Patient Label Here



ADMISSION TAB	Maternal Transfer from: (Select One)
DEMOGRAPHICS : Per patient label <i>or</i>	□ No transfer □ Hospital □ Planned Home or Clinic Birth
Family Name:	□ Nursing station □ Birthing Center
Given Name:	□ Other unit same hospital □ Other
Maternal Date of Birth: dd / mm /yyyy	IF TRANSFER
Chart Number/Client ID:OHIP:	IF TRANSFER:
Address:	Maternal Transfer from Hospital (name):
Postal Code:Phone:	
□ No Fixed Address	Maternal transfer from Birth Centre (name):
Estimated Date of Birth (EDB): dd/mm/yyyy	
Primary Language: (Select One)	Reason for Maternal Transfer From: (Select One)
□ English □ French □ Unknown	☐ Fetal health concern ☐ Lack of nursing coverage
□ Other (specify):	□ Lack of physician coverage
	. □ Maternal medical/obstetrical problem
MATERNAL ADMISSION TO HOSPITAL	□ No beds available □ Organization evacuation
Admission date: dd / mm / yyyy Admission Time:	☐ Birth outside of hospital prior to admission
Admission by Healthcare Provider: (Select One) Obstetrician	□ Other □ Unknown



HISTORY TAB Pre-existing Health Conditions (Outside of Pregnancy): (List All) Mental Health Concerns: (Select All That Apply) □ None □ Anxiety □ Depression ☐ History of Postpartum Depression ☐ Addiction ☐ Bipolar □ Schizophrenia □ Other □ Unknown **Domestic/Intimate Partner Violence:** (Select One) □ No Disclosure □ Disclosure □ Unable to ask Obstetrical History: Gravida (G): # of Previous Term Pregnancies (T): # of Previous Preterm Pregnancies (P): # of Previous Abortions (A): # of Living Children (L): # of Previous Stillbirths (S): # of Previous Vaginal Births: # of Previous C/S Births: # of Previous VBACs: _____ Parity: Auto calculates

PREGNA	ANCY TAB		
Materna	l Height:	(in, ft & in, cm)	□ Unknown
Pre-preg	ınancy weight	:(lb/kg)	□ Unknown
Pre-preg	nancy BMI: C	alculates	
Maternal Weight at end of Pregnancy:(lb/kg)			(lb/kg)
□ Unknov	vn □ Declined	l weight check	
Maternal Weight Gain at end of Pregnancy: Calculates			
Number of Fetuses:			
Is the pregnant person a gestational carrier? (Select One)			
□ Yes □ No □ Unknown			
Estimated Date of Birth (EDB): dd / mm / yyyy			
Conception type: (Select One)			
□ Spontaneous			
□ Intrauterine Insemination alone			
\square Intrauterine Insemination (IUI) with ovulation induction but no IVF			
□ Ovulation induction without IVF (i.e. Clomid, FSH)			
□ IVF □ Vaginal insemination □ Unknown			

□ Unknown if patient/client received prenatal education



	Was an adul as a the same and a self-
EDB determined by: (Select One)	Was prenatal genetic screening offered, as indicated on the OPR?: (Select One)
□ Last Menstrual Period	□ Yes, screening was offered
□ First trimester dating ultrasound	□ No, screening was not offered
□ Second trimester ultrasound	□ Counselled and declined screening
□ Assisted reproductive technology	□ Unknown if screening was offered – no access to the OPR
□ Obstetrical clinical estimate (includes S-F height)	□ Unknown if screening was offered – other reason
□ Unknown First Trimester Visit: (Select One) □ Yes □ No □ Unknown	Folic Acid Use: (Select One) □ None □ Pre-conception only □ During pregnancy only □ Pre-conception and during pregnancy □ Unknown
Antenatal Health Care Provider: None Obstetrician Family Physician Midwife Nurse Nurse Practitioner (APN/CNS) Other Unknown	Intention to Breastfeed: (Select One) Select One) Select One Select One) Select One) Select One) Select One)
Prenatal Education: (Select One) Select One) Yes - In-person prenatal education only Yes - Online prenatal education only	milk substitute) □ No, does not intend to breastfeed □ Mother unsure □ Unknown, intent not collected
☐ Yes - Combination of in-person and online prenatal education	
□ Yes - Unknown method of education delivery	
□ No - Patient/client did not receive prenatal education	



