



North Carolina State Health Director’s Standing Order for Oral and Transdermal Self-Administered Combined Hormonal and Progestin-Only Contraceptives

Revised April 4, 2023

Pursuant to S.L. 2021-110, this standing order signed by the North Carolina State Health Director, authorizes immunizing pharmacists practicing in the state of North Carolina and licensed by the North Carolina Board of Pharmacy to dispense, deliver, or administer the following contraception products as directed below.

Immunizing pharmacists who provide contraception products in accordance with this standing order must also complete North Carolina Hormonal Contraception Training Program

Contraception Dispensing Protocol			
Eligible Candidates	<ul style="list-style-type: none"> ▪ Persons of reproductive age, who voluntarily request contraception, and are at risk of experiencing unintended pregnancy and that the patient is, within reasonable certainty, not pregnant. ▪ This standing order may be used for persons < 18 years of age with a parent or legal guardian consent. ▪ Persons of reproductive age may be provided any contraceptive allowed by this standing order that is a US Medical Eligibility Criteria (USMEC) category 1 or 2 agent based on completion of a patient assessment and evaluation consistent with the USMEC linked below and/or the Oral and Transdermal Self-Administered Combined Hormonal and Progestin-Only Contraceptives Patient Questionnaire and Pharmacist-Initiated Hormonal Contraception Assessment and Treatment Care Pathway for this standing order. An alternative questionnaire, assessment and evaluation may be completed, in a format of the immunizing pharmacists’ choosing, as long as it is consistent with USMEC. A patient questionnaire document may be completed by the patient prior to, or at the time of, the visit and then reviewed with the patient by the pharmacist. ▪ Patient has a seated blood pressure (< 140/90 mmHg) measured by a qualified health care provider at the time of assessment. This may be done manually or by a blood pressure machine. If the initial blood pressure reading is 140/90 mmHg or greater, reassess the blood pressure after the patient has been seated for five or more minutes. If blood pressure remains high, then do not dispense, deliver or administer and refer to a medical care provider. ▪ Refer to the following guidance regarding eligibility criteria and when a person should start using specific contraceptive methods: <ul style="list-style-type: none"> ▪ CDC When to Start Using Specific Contraception ▪ Summary Chart of U.S. Medical Eligibility Criteria for Contraceptive Use ▪ U.S. Medical Eligibility Criteria for Contraceptive Use, 2016 		
Combined Hormonal Contraceptive (CHCs)			
Route(s) of Administration	Combined Oral Contraceptive (COC)	Transdermal (TD)	Progestin Only Pill (POP)
Medication	<ul style="list-style-type: none"> • estradiol valerate/dienogest • estetrol/drospirenone • ethinyl estradiol/desogestrel • ethinyl estradiol/drospirenone • ethinyl estradiol/drospirenone/levomefolate • ethinyl estradiol/ethynodiol diacetate • ethinyl estradiol/levonorgestrel • ethinyl estradiol/norethindrone • ethinyl estradiol/norgestimate • ethinyl estradiol/norgestrel • mestranol/norethindrone 	<ul style="list-style-type: none"> • ethinyl estradiol/levonorgestrel • ethinyl estradiol/norelgestromin 	<ul style="list-style-type: none"> • drospirenone • norethindrone
Directions for Use	Take one tablet by mouth daily.	Apply one patch to the skin once a week x 3 weeks. Then remain patch-free for one week.	Take one tablet by mouth daily.
Follow guidance for initiation, modification, and discontinuation as set out in the Pharmacist Initiated Hormonal Contraception Assessment and Treatment Care Pathway (Appendix B).			
Refills	As needed up to a one-year supply. Refills may be provided in monthly or extended supplies, as allowed by the patient’s insurance. Patient screening questionnaire must be completed at least annually.		

