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| Title (same as consent form) | Canadian Sexual Health Survey Study |
| Principal Investigator: | Dr. Wendy Norman & Dr. Stirling Bryan |
| Primary Contact:(name, title, phone, email) | Eva McMillan, Research Coordinator, 604-875-2424 x 4871, eva.mcmillan@ubc.ca |
| About the Study:(plain language summary) | This study developed and validated the Canadian Sexual Health Survey (CSHS) that aimed to collect data on sexual and reproductive health among women aged 14-49 years throughout BC. The survey was then administered door-to door by trained health professional surveyors, who finished 1677 surveys in total, within 21 Local Health Authorities, across all five Regional Health Authorities.   |
| Why is this research important? | Data collected through the Canadian Sexual Health Survey will be used to develop Canada’s first contraception Cost-effectiveness Model. This model will be used to create decision-making tools capable of supporting evidence-informed policies for optimal health system strategies, such as the potential for universal subsidy of contraception, in BC and for adaptation to other Canadian provinces, territories and federal health jurisdictions.  |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) | Survey administration and data collection were completed in Fall 2015. The research team is cleaning the survey data and will conduct data analysis to help build the cost-effectiveness model and answer research questions.  |
| Who can participate?(short description, attach consent form) | Participant enrollment for this study has finished.  |
| Study Results/Publication: | Pending |
| Co-Investigators: | Drs. Perry Kendall, Rollin Brant, Janusz Kaczorowski, Steven Shechter, Saied Samiedaluie, Gina Ogilvie, Jean Shoveller, and Sheila Dunn.  |
| Funded by: | Canadian Institutes of Health Research |
| Partners: | Michael Smith Foundation for Health Research; BC Women’s Hospital. |
| Other Attachments (e.g. Newsletters, videos): |  |

**WHRI Website: Research Project Template**

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| Title (same as consent form) | Better Contraceptive Choices: Insertion timing and effectiveness of intrauterine and other contraceptive methods post-abortion |
| Principal Investigator: | Dr. Wendy Norman |
| Primary Contact:(name, title, phone, email) | Weihong Chen, Research Coordinator, 604-875-2424 x 4894, Weihong.chen@ubc.ca |
| About the Study:(plain language summary) | This project includes two randomized controlled trials: 1) Immediate vs. delayed insertion of intrauterine contraception after second trimester abortion; and, 2) Is FlexiT non-inferior to NovaT when inserted immediately after first-trimester abortion? Comparing the effectiveness of copper intrauterine devices available in Canada.BC has a higher rate of abortion per thousand women of reproductive age than any Canadian province other than Quebec. We seek to find the most effective method to prevent unintended pregnancies among women seeking abortion in British Columbia. We hope to see that the interventions investigated in the project, such as inserting the IUC immediately after the second trimester abortion compared to the standard practice that delays the insertion to 4-6 weeks post abortion, will result in fewer unintended pregnancies.  |
| Why is this research important? | This research should produce compelling evidence of the highest scientific quality, permitting a change to the current device indication for world-wide use. We aim for international improvement in the delivery of highly effective long lasting contraception following both first and second trimester abortion. |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) | The uniqueness of this research project is that we have obtained consent from the participants to link the study data to the provincial health administrative data. This allows the team to have a close to 100% follow-up rate of the study outcomes for almost all the participants. The team has been busy analyzing the data and writing the manuscripts for publication.  |
| Who can participate?(short description, attach consent form) | Participant enrollment for this project has finished.  |
| Study Results/Publication: | Pending |
| Co-Investigators: | Study 1: Janusz Kaczorowski, Judith Soon, Stirling Bryan, Rollin Brant, Lyda Dicus, Konia Trouton. Study 2: Jessica Chiles , Caroline Turner. |
| Funded by: | Study 1: Canadian Institutes of Health Research. Study 2: Society of Family Planning |
| Partners: | BC Women’s Hospital. |
| Other Attachments (e.g. Newsletters, videos): | Published Study 1 Protocol: <http://www.ncbi.nlm.nih.gov/pubmed/21672213>Published Study 2 Protocol: <http://www.ncbi.nlm.nih.gov/pubmed/22920273>  |

**WHRI Website: Research Project Template**

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| Title (same as consent form) | Personalized Contraception for Canadians- Decision analytic modelling to support women-centered choices |
| Principal Investigator: | Dr. Flora Teng |
| Primary Contact:(name, title, phone, email) | Weihong Chen, Research Coordinator, 604-875-2424 x 4894, Weihong.chen@ubc.ca |
| About the Study:(plain language summary) | This study will develop and pilot test a Canadian woman-centered contraception decision-making mobile application. This app will be “effectiveness-based” and will incorporate factors women consider important when choosing a contraceptive. Mixed methods will be used including focus groups and interviews to select design elements and feature priorities and to inform iterative pilot testing and improvement cycles of the application. Operations Research methods will be utilized to design decision pathways, and software engineering will be used to enhance usability and human-computer interaction. |
| Why is this research important? | This study aims to enable women to make contraceptive choices that are highly effective and optimal fit to personal preferences. Therefore, they will be able to continue their chosen method, achieve their reproductive goals, and minimize risk of unintended pregnancy. |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) | We have conducted several focus groups to understand the most important and relevant decision criteria in choosing contraception from the perspective of Canadian women. This qualitative data is being analyzed and will be used to create a survey that will specifically rank the contraception priorities of women. The survey will be pilot tested with 120-150 women aged 14-49. Results of the survey will be translated into the Decision-Making model to build the electronic application. |
| Who can participate?(short description, attach consent form) | Reproductive age, English-speaking women from 14 to 49 years old. |
| Study Results/Publication: | Pending |
| Co-Investigators: | Wendy Norman, Saied Samiedaluie, Tymarah Colewah. |
| Funded by: | Women’s Health Research Institute |
| Partners: |  |
| Other Attachments (e.g. Newsletters, videos): |   |

**WHRI Website: Research Project Template**

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| Title (same as consent form) | The CONNECT Project: Integrating family planning and community services for women experiencing intimate partner violence.  |
| Principal Investigator: | Dr. Wendy Norman |
| Primary Contact:(name, title, phone, email) | (For this study, we just want to mention the title, and give the link to the report of the study) |
| About the Study:(plain language summary) |  |
| Why is this research important? |  |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) |  |
| Who can participate?(short description, attach consent form) |  |
| Study Results/Publication: |  |
| Co-Investigators: |  |
| Funded by: |  |
| Partners: |  |
| Other Attachments (e.g. Newsletters, videos): | <http://med-fom-cart-grac.sites.olt.ubc.ca/files/2015/08/CART-Connect-Report-v2-2015-12-22.pdf>  |

**WHRI Website: Research Project Template**

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| Title (same as consent form) | Systematic Review Study: What is the relationship between inter-pregnancy interval and maternal and perinatal outcomes among women who delayed childbearing to 30 years or older? |
| Principal Investigator: | Dr. Wendy Norman |
| Primary Contact:(name, title, phone, email) | (For this study, we just want to mention the title, and give the link to the description of the study at PROSPERO) |
| About the Study:(plain language summary) |  |
| Why is this research important? |  |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) |  |
| Who can participate?(short description, attach consent form) |  |
| Study Results/Publication: |  |
| Co-Investigators: |  |
| Funded by: |  |
| Partners: |  |
| Other Attachments (e.g. Newsletters, videos): | <http://www.crd.york.ac.uk/PROSPERO/display_record.asp?ID=CRD42015019057> |

**WHRI Website: Research Project Template**

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| Title (same as consent form) | Systematic Review of outcomes related to asking about pregnancy intention among reproductive age women presenting for primary care |
| Principal Investigator: | Dr. Wendy Norman |
| Primary Contact:(name, title, phone, email) | (For this study, we just want to mention the title, and give the link to the description of the study at PROSPERO) |
| About the Study:(plain language summary) |  |
| Why is this research important? |  |
| Study Status:(e.g. recruiting, data analysis, manuscript development, complete) |  |
| Who can participate?(short description, attach consent form) |  |
| Study Results/Publication: |  |
| Co-Investigators: |  |
| Funded by: |  |
| Partners: |  |
| Other Attachments (e.g. Newsletters, videos): | <http://www.crd.york.ac.uk/PROSPERO/display_record.asp?ID=CRD42015019726>  |