□ None □ < 1 drink/month □ 1 drink/month □ 2-3 drinks/month □ 1 drink/week □ More than 1 drink/week □ Episodic excessive drinking (binging) □ Exposure prior to pregnancy confirmed, amount unknown □ Unknown
Cannabis Exposure in Pregnancy: (Select One) Never
□ Some use, but frequency unknown □ Usage unknown Drug and Substance Exposure in Pregnancy: (Select All That Apply) □ None □ Amphetamines
□ Cocaine □ Gas/Glue □ Hallucinogens □ Opioids □ Other □ Unknown



ANTENATAL EXPOSURE TO MEDICATION:

(0	T None	Opiola Agonist Therapy:	
(Select All That Apply)	□ None	□ Methadone	□ Buprenorphine monoproduct (Subutex)
OTC/Vitamins/Homeop	pathic:	□ Buprenorphine	– naloxone (Suboxone)
□ Prenatal Vitamins (inc	luding folic acid)	□ Slow-release m	norphine for opioid use disorder
□ Probiotics □ Iron Sup	pplements		
□ Anti-emetics (OTC)	□ Antihistamines (OTC)	Other Medicatio	ns:
□ Herbal or homeopathi	c remedies	□ Psychotropics	□ Selective Serotonin Reuptake Inhibitors
□ Other over the counte	r medications	□ Thyroid medico	ations 🗆 Other prescription
Prescribed Medication	s:	□ Unknown presc	ription or OTC exposure
□ Anticonvulsants (NOT : □ Anti-emetics □ Anti-inflammatory □ □ Anti-inflammatory □ Anti-rheumatic □ Anti-r	atihistamines	□ None □ C-Diff □ Group B Strept □ Hepatitis B □ □ HPV □ Seasor □ Methicillin-resis	REGNANCY: (Select All That Apply) ficile
		If Yes To Covid In	
		·	COVID-19 Diagnosis: dd/mm/yyyy
		was the patient r	nospitalized due to COVID-19 specifically?

□ Yes

□ No

□ Unknown



GBS Screening Results (35-37 wks): (Select One)	BLOOD TYPING AND IMMUNOGLOBULIN
□ Not Done □ Done, negative result □ Done, positive result □ Done, result unknown □ Unknown if screened GBS Screening Date (if done): <u>dd/mm/yyyy</u>	Blood group and type of pregnant individual, ABO/Rh(D): (Select One)
Reason GBS Screening Not Done: (Select One)	What was the antibody screen result?:
□ Previous baby with GBS disease	□ Negative □ Positive □ Unknown
□ Previous GBS screening done in this pregnancy □ Urine positive for GBS □ Declined Screening □ Other □ Unknown	For Rh(D) negative patients, was Rh(D) immunoglobulin (RhIG/Rhogam/WinRho) given in pregnancy?: □ No □ Yes, 1 dose □ Yes, 2 doses □ Yes, 3 or more doses
Progesterone taken daily for spontaneous preterm birth prevention, any time after 16 weeks gestation:	☐ Yes, number of doses unknown ☐ Unknown
□ Yes □ No □ Unknown	
(Do NOT include if progesterone is used only in first trimester) ASA (aspirin) taken daily for preeclampsia prevention, any time after 12 weeks' gestation: Yes No Unknown	Date of Rh(D) Immunoglobulin Dose (latest prior to birth): dd/mm/yyyy
(Do NOT include if aspirin is used only in first trimester)	



