

~ *List of Clinical Studies* ~

MEDICAL ABORTION

Simplified mifepristone-misoprostol medical abortion

- *Testing 200 mg mifepristone dose followed by home or clinic administration of 400 mcg misoprostol orally two days later up to 56 days LMP.*
Project countries: Ghana, India, Moldova, Mozambique, Nepal, Nigeria, Thailand, Tunisia, Turkey, Ukraine, Vietnam

Expanding options with mifepristone for early medical abortion

- *Comparing two doses of 800 mcg buccal misoprostol to the combined regimen of 200 mg mifepristone followed by 800 mcg buccal misoprostol for gestations up to 63 days' LMP.*
Project countries: Tunisia, Vietnam
- *Comparing sublingual to buccal administration of 400 mcg misoprostol following a 200 mg mifepristone dose for gestations up to 63 days.*
Project country: Moldova
- *Testing sublingual administration of 400 mcg misoprostol following 200 mg mifepristone up to 63 days LMP.*
Project country: Ukraine
- *Comparing 800 mcg buccal misoprostol with 800 mcg oral misoprostol following 200 mg mifepristone up to 63 days LMP.*
Project country: United States
- *Comparing sublingual to oral administration of 400 mcg misoprostol following a 200 mg mifepristone dose.*
Project countries: Moldova, Tunisia and Turkey
- *Testing buccal administration of 800 mcg misoprostol following 200 mg mifepristone up to 63 days LMP.*
Project country: Armenia, Azerbaijan, Mexico and Puerto Rico
- *Comparing 400 mcg buccal misoprostol with 800 mcg buccal misoprostol following 200 mg mifepristone up to 63 days LMP.*
Project country: Georgia
- *Testing one versus two doses of 400 mcg oral misoprostol following mifepristone administration up to 56 days LMP.*
Project countries: India and Vietnam
- *Assessing the need for routine ultrasonography in medical abortion.*
Project country: United States
- *Testing 200 mg mifepristone dose followed by clinic administration of 800 mcg misoprostol vaginally and repeated additional doses of 400 mcg oral misoprostol two days later for gestational ages 10-12 weeks LMP.*
Project countries: India, United States and Vietnam

Medical abortion regimens for pregnancy termination in the second trimester

- *Comparing a combination of mifepristone and misoprostol to misoprostol used alone for termination of pregnancies 14-21 weeks' gestation.*
Project countries: Moldova, Puerto Rico, Tunisia, Turkey and Vietnam

Over

PREGNANCY FAILURE AND MISCARRIAGE

Misoprostol for post-abortion care

- *Comparing 600 mcg dose of oral misoprostol to standard surgical care for treatment of incomplete abortion.*
Project countries: Burkina Faso, Ecuador, Ghana, India, Mozambique, Tanzania, Uganda and Venezuela
- *Comparing misoprostol administered either orally (600 mcg) or sublingually (400 mcg) for treatment of incomplete abortion.*
Project countries: Guatemala, Madagascar and Moldova
- *Comparing 400 mcg dose of sublingual misoprostol to standard surgical care for treatment of incomplete abortion.*
Project countries: Burkina Faso, Egypt, Mauritania, Niger and Senegal

Misoprostol for second trimester intra-uterine fetal death

- *Comparing two different doses of misoprostol (200 mcg vs. 100 mcg) administered buccally as a treatment for fetal death at 14 – 28 weeks' gestation.*
Project country: United States

POSTPARTUM HEMORRHAGE

Misoprostol for prevention of postpartum hemorrhage

- *Comparing 600 mcg dose of oral misoprostol to standard care for prevention of postpartum hemorrhage in home birth settings.*
Project countries: The Gambia and Pakistan

Misoprostol for treatment of postpartum hemorrhage

- *Comparing 800 mcg sublingual dose of misoprostol to oxytocin for treatment of PPH in two circumstances: 1) after prophylactic uterotonics given in the third stage of labor fail to prevent excess bleeding; and, 2) where no prophylactic uterotonics have been given.*
Project countries: Burkina Faso, Ecuador, Egypt, Turkey and Vietnam

Misoprostol for adjunct treatment of postpartum hemorrhage

- *In collaboration with the World Health Organization, Gynuity is conducting a study of misoprostol for adjunct treatment of post-partum hemorrhage (along with oxytocin). Women who are treated for PPH with oxytocin will be randomized also to receive 600 mcg misoprostol sublingually or placebo as an adjunct treatment.*
Project countries: Argentina, Egypt, South Africa, Thailand and Vietnam
- *A similar protocol is being conducted with Aga Khan University in Pakistan in 4 hospitals.*

Blood measurement

- *Comparing measurement techniques for assessing blood loss during normal delivery.*
Project country: India

PRE-ECLAMPSIA

Magnesium sulfate for treatment of pre-eclampsia

- *Testing the low-tech SpringFusor pump for introduction of IV magnesium sulfate.*
Project country: India

STIs/HIV/ INFECTIOUS DISEASE

Woman-controlled products for vaginal health

- *Testing a topical estrogen cream for vaginal self-protection.*
Project country: United States

Clostridium sordellii and perfringens infection

- *Estimating prevalence of vaginal and rectal colonization in women of reproductive age.*
Project countries: United States & Canada