



PROGRAM UTILIZATION FORM

This Form must be completed if your research study impacts a BC Women's Hospital + Health Centre (BCWH) program or clinic. Refer to the <u>BCWH Program Utilization Form Guidance Notes</u> for information on institutional approval, program utilization, and the submission process. Note that this process generally takes at least 6-8 weeks.

The Programs/Clinics are responsible for determining if these services will have sufficient impact as to require cost recovery. It is the responsibility of the Principal Investigator/Project Lead to ensure proper consultation is done with the Programs/Clinics prior to finalizing the project budget.

Principal Investigator/Project Site Lead Declaration

It is the responsibility of the Principal Investigator (PI)/Project Site Lead to inform the program/clinic and the Women's Health Research Institute (<a href="white:whit:white:white:white:white:white:white:white:white:white:white:whit

If a change in privileges or appointment may occur or has occurred, study approval will be re-reviewed by the program/clinic and by the Women's Health Research Institute.

Please select the declaration option below that best fits with the current research study:

ш	Health Centre of British Columbia.
	As Principal Investigator, I understand it is my responsibility and agree to inform the program/clinic and the WHRI within 4 weeks of any potential or actual change in my BC Women's Hospital + Health Centre medical staff privileges or appointment during the study period.
	Principal Investigator Signature:
	Print Name
	Date
	The Principal Investigator has designated a Project Site Lead to oversee study activities who holds an appointment with the Children's & Women's Health Centre of British Columbia.
	As designated Project Site Lead, I understand it is my responsibility and agree to inform the program/clinic and the WHRI within 4 weeks of any potential or actual change in my BC Women's Hospital + Health Centre medical staff privileges or appointment during the study period.
	Project Site Lead Signature:
	Print Name
	Date

Section 1: Project Information

Study Title:		
REB#:	REB Approval Date:	☐ In progress
Principal Investigator Name:	PI Email:	
Primary Contact Name:	Primary Contact Email:	
Primary Contact Role: (E.g., Researcher, learner-student, resident)	Study Sponsor (if applicable):	
Anticipated start date (in program):	Anticipated end date (in program):	
Summarize the research proposal, including study purp research method (please be brief and use lay language)	,	

Section 2: Supporting Documents

Include the following documents (if applicable) with your PU Form before the signatories can review your request:

Study/Project Protocol RISe (Research Ethics) Application

Research Ethics Approval Certificate

Consent Form(s)/ Waiver of consent

Patient Information Sheet

Recruitment Material (e.g., posters)

Service agreements (e.g., lab services, imaging, pharmaceutical)

Section 3: BC Women's Hospital Program and/or Specific Clinic

One form must be submitted for each program that is impacted by your study.

ACUTE	PROGRAMS
☐ Maternal Newborn Program: ☐ Antepartum/Postpartum Specify Unit(s): (Evergreen, Dogwood, Arbutus, Balsam) ☐ Cedar Birthing Suites ☐ Teck L&D, OB Surgical Services, UCC Specify Area(s): ☐ Perinatal Substance Use (Fir square)	□ Neonatal Program: □ NICU □ Neonatal Follow-up □ MBC
AMBULATO	DRY PROGRAMS
☐ Maternity Ambulatory Program Specify Clinic(s): (I.e., Anesthesia, Antepartum Homecare, Diabetes in Pregnancy, Fetal Assessment, Fetal, Diagnosis Service, Hematology, Infectious Diseases, Internal Medicine, Iron Infusions, Lactation Consultation, Maternal Fetal Medicine, New Beginnings Maternity, Prenatal/Special Procedures, Social Work, Ultrasound).	■ Nurse Practitioner Services Specify Clinic(s): (I.e., After Breast Cancer, Aboriginal Mother's Centre (AMC), Vancouver Women's Health Collective (VWHC), WISH drop-in Centre, Sisterspace Overdose Prevention Site (OPS), Heart Health, Newcomer Services).
☐ Gynecology and Sexual Health Program Specify Clinic(s): (I.e., Chronic Pelvic Pain and Endometriosis, Early Pregnancy Assessment Clinic (EPAC), Recurrent Pregnancy Loss (RPL), ACCESS, Continence, CARE Program)	☐ Gynecology Daycare Surgical Services
☐ Breast Health Program	☐ Oak Tree Clinic
☐ Sexual Assault Service	☐ Provincial Medical Genetics Program
☐ Complex Chronic Diseases Program	☐ Penicillin Allergy Clinic
☐ Other, please specify:	
For a full list of BCWH Services:	http://www.bcwomens.ca/our-services