DIABETES AND PREGNANCY: (Select One)	Complications of Pregnancy – Maternal:
	□ Anemia unresponsive to therapy
□ None □ Gestational - Insulin □ Gestational - No Insulin	□ Antepartum bleeding (persistent and unexplained)
□ Gestational - Insulin status unknown □ Type 1	□ Cancer – diagnosed in this pregnancy
□ Type 2 Insulin □ Type 2 No Insulin	□ Haemotology – Gestational Thrombocytopenia
□ Type 2 Insulin Usage Unknown □ Type Unknown	□ Hyperemesis Gravidarum (Requiring Hospital Admission)
□ Declined Testing □ Unknown	□ Liver/Gallbladder – Intrahepatic Cholestasis of Pregnancy
LIVERTENANT DISCRETE OF PRECNANCY (C. L. C.)	□ Liver/Gallbladder – Acute Fatty Liver of Pregnancy
HYPERTENSIVE DISORDERS OF PREGNANCY: (Select One) □ None □ Gestational Hypertension □ Preeclampsia	□ Neurology – Epilepsy/Seizures – Seizure occurred during current pregnancy
□ Pre-existing Hypertension with superimposed preeclampsia	□ Prelabour rupture of membranes (PROM)
□ Eclampsia □ HELLP syndrome □ Unknown	□ Preterm labour
	□ Preterm pre-labour rupture of membranes (PPROM)
COMPLICATIONS OF PREGNANCY, NOT INCLUDING HYPERTENSION OR DIABETES: (Select All That Apply)	□ Pulmonary – asthma occurred during current pregnancy
TITE EXTENSION ON DIABETES. (Select All That Apply)	□ Other
Complications of Pregnancy, not including Hypertension or Diabetes: None Unknown	Complications of Pregnancy – Placental:
Complications of Pregnancy – Fetal:	□ Placenta accreta □ Placenta Increta □ Placenta percreta
□ Anomaly(ies) □ Isoimmunization/Alloimmunization	□ Placenta Previa □ Placental abruption □ Other
□ Intrauterine Growth Restriction (IUGR)	
□ Oligohydramnios □ Polyhydramnios □ Other	



Type of Labour: (Select One) **INTRAPARTUM TAB** ☐ Active labour achieved without any intervention **Antenatal Steroids:** (Select One) □ Induced labour in latent phase □ None □ 1 dose < 24 hours (before the time of birth) □ Induced labour prior to onset of contractions □ 2 doses: Last dose < 24 hours (before the birth) ("cold induction") □ 2 doses: Last Dose > 24 hours (from the time of the last dose □ No labour or latent phase to the time of birth) **Cervical ripening/induction methods:** (Select All That Apply) □ Unknown □ None □ Prostaglandin (PGE2) Fetal Surveillance: (Select All That Apply) □ Mechanical (Foley catheter) □ Laminaria tents ☐ Misoprostol (PGE1) ☐ Other ☐ Unknown □ Admission EFM Strip □ Auscultation □ Intrapartum EFM (external) □ Intrapartum EFM (internal) Was oxytocin used any time before birth? ☐ Yes ☐ No □ No Monitoring □ Unknown Cervical dilation at start of oxytocin: **Group B Strep Antibiotics:** (Select One) Start date of oxytocin: ☐ Yes ☐ No ☐ Declined antibiotics ☐ Unknown Start time of oxytocin: □ Unknown Membrane Rupture: (Select One) Initial cervical dilation (cm) upon hospital admission for ☐ Artificial rupture of membranes labour and birth: ☐ Spontaneous rupture of membranes ☐ Unknown Date of Membrane Rupture:

Time of Membrane Rupture:



STAGES OF LABOUR	IF INDUCED LABOUR:
First Stage	All Indications for Induction of Labour: (Select All That Apply)
Date of latent phase onset:	Fetal Indications:
Time of latent phase onset:	□ Atypical or abnormal fetal surveillance
□ Unknown	□ Fetal anomaly/ies □ Intrauterine Fetal Death (IUFD)
Date of active phase onset:	□ Isoimmunization/alloimmunization □ IUGR □ Macrosomia
Time of active phase onset:	☐ Multiple gestation ☐ Other fetal complication ☐ Post dates
□ Unknown	□ Termination of pregnancy
Second Stage	Maternal Indications:
•	□ Abnormal Biomarkers (eg. PAPP_A, PIGF, and HCG)
Date fully dilated:	- Cholesiasis of Fleditatics
Time fully dilated:	□ Diabetes □ Elevated BMI
□ Unknown	□ Hx of Precipitous Birth
Date started pushing:	☐ Hx of Previous of Intrauterine Fetal Death
Time started pushing:	
□ Unknown	Other obstetrical complications/concerns
	□ Polyhydramnios □ Preeclampsia/Hypertension
	□ Pre-existing maternal medical conditions
	□ Pregnant individual age >= 40
	□ Pre-labour rupture of membranes (PROM)
	□ Preterm Pre-labor rupture of membranes (PPROM)
	□ Prolonged Latent Phase Labour



Other Indications:	* If Transfer to Other Hospital, ICU/CCU,
	or Other Non-Obstetrical Unit, same hospital:
□ Accommodates care provider/organization	Reason for Maternal Transfer: (Select One)
□ Distance from birth hospital/safety precaution	□ Fetal Health Concern □ Lack of Nursing Coverage
□ Maternal request □ Unknown	
Primary Indication for Induction of Labour:	□ Lack of Physician Coverage□ Maternal medical/obstetrical problem□ No beds available
Dish on Cooper City	□ Organization evacuation □ Other □ Unknown
Bishop Score: Circle 0 1 2 3 4 5 6 7 8 9 10 11 12 13 □ Unknown	Maternal Transfer Date: dd / mm / yyyy
	Maternal Transfer Time:
	* If Transferred:
ALL LABOUR TYPES - SPONTANEOUS, INDUCED	Pharmacologic Pain Management: (Select All That Apply)
AND NO LABOUR	□ None □ Nitrous oxide □ Opioids □ Epidural □ Spinal
Maternal Outcome (prior to birth): (Select One)	□ Spinal-epidural combination □ Pudendal □ Unknown
□ No Transfer □ Transfer to other organization	Labour and Birth Complications: (Select All That Apply)
□ Transfer to ICU/CCU	□ None □ Atypical or abnormal fetal surveillance
□ Transfer to other non-obstetrical unit, same hospital	□ Meconium □ Cord prolapse □ Shoulder dystocia
□ Maternal Death—Not Related to Pregnancy or Birth	□ Fever>38.5 C □ Non-progressive first stage of labour
□ Maternal Death—Related to Pregnancy or Birth	□ Non-progressive second stage of labour
Ç ,	□ Placental abruption □ Uterine rupture
* If Transfer to Other Organization:	□ Uterine dehiscence □ Retained placenta-manual removal
Maternal Transfer to [hospital name]:	□ Retained placenta-surgical removal
	□ Postpartum hemorrhage □ Uterine atony
	□ Perineal hematoma □ Amniotic fluid embolism
	□ Pulmonary embolism □ Hysterectomy □ Other □ Unknowi



BIRTH TAB	Birth Location: (Select One) □ Hospital □ Home
Type of Birth: (Select One) □ Vaginal Birth □ Cesarean Birth	□ Birth Centre □ Clinic (Midwifery) □ Nursing Station □ Other Ontario location □ Outside of Ontario
PRESENTATION TYPE (Select One) Cephalic: □ Vertex □ Brow □ Face □ Compound □ Cephalic type unknown	Birth Hospital name: Date placenta delivered: dd / mm / yyyy Time placenta delivered:
Breech: □ Frank □ Complete □ Incomplete	IF CESAREAN BIRTH:
□ Footling □ Compound □ Breech type unknown	Type of Cesarean birth: (Select One)
Other: □ Transverse/Malpresentation □ Unknown	□ Planned (as scheduled) □ Planned (not as scheduled)
Newborn DOB: dd/mm/yyyy	□ Unplanned
Time of birth:	Dilation at Cesarean Birth (cm):
Forceps/Vacuum used vaginally: (Select One) □ None	Anesthesia for Cesarean birth: (Select One)
□ Vacuum □ Forceps □ Vacuum and Forceps □ Unknown	□ Epidural □ Spinal □ Spinal-Epidural Combination
Episiotomy: (Select One)	□ General □ Other □ Unknown
□ None □ Medio-lateral □ Midline □ Unknown	ALL INDICATIONS FOR CESAREAN BIRTH:
Perineal Laceration: (Select All That Apply) □ None	(Select All That Apply)
□ 1st degree □ 2nd degree □ 3rd degree □ 4th degree □ Cervical tear □ Other □ Unknown	Fetal: □ Anomaly(ies) □ Atypical or Abnormal Fetal Surveillance □ Cord prolapse □ Intrauterine Growth Restriction (IUGR) □ Macrosomia

□ Malposition/Malpresentation □ Other Fetal Indication



Maternal: □ Cholestasis of pregnancy	Primary indication for Cesarean birth:
□ Failed forceps/vacuum □ Failed induction	
□ Gestational hypertensio	Labour and/or Birth Complications: (Select All That Apply)
□ HIV – Human immunodeficiency Virus	□ None
□ HSV – Herpes Simplex Virus	□ Atypical or abnormal fetal surveillance □ Meconium
□ Hypertensive Disorders of Pregnancy – Eclampsia	□ Cord prolapse □ Shoulder dystocia □ Fever>38.5 C
□ HELLP □ Preeclampsia □ Maternal Health condition(s)	□ Non-progressive first stage of labour
□ Multiple gestation □ Non-progressive first stage of labour	□ Non-progressive second stage of labour
□ Non-progressive second stage of labour □ Obesity	□ Placental abruption □ Uterine rupture
□ Other Obstetrical complication	□ Uterine dehiscence □ Retained placenta-manual removal
□ Placenta Increta/Acreta/Percreta □ Placenta previa	□ Retained placenta-surgical removal
□ Placental abruption	□ Postpartum hemorrhage □ Uterine atony
□ Prelabor rupture of membranes (PROM) in pregnant	□ Perineal hematoma □ Amniotic fluid embolism
individuals with a planned cesarean birth	□ Pulmonary embolism □ Hysterectomy □ Other □ Unknown
□ Preterm pre-labor rupture of membrances (PPROM) in	Intrapartum Medications Administered: (Select All That Apply)
pregnant individuals with a planned cesarean birth — Previous cesarean birth	□ None
	□ Magnesium Sulfate for preeclampsia
□ Previous T incision/classical incision/uterine surgery	□ Magnesium Sulfate for fetal neuroprotection
□ Previous uterine rupture □ Suspected chorioamnionitis □ Uterine rupture □ Declined VBAC □ VBAC - Failed Attempt □ VBAC - Not Eligible	□ Antibiotics (not for GBS) □ Antihypertensives
	□ Anti-emetics □ Antipyrexics (example: acetaminophen)
	□ Diuretics □ Insulin
Other: Accommodates care provider/organization	□ Tocolytics (Mag sulfate/indomethecine/nifedipine/ritodrine etc)
□ Maternal request □ Unknown	□ Other □ Unknown



Pharmacologic Pain Management: (Select All That Apply) None Nitrous oxide Opioids Epidural Spinal Spinal-epidural combination Pudendal Unknown Supportive Care: (Select All That Apply) None 1:1 Supportive care by clinical staff/care provider Breathing exercises Hypnobirthing/guided imagery	Other Care Providers Present at time of Labour and/or Birth: (Select All That Apply) Family Physician
□ Massage □ Shower □ Sterile water/saline injections □ Support partner or doula □ TENS □ Tub □ Other □ Unknown	□ Clinical Nurse Specialist/Nurse Practitioner □ Doula □ Other Care Provider □ None □ Unknown OUTCOME TAB
Healthcare Provider Who Caught/Delivered Baby: (Select One) Family Physician Registered Midwife Obstetrician Resident Surgeon Registered Nurse Nurse Practitioner (CNS/NP) Aboriginal Midwife Midwifery Student Unattended (None) Other Health Care Provider Unknown ID of Healthcare Provider Attending Birth: (Optional Field)	Pregnancy Outcome (Complete for each fetus if multiple pregnancy): (Select One) Live birth Stillbirth >= 20 wks or >= 500 gms - Spontaneous - occurred during antepartum period Stillbirth >= 20 wks or >= 500 gms - Spontaneous - occurred during intrapartum period Stillbirth >= 20 wks or >= 500 gms / Termination Stillbirth >= 20 wks or > = 500 gms / Termination Pregnancy loss < 20 wks and < 500 gms / Spontaneous miscarriage Pregnancy loss < 20 wks and < 500 gms / Termination

Gestational age at birth: Auto-calculates





Maternal Birth Outcome: (Select One)
□ No Transfer □ Transfer to other organization
□ Transfer to ICU/CCU
□ Transfer to other non-obstetrical unit, same hospital
□ Maternal Death—Not Related to Pregnancy or Birth
□ Maternal Death—Related to Pregnancy or Birth
*IF TRANSFER TO OTHER HOSPITAL:
Maternal Transfer to [hospital name]:
*IF TRANSFER TO OTHER HOSPITAL, ICU/CCU, OR OTHER NON-OBSTETRICAL UNIT, SAME HOSPITAL:
Reason for Maternal Transfer To: (Select One)
□ Fetal Health Concern □ Lack of Nursing Coverage
□ Lack of Physician Coverage
□ Maternal medical/obstetrical problem □ No beds available
□ Organization evacuation □ Care Closer to Home
□ Other □ Unknown
Maternal Transfer Date: dd / mm / yyyy
Maternal Transfer Time:or
Maternal Discharge Date: dd / mm / yyyy
Discharge Time:



Prior to birth, when was the maternal transport to hospital?

(Select one) □ First stage □ Second stage

MIDWIFERY TAB	Reason(s) for Transport: (Select all that apply)
	□ Fetal well-being concerns □ Pain management
Was care of client transferred back to Midwifery during intrapartum?*: □ Yes □ No	□ Prolonged labour □ Maternal request
If there was transfer of care (w/o a return to care) in a previous encounter	□ Neonatal condition/complication □ Provider preference
	□ Other maternal condition/complication
	□ Other fetal condition/complication
NTRAPARTUM	Primary Reason for Transport:
Began Intrapartum Period intending to give birth at: (Select One) Hospital Home Birth Centre	Did EMS attend the labour? (Select one)
□ Clinic (Midwifery) □ Other □ Nursing Station	□ Yes □ No □ Unknown
□ Undecided	Was EMS used to transport to hospital prior to birth?
Actual Location of Labour: (Select One)	(Select one) □ Yes □ No □ Unknown
□ Hospital □ Home □ Birth Centre □ Clinic (Midwifery)	Barrier to Transport: (Select all that apply)
□ Other □ Nursing Station	□ None □ Delayed arrival time of EMS
Birth Centre of Labour: (Select One)	□ Delayed Departure of EMS □ Delay on route □ Other
(select only if labour at Birth Centre)	
□ Ottawa Birth and Wellness Centre	Did midwife attend client at home at any point during labour? (Select one) □ Yes □ No □ Unknown
□ Toronto Birth Centre, Inc.	
Was there unplanned Maternal transport to hospital at any part of the labour? (Select one) Yes No Unknown	

 \square Other fetal condition/complication



FOR BIRTHS THAT TOOK PLACE AT HOME:	Primary Reason for Transport: (indicate)
Initial cervical dilation (cm) upon midwife's arrival at the home to attend labour and birth (home births only):	Did EMS attend the actual location of labour at any part of the birth or immediate postpartum?
Other care providers present at time of labour and/or birth: (Select all that apply)	Was EMS used to transport to hospital? Yes No Unknown Barrier to Transport: (Select all that apply)
*If there was transfer of care (w/o a return to care) in a previous encounter:	□ None □ Delayed arrival time of EMS □ Delayed Departure of EMS □ Delay on route □ Other IF VAGINAL BIRTH:
Was care of client transferred back to Midwifery during birth? Yes No	Maternal Position at Time of Birth: (Select one) Supine Semi-fowler's Lateral Standing
Was there unplanned Maternal transport to hospital at any part of the birth or immediate postpartum?	□ Squatting □ Kneeling □ All-fours □ Lithotomy □ McRoberts □ Birth Stool □ Other □ Unknown
□ Yes □ No □ Unknown	IF SPONTANEOUS VAGINAL BIRTH:
Reason(s) for Transport: (Select all that apply) □ Fetal well-being concerns □ Pain Management	Was the baby born in the water? □ Yes □ No □ Unknown
□ Prolonged labour □ Maternal request □ Neonatal condition/complication □ Provider preference □ Other maternal condition/complication	Was this a planned water birth? □ Yes □ No □ Unknown

of practice? □ Yes □ No



IF VAGINAL BIRTH:	Labour/Birth Transfer of Care?: □ Yes □ No
Components of third stage management employed (unrelated to corrective measures for bleeding):	IF YES:
(Select all that apply)	Was rationale for transfer of care only because of hospital physician protocol, and not because of midwifery judgeme or scope of practice? □ Yes □ No
□ None □ Breastfeeding □ Controlled cord traction	
□ Early cord clamping □ Prophylactic uterotonic	
□ Unknown – Midwife was NOT most responsible provider during birth	AND:
□ Unknown – Other	Was the transfer of care returned anytime from onset of active labour to approximately 1 hour post-birth? Yes No Was the client discharged from midwifery care during intrapartum / immediate postpartum? (Select yes to discharge client from Midwifery Care and/or bill for Course of Care) Yes No
Was there a known midwife or midwifery student at the birth? □ Yes □ No □ Unknown	
Were there any labour/birth consults, transfers of care, from the onset of active labour to approximately 1-hour post-birth? ☐ Yes ☐ No	
Reason(s) for labour/birth consultation/transfer of care?	
Labour/Birth Consult with Physician? ☐ Yes ☐ No	
IF YES:	
Was rationale for consult only because of hospital/physician protocol, and not because of midwifery judgement or scope	