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Two-week postpartum intrauterine contraception insertion: A study of feasibility, patient acceptability, and short-term outcomes

Matthew L. Zerden, Gretchen S. Stuart, Samantha Charm, Amy Bryant, Joanne Garrett, Jessica Morse

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Title: Two-week postpartum intrauterine contraception insertion: A study of feasibility, patient acceptability, and short-term outcomes

Authors: Matthew L. Zerden^{a, b}; Gretchen S. Stuart^a; Samantha Charm^a; Amy Bryant^a; Joanne Garrett^{a, c}; Jessica Morse^a;

^a Division of Family Planning in the Department of Obstetrics & Gynecology, University of North Carolina School of Medicine. 4002 Old Clinic Building, Chapel Hill, NC, 27599-7570

^b WakeMed Health & Hospitals, 10,000 Falls of Neuse Rd. Raleigh, NC, 27614

^c Department of Epidemiology in the Gillings School of Public Health, University of North Carolina, 135 Dauer Dr., Chapel Hill, NC, 27599

Corresponding Author, present address: Matthew Zerden 4002 Old Clinic Bldg, CB #7570, Chapel Hill, NC 27599-7570 mzerden@gmail.com

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Clinical Trial Registration: ClinicalTrials.gov, www.clinicaltrials.gov, NCT02121067

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Objective: To determine the feasibility and acceptability of inserting the Levonorgestrel Intrauterine System, LNG 52 mg IUS (LNG IUS), at two weeks postpartum.

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