Section 4 PROGRAM UTILIZATION REQUEST

a) What BCWH Program/Clinic resource(s) are you requesting? Check all that apply.	☐ Staff (e.g., booking clerk, nurse, health records tech) ☐ Infrastructure (e.g., Exam Room, Equipment) ☐ Clinic or Program Records ☐ Parent Advisors (NICU) ☐ Other, please list: ☐ None
b) What tasks are being requested of Hospital Staff for this study?	☐ Introduce research study/staff to patient ☐ Chart flagging ☐ Chart access ☐ Data entry ☐ Sample collection ☐ Other ☐ None
c) How many research participants will be participating at BCWH (in this program specifically)?	
d) Describe what is being requested of Program Staff and/or Program resources for this study. For Acute programs, if more than one clinic area was selected in Section 3, list requests for each area separately. For Ambulatory programs, where applicable, include the following: - Type of resource - Duration (i.e., minutes/hours) - Time (of day) - Frequency (weekly, ad hoc) - Start Date - End Date e) Describe study activities conducted in the Program by non-Program Staff. e.g., Research staff, trainees, research nurse	
f) If your study requires participant recruitment within a program, how will your study representative be introduced to the patient or family member?	
g) How will program staff be oriented to the study (or trained) if necessary?	

h) How will the research results be shared with the program?	
i) If required by the program, is funding available to support any requested BCWH Program/Clinic resources?	Yes No
j) Please include any additional information about your study that would help during our review.	
k) Would you like to promote your study on the BC Women's Hospital website?	Yes No

Please see next page for required signatures:

For Acute Programs, please see Section 5.1

<u>For Ambulatory Programs</u>, please see Section **5.2.A**; for <u>Provincial Medical Genetics Program</u> see Section **5.2.B**

To obtain signatures, please submit your PU Form request to:

Acute: Maternal Newborn Programs

• Submit completed form and supporting documentation to Jesse Veenstra (<u>jesse.veenstra@bccdc.ca</u>) who will assist with obtaining all necessary signatures.

Acute: Neonatal Programs

- Step 1: Contact Lindsay Richter (<u>lindsay.richter@cw.bc.ca</u>) prior to submission of the PU Form to schedule a presentation at the NICU Research and Quality rounds.
- Step 2: Submit the completed form and supporting documentation to Lindsay who will assist with obtaining all necessary signatures.

Ambulatory Programs (including the Provincial Medical Genetics Program)

• Submit completed form and supporting documentation to the appropriate Program Manager as identified in the <u>Signatories List.</u> If you have any questions about your submission, please contact Carola Muñoz (<u>carola.munoz@cw.bc.ca</u>).

Section 5.1: Required Signatures (ACUTE PROGRAMS) For a full list of signatories, click <u>here</u>

	nandwritten, scanned signature, or signature line in box below:	
Print		
Date		
Progra	ram Medical Lead Signature	
Add h	nandwritten, scanned signature or signature line in box below:	
Drint	Name	
Date	ivame	
	r Director	
Add h	nandwritten, scanned signature or signature line in box below:	
Print Date	Name	
	r Medical Director	
Add n	nandwritten, scanned signature or signature line in box below:	
Date	Name	
	Senior Director/Senior Medical Director signature is obtained, please submit to the vertice of the Director (Rm H214 c/o Lori Brotto)	ne office of the
	ntive Director, Women's Health Research Institute Signature nandwritten, scanned signature or signature line in box below:	
Print Date	Name	
Dute		

For program use only. Notes/ Comments/Additional Information Required:

Section 5.2.A: Required Signatures (AMBULATORY PROGRAMS) For a full list of signatories, click <u>here</u>

Print Name		
Date		
Program Medical Lead Add handwritten, scan	Signature ned signature or signature line ir	n box below:
[
Print Name		
Date		
Senior Patient Services	Director	
	ned signature or signature line ir	n box below:
Print Name		
Date		
Senior Medical Directo		
	ned signature or signature line ir	n box below:
Print Name		
Date		
Once Senior Director/See	nior Medical Director signature is a	obtained, please submit to the office of th
Executive Director (Rm H		obtained, piedse submit to the office of the
	men's Health Research Institute ned signature, or signature line i	
rada nandwirtten, scam		

Section 5.2.B: Required Signatures (Provincial Medical Genetics Program) *For a full list of signatories, click <u>here</u>*

	Program Operations Director Signature
	Add handwritten, scanned signature, or signature line in box below:
	Print Name
	Date
	Program Medical Director Signature
	Add handwritten, scanned signature or signature line in box below:
	Print Name
	Date
<u></u>	
	Senior Medical Director
	Add handwritten, scanned signature or signature line in box below:
	Print Name
	Date
Г	Chief O and a Office PCW and black to the Health Control
	Chief Operating Officer, BC Women's Hospital + Health Centre Add handwritten, scanned signature or signature line in box below:
	And Hallawritterl, Scarlined Signature of Signature life in Box Below.
	Print Name Date
	Date
е.	Senior Director/Senior Medical Director signature is obtained, please submit to the office of the WHRI
ıti	ive Director (Rm H214 c/o Lori Brotto)
	Executive Director, Women's Health Research Institute Signature
	Add handwritten, scanned signature, or signature line in box below:
	Print Name

For program use only. Notes/ Comments/Additional Information Required: