Study to evaluate the pericoital oral contraceptive use of Levonorgestrel 1.5 mg

**Objectives and Background**

**Overall goal:**
This is a study to provide a “proof of concept” for the use of a levonorgestrel oral contraceptive pill that can be taken 24 hours before or after intercourse (pericoital use), up to six times a month. The study recruited 330 women in four country sites to evaluate pregnancy rates, safety information, and acceptability parameters on using on demand contraception, up to six times a month. The last participant follow up ended in July 2014 and analyses was conducted up to January 2015.

**Geographic location**
Bangkok, Thailand; Singapore; Szeged, Hungary; Campinas, Brazil

**Main deliverables**
The technical report and a scientific paper for journal submission will be prepared after the investigators meeting in January 2015.

**Partners**
HRA Pharma

**Sources of funding**
HRP
Gynuity and BMGF for support for ensuring quality data monitoring

**Date Issued**
15 January 2015