Manual on

Informed Consent

For

Gynaecologic & Obstetric Surgical Procedures



Association of Fellow Gynaecologist, Mumbai

Manual on Informed consent For Gynaecologic & Obstetric Surgical Procedures

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From Editor's Desk



Rajendra M Jain

Significance of Informed consent

It is important to understand the concept of "informed consent" and the "legal theories" upon which the law is based. The model consent forms must be sufficiently generic to be acceptable in most of jurisdictions. The readers are cautioned to thoroughly investigate their respective state laws that pertain to informed consent or consult with legal counsel or adviser before attempting to implement the forms or draft policies and procedures dealing with consent issues.

Informed consent is a legal doctrine that requires physician to obtain consent for treatment (diagnostic / therapeutic or medical / surgical, and invasive / noninvasive).

Without informed consent the doctor can be held liable for violating patient's rights regardless of whether the treatment was appropriate and rendered within the standard of care. The failure to obtain informed consent may result in the physician being accord to battery under common law. It is primary charge in approximately 2% cases of malpractice suits according to 2003 ACOG survey but it is secondary issue in nearly one third cases.

Informed consent is a continuous ongoing process that include exchange of information and development of choices therefore is not mere signing of a consent sheet but involve four basic components which are "voluntariness", "competence", "information" and "understanding". It is recommended that consent issues be addressed on the basis of the substance of the communication and the quality of the evidence.

Patient's autonomy and patient's moral right to self-body integrity and self-determination are to be respected but beneficence is also important ethical principle in informed consent. While physicians treating patients always keep in mind about "limited medical resources" available to community.

Healthcare professionals must merge the concepts of "communication and evidence" of consent so that when a challenge arises about an individual case the consent form itself will create a strong presumption that informed consent was obtained.

Informed consent is based on concept of freedom, which means "ability to choose and ability to refuse" treatment therefore recognition of different values, preferences, and alternative treatment options are also basic element of consent sheet.

There are some "practical limitations" such as time limitation, technical limitation, language barriers, communication ability of individual physician, patient's right, confidentiality some time create problems while making ethical decision.

There are "certain concerns" such as physician's financial interest, hospital treatment protocols, medical care plans, good practice guidelines while taking informed consent.

According to courts there are "three different degree of disclosure" in informed consent which is following:

- Reasonable physician standard
- Reasonable patient viewpoint standard
- Subjective patient viewpoint (individual 's personal need and particular requirement) this is rarely held by court

<u>Procedure of informed consent is suspended in following emergency situations:</u> (Implied Consent Rule Enables Provider to Act in an Emergency)

- Unconscious patient
- Life threatening conditions

Patient has right to refuse treatment: in such situation, the physician is supposed to document the reason given by the patient for refusal of the proposed treatment and also document possible adverse consequences to future health and wellbeing by refusal, and obtain signature of patient for refusal of treatment or procedure.

Informed consent is better to be obtained in office setting because time for adequate consideration and adequate discussion is provided, where presence of family member can also witness the discussion and sign the consent form as witness. The pamphlets or information leaflet, audiovisual, interactive visuals, greatly assist this whole process of taking informed consent because people usually retain 30% of verbal communication.

Informed consent usually covers six areas which are following:

- a. Diagnosis
- b. Nature and purpose of procedure
- c. Risk of procedure (general and specific)
- d. Likelihood of success of procedure
- e. Reasonable alternative options
- f. Consequence if procedure is refused and prognosis

Consent forms usually include words like:

- "request for surgery"
- "general risk for surgery"
- "specific risk of surgery"
- "may include but not limited to such complications"
- "reasonable alternatives of treatment such as"
- "Likelihood of success"

Well informed patients understand better about realistic outcome and medicine is not purely science therefore expectations are reasonable and disclosed complications are not cause of legal actions if it is not due to negligence.

Regulatory agencies statutes may require additional information or more details such as the date, patient's identity, names of the individuals, who will perform the procedure, specific authorization for anaesthesia, and disposition/disposal of any tissues removed.

"Anaesthesia consent forms" differ from surgical consent forms in that they do not contain the diagnosis, or the surgical or diagnostic procedures. However, in the first paragraph of consent form, the patient acknowledges that the surgical consent process occurred and that he/she understands the reason for anaesthesia. This acknowledgment by the patient is included to protect all parties and assure that appropriate informed consent took place.

Obtaining Informed Consent from Impaired Patients:

Among the circumstances that can diminish or impair a patient's capacity to understand the nature and risk of the proposed treatment, as well as alternatives to it are:

- The inability to speak or understand English
- The patient's physical condition adversely affects his/her capacity to decide
- Senility or another mental or emotional condition adversely affects the patient's capacity to comprehend
- Medication, alcohol or drugs prevent comprehension
- The patient must also have reached legal majority(which usually is 18 years old)

The following are commonly accepted substitutes:

- A parent (usually only one is necessary) for a minor child
- A husband or wife for a spouse

- A guardian for a ward
- Any adult standing in loco parentis (in place of the parent) for a minor; example, the principal of a boarding school.

Professional liability claims are increasing every year may be due to following factors:

- Fees are charge for services
- Reimbursement of doctor's fee by various medical care agencies or govt. programs
- Third party payer interventions
- Declining income and increasing overhead cost of hospitals / nursing home
- Longer