SUBCHAPTER 09M - DRUGS

02 NCAC 09M .0101 MANUFACTURER REGISTRATION

(a) Every person doing business in North Carolina and operating as a prescription drug manufacturer, repackager or wholesaler shall submit a completed prescription drug registration form to the department. A separate registration form shall be submitted for each establishment operating in the State of North Carolina. Each registration form shall be signed by the owner or individual in charge.
(b) A fee of five hundred dollars ($500.00) for manufacturers or repackagers and a fee of three hundred fifty dollars ($350.00) for wholesalers shall be submitted with each registration or renewal form.
(c) On or before December 31 of each year, every person registered in accordance with Paragraph (a) of this Rule shall submit a renewal form furnished by the division.
(d) Prescription Drug Registration Forms may be obtained from the Food and Drug Protection Division.

History Note: *Authority G.S. 106-140.1;*
Eff. June 1, 1988;
Amended Eff. January 1, 1992;
Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. March 22, 2015.

02 NCAC 09M .0102 ADOPTION BY REFERENCE

History Note: *Authority G.S. 106-145.12; 106-145.10;*
Eff. July 1, 2010;
Pursuant to G.S. 150B-21.3A, rule Expired April 1, 2015.

02 NCAC 09M .0103 DUTY TO VERIFY SUPPLIERS

Wholesale prescription drug distributors that have distribution facilities in North Carolina shall not purchase or accept delivery of a prescription drug from suppliers that are not licensed or registered to ship or sell in or into North Carolina.

History Note: *Authority G.S. 106-145.12; 106-145.1;*
Eff. July 1, 2010;
Pursuant to G.S. 150B-21.3A, rule is necessary without substantive public interest Eff. March 22, 2015.