

LOUISIANA BOARD OF WHOLESALE DRUG DISTRIBUTORS

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INSPECTION GUIDELINES

The following areas will be reviewed when performing the inspection of the wholesaler:

1. Responsibility for Operation

- Check for posting of license
 - Names of officers
 - Ownership information
 - Administration Who is responsible for operations?

2. Policies and Procedures

- Review written policies and procedures these should cover the following (minimums):
 - Receipt of product
 - Inventory and distribution
 - o Inspection of incoming and outgoing shipments of legend drug, device, and/or medical gas products
 - Identification, recording, and correction of inventory variances and adjustments for legend drug, device, and/or medical gas products
 - Assure the legend drug, device, and/or medical gas product is shipped only to authorized recipients (e.g., pharmacies, hospitals, physicians, dentists, veterinarians, other wholesale distributors)
 - Check for valid licensure
 - o Product stock rotation
 - o Excess or unusual purchases of legend drug, device, and/or medical gas product
 - o Security
 - Security system
 - o Storage
 - o Temperature monitoring and logs (when required)
 - Facility sanitation and pest control
 - o Handling of recalls and withdrawals of legend drug, device, and/or medical gas products
 - Handling of outdated, damaged, deteriorated, misbranded or adulterated drug, device, and/or medical gas product
 - Product segregation
 - o Handling of returned legend drug, device, and/or medical gas product
 - Product destruction
 - o Reporting losses, theft, contraband, misbranded, or counterfeit product
 - Crisis handling
 - o Recordkeeping
 - o Personnel qualification/training and responsibility

3. Suspicious and Excessive Purchases and Sales (Sales personnel should provide input)

- What criteria is used?
- To whom do they report?

4. Record Keeping

- Records readily available
- Inventories and transactions current
- Sales and receipts records
- Temperature monitoring logs current (not required of licenses who distribute medical gas product only)
- Customer licenses
 - Customer list must include customer name, address, authorizing license, permit, certificate number and expiration date
 - Copies of valid authorizing license, permit, certificate for each customer must be on file
 - Cross-check records select some records for verification of customer license

- Suppliers/vendors are verified as current, valid LBWDD licensees
- Check on excess returned legend drug, device, and/or medical gas product
- Maintenance of records for a minimum of three years.

5. Recalls

- Where are recalled legend drugs, devices, and/or medical gases stored?
- Are they in a signed and separated (quarantined) location?

6. Returned Goods

- How are returned products from the customer handled?
- Where are returned items stored?
- How are outdated products checked?
- How is it determined which returned items may be resold?

7. Facilities

- Is the facility suitable for wholesale/distribution operations
 - Suitable size and construction
 - o Clean and orderly
 - Free of infestation from insects, rodents, birds, etc.

8. Storage Area

- Adequate lighting
- Ventilation
- Product stored at appropriate temperature and humidity conditions in accordance with USP or labeled requirements
- Sanitary
- Spacious
- Properly equipped
- Secure
- Has a designated and marked quarantine area
- Product stored away from poisons or harmful substances such as cleaning solutions, vermin repellant, and chemicals.

9. Security

- Working monitored alarm system; or for licensees who distribute medical gas product only, product is stored under lock and key in Board approved facility and/or working monitored alarm system on open dock area
- Working security system against theft or diversion
 - How does security system operate?
 - Are legend drugs, devices, and/or medical gases stored in secure area? Is the shipping / receiving area secure?
 - o Who has access to legend drugs, devices, and/or medical gases?
 - o Is access to legend drugs, devices, and/or medical gases restricted?
 - Who has access to facility after operating hours?
 - Has there ever been any lost or stolen legend drug, device, and/or medical gas product at the facility.

ARE YOU READY FOR INSPECTION? TEST INSPECTION - CHECKLIST

□ Current LBWDD license displayed [46:XCI.301.E] (not for initial (new) applications) \Box Has off-site storage facility(s) [46:XCI.301.J] □ Not Applicable □ Facility is of suitable size and construction [46:XCI.309.A.1.a] □ Facility is clean and orderly [46:XCI.309.1.d] □ Facility is free from infestation by insects, rodents, birds, etc. [46:XCI.309.1.e] □ Storage area provides adequate: [46:XCI.309.1.b] □ Lighting □ Humidity □ Ventilation □ Space □ Temperature □ Equipment □ Security Sanitation □ Storage area has a designated and clearly marked guarantine area [46:XCI.309.1.c] □ Storage facility has working monitored alarm system <u>OR</u> (medical gas only) is a Board-approved facility kept under lock and key [46:XCI.309.2.b/c] □ Medical gases stored on open dock area has a working monitored alarm system [46:XCI.309.2.d] □ Facility is equipped with working security system suitable for protection against theft or diversion of product and tampering with computer or electronic records [46:XCI.309.2.e] Drug or Device products are stored at appropriate temperature [46:XCI.309.3] □ Inventory records readily available for inspection [46:XCI.311.B/C] □ Perpetual inventories and records of all transactions for all drugs, devices, and/or medical gases are current [46:XCI.311.A] □ Temperature monitoring logs are maintained and current (excluding medical gases only) [46:XCI.309.A.3.b] Copies of customer licenses verifying authority to purchase drug products on file and current [46:XCI.311.D] □ Has verified all suppliers are licensed by LBWDD [46:XCI.311.F] □ Has established and maintains policies and procedures covering: [46:XCI.313] □ Product returns or destruction [45:XCI.309.A.5] □ Receipt of drug, devices, and/or medical gases □ Reporting of loss and theft □ Storage □ Security and crisis handling □ Validation of customer licenses [46:XCI.311.D] □ Inventory and distribution □ Correction of inventory errors and □ Notification of theft & diversions; findings of inaccuracy contraband, counterfeit or misbranded drugs □ Inspection of all incoming & outgoing □ Verification of suppliers [46:XCI.311.F] □ Review of excessive or suspicious purchases shipments [45:XCI.309.A.4] □ Product rotation □ Monitoring & recording of storage temperatures □ Recalls and withdrawals [46:XCI.309.A.3] □ Out-dated product segregation □ Has current list of responsible persons – owners, officers, directors, and responsible party [46:XCI.315] □ List of employees with access to product List of employees with access to building after hours □ Facilities handling controlled substances = [46:XCI.317.2] □ Has current CDS registration In Not Applicable □ Has current DEA registration D Not Applicable