

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.