

Government of Western Australia North Metropolitan Health Service Women and Newborn Health Service



#### OBSTETRICS AND GYNAECOLOGY CLINICAL PRACTICE GUIDELINE

# Intrauterine Pressure Catheter

Scope (Staff):	WNHS Obstetrics and Gynaecology Directorate staff		
Scope (Area):	Obstetrics and Gynaecology Directorate clinical areas at KEMH, OPH and home visiting (e.g. Visiting Midwifery Services, Community Midwifery Program and Midwifery Group Practice)		

#### This document should be read in conjunction with this Disclaimer

# Aims

- To measure the frequency, duration and pressure of uterine contractions in cases where uterine activity is not readily palpable e.g. maternal obesity.<sup>2</sup>
- To collect an amniotic fluid sample for laboratory analysis.
- To perform <u>amnioinfusion</u>

# Key points

- 1. Intrauterine pressure catheters are not recommended in routine intrapartum fetal surveillance.<sup>2</sup>
- 2. Insertion of intrauterine pressure catheter (IUPC) is to be done by Obstetric Registrar, Senior Registrar or Consultant.
- 3. The IUPC should not be left in situ for longer than 24 hours.
- 4. A pressure sensor in the tip of the IUPC measures changes in the amniotic fluid pressure in response to contractions.<sup>1</sup>

# Risk factors / complications

- Increased risk of postpartum haemorrhage and need for blood transfusion<sup>3</sup>
- Uterine perforation<sup>5</sup>

• Extramembranous placement occurs 14-38% of the time, with adverse events occurring in 1/1400 placements.<sup>4</sup>

## Contraindications

Do not insert if there is bleeding of unknown origin, placenta previa or non-ruptured amniotic membranes (as per manufacture).

# Equipment

- Sterile catheter pack
- Sterile gloves
- IUPC cable for CTG
- Lubricating gel
- Cardiotocograph (CTG) monitor
- Adhesive tape, optional

## Procedure

#### Prior to the procedure

- 1. The obstetric Consultant or Senior Registrar should assess the suitability for insertion of an IUPC.
  - Including confirmation of placental location
- 2. Obtain verbal consent from woman.
- 3. Ensure the catheter, cable and CTG monitor are compatible **before** insertion.
  - Plug the IUPC cable into the CTG monitor.
- 4. Read the manufacturers instruction in or on the packaging of the IUPC regarding insertion.
- 5. Place woman in dorsal position with wedge below right buttock.

#### Procedure

- 1. Remove catheter from package using aseptic technique.
- 2. Zero the system.
- 3. Perform vaginal examination to;
  - Confirm cervical dilatation
  - Confirm or perform rupture of membranes
- 4. Insert the introducer and catheter into the vagina and to the cervical os. Do not advance the introducer through the cervix.
- 5. Attempt to insert the catheter opposite to the placental site.

- 6. Gently advance the catheter into the uterus. If resistance is met at any time during insertion.
  - Pull the catheter tip back to the introducer and alter the direction of the catheter by changing direction of the introducer
  - Determine an alternative position for placement
  - If resistance continues cease insertion of transducer
- 7. Remove the introducer by gently sliding back out of the vagina.
- 8. Secure the catheter to the woman's leg.
  - The catheter should be secured as close as possible to the introitus to prevent the catheter from working its way out of the uterus when it is flexed.
- 9. Zero the CTG monitor as required.
- 10. Connect the catheter to the cable.
- 11. Instruct the woman to cough.

12. A spike on the CTG tracing in response to a cough indicates correct positioning.

#### After the procedure

- 1. Document the time of insertion in the medical record or Phillips Intellispace Perinatal (PIPS).
- 2. If an amniotic fluid sample is required remove the cap from the amnio port and collect the sample.

# Troubleshooting

If the IUPC is not recording:

- Ensure the catheter, cable and CTG monitor are compatible before insertion.
- Check the cables are plugged in and all connections are correct.
- Disconnect the catheter from the cable and inject 10mL of sterile 0.9 % sodium chloride through the amnioport. Reconnect the cap and cable.
- Liaise with the Medical Officer who may decide to disconnect the catheter from the cable, rotate, retract or advance the catheter. Wait 15 seconds before reconnection.

### Removal of IUPC

- The IUPC can be removed by midwifery or medical staff.
- Grasp the catheter and gently pull until fully withdrawn.
- Disconnect the catheter from the cable.

#### References

- 1. Cunningham F, Leveno J, Bloom S, Dashe J, Hoffman B, Spong C, Casey, B. **Williams Obstetrics.** 26th. United States: McGraw Hill; 2022
- 2. Royal Australian and New Zealand College of Obstetricians and Gynaecologists. Intrapartum Fetal Surveillance Clinical Guideline. Fourth Edition. 2019.
- 3. Haizler-Cohen L, Baxter D, Sanghavi K, Fahimuddin F, Chornock R, Saeed H, Iqbal S, Mokhtari N. Intrauterine pressure catheter use and risk of placental abruption and postpartum hemorrhage. **American Journal of Obstetrics and Gynaecology.** 2023. Available from: <u>https://doi.org/10.1016/j.ajog.2022.11.902</u>
- 4. Bakker J, Janssen P, van Halem K, van der Goes B, Papatsonis D, van der Post J, Mol B. Internal versus external tocodynamometry during induced or augmented labour. **Cochrane Database of Systematic Reviews**. 2013 (3). Available from: <u>Internal versus external tocodynamometry</u> <u>during induced or augmented labour - Bakker, JJH - 2013 | Cochrane Library</u>
- 5. Richards E, Rehmer J, Falcone T. Perforation During Gynecological Procedures. **JAMA.** 2023. <u>Perforation During Gynecological Procedures | Health Care Safety | JAMA | JAMA Network</u>

#### Related WNHS policies, guidelines and procedures

WNHS Obstetrics and Gynaecology Clinical Guideline: Amnioinfusion

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NSQHS Standards (v2) applicable:	<ul> <li>I: Clinical Governance</li> <li>I: Clinical Governance</li> <li>I: Partnering with Consumers</li> <li>I: Preventing and Controlling Healthcare Associated Infection</li> <li>I: Medication Safety</li> </ul>	olling			
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Version History

Version Number	Date	Summary		
1.0 - 5.0	July 2003; April 2008; February 2011; July 2014 Issue date unknown; regular review as above			
6.0	10 May 2018	Added clinical circumstances that intrauterine pressure transducer is required; updated procedure		
7.0	14 October 2024	Updated to new template; added contraindications; revised risk factors/complications; reviewed references and updated guideline accordingly; minimal changes to procedure itself; expanded removal of IUPC section.		

The health impact upon Aboriginal people has been considered, and where relevant incorporated and appropriately addressed in the development of this policy.

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