a)(10) - Maintain all electrical components so they do not present a fire or shock hazard.
72)	Unanswered	46.2611(a)(11) - Ensure that all appropriate warning labels or labeling, including tags are present on the equipment provided.
73)	Unanswered	46.2611(a)(12) - Function and safety check prior to set up.
74)	Unanswered	46.2611(a)(13) - Protocol for cleaning and disinfecting equipment which addresses both aerobic and anaerobic pathogens including procedures to prevent cross contamination.
75)	Unanswered	46.2611(a)(14) - Clean and disinfect equipment according to manufacturers' specification.
76)	Unanswered	46.2611(b)(1) - Problem reporting, tracking, recall, and resolution.
77)	Unanswered	46.2611(b)(2) - Performance of service as specified by the manufacturer and the documentation of such performance in the service records.
78)	Unanswered	46.2611(b)(3)(A-G) - Equipment suppliers shall maintain repairs logs to document repairs and maintenance of equipment. Documentation must include type of equipment, manufacturer, model, serial number, date of repair, type of repair, who performed the repair.
79)	Unanswered	46.2611(c)(1) (A-E) - Counseling requirements, orientation checklist, instructions, safety, precautions, cleaning procedures, maintenance procedures, and return demonstrations on equipment delivered.
80)	Unanswered	46.2611(c)(2) - Instruct patient about emergency & routine contact procedure.
81)	Unanswered	46.2611(c)(3) - Written instructions to properly operate equipment.