

Pharmacist experiences delivering contraceptive care on the PRISM platform in a community pharmacy setting

Thank you for considering participating in this important study. This quick reference guide will provide you with the information you need to decide if joining the study is right for you.

Scope of the Study

This study is a 12-week pilot program conducted at select pharmacies across the United States. Participating pharmacists will use the PRISM software platform to deliver contraceptive services directly in a community pharmacy setting. The platform is intended to streamline workflows and provide clinical decision support to facilitate pharmacist-led care. Your participation will not only give patients in your community convenient access to contraceptive services but will also generate valuable real-world data on the feasibility of implementing these services using the PRISM platform. Furthermore, the insights gained from this study may help fill a critical gap in the literature regarding pharmacist-led contraceptive care, potentially informing strategies to broaden adoption nationwide, which could enhance patient access and support future efforts to expand pharmacists' scope of practice.

What's Involved

- **Your Role:**
 - Provide patient-centered contraceptive consultations in the pharmacy using the PRISM platform.
 - Ensure prescriptions generated during encounters are compliant with the CDC's United States Medical Eligibility Criteria (USMEC) guidelines.
 - Provide contraceptive education and counseling for each patient.
 - Provide non-emergent contraceptive post-encounter guidance for patients that have received services at the pharmacy on an as-needed basis.
 - Complete 5 surveys to contribute to pharmacy-led services implementation science and provide feedback on the PRISM platform.

- **Time Commitment:**
 - Complete the study enrollment process and create a PRISM account (about 30 minutes).
 - Complete a pharmacist contraceptive CE course - not needed if previously completed (about 240 minutes).
 - Study introduction & PRISM training session (about 60 minutes).
 - Provide contraceptive services during the 12-week pilot – 25 total patients per participating pharmacy (time is variable).
 - Complete a digital survey before the pilot begins (about 20 minutes). Complete digital surveys at weeks 4, 8, and 12 during the pilot (about 20-35 minutes per survey).
 - Complete a digital survey 4 weeks after the pilot ends (about 35 minutes).

What You Get

- **Compensation:**
 - Pharmacist contraceptive prescribing CE course paid for by the study - \$250.00 value
 - \$100.00 gift card to attend the study introduction & PRISM training session
 - \$50.00 gift cards for each of the 5 surveys that are completed – total of \$250.00 to complete the study.
 - Full participation value is \$600.00, about \$80.00 per hour for non-clinical participation.
 - Additionally, your pharmacy will be reimbursed \$50 for every contraceptive patient encounter completed and get \$3,000.00 in patient recruitment outreach.
 - **No Cost Training:**
 - Access to a state-approved contraceptive education course providing 0.4 CEUs upon completion.
 - Training on implementing clinical workflows for contraceptive services and how to use the PRISM platform.
 - **Enhanced Skills:**
 - Gain hands-on experience with digital tools for providing contraceptive care.
 - Learn best practices in patient-centered counseling and clinical service implementation.
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Frequently Asked Questions

1. **What is the primary goal of the Clinical trial?**
 - To assess the PRISM platform's impact on pharmacist-led contraceptive services implementation, efficiency, patient outcomes, and usability in real-world pharmacy settings.
 2. **What training will I need to complete?**
 - Complete the contraceptive education course and attend the study introduction and PRISM platform training session.
 3. **How will patient encounters be conducted?**
 - Patient encounters will occur on the PRISM platform, with pharmacists ensuring that the care is patient-centered and compliant with the USMEC guidelines.
 4. **What are the expectations for surveys?**
 - Complete five surveys before, during, and after the pilot to provide feedback on the platform and your experience implementing contraceptive services.
 5. **What if I encounter technical issues?**
 - The OvaryIt team will provide full technical and clinical support throughout the 12-week pilot.
 6. **Can I withdraw from the study?**
 - Yes, participation is voluntary, and you may withdraw at any time without penalty.
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Next Steps:

1. Once your organization joins, you can complete the study onboarding and informed consent process.
2. Complete the contraceptive prescribing CE course (unless already completed).
3. You will receive an email about your new PRISM account and attend the required training session.
4. Use the PRISM platform to provide contraceptive services during the 12-week pilot.
5. Share your valuable feedback through five surveys.

About OvaryIt

OvaryIt, LLC was born from personal experience when our founder faced barriers accessing contraceptives through traditional means, leading to harm when relying on a retail telehealth platform. This experience ignited our mission to revolutionize equitable access to safe women's health services by breaking down barriers in healthcare. We leverage innovative technology and pioneering delivery models to bring essential health services, like contraceptives, directly to those that need them most. Funded by a grant from the National Institutes of Health, we are building the PRISM platform to overcome barriers to the widespread adoption of pharmacist-led contraceptive services, a critical strategy for increasing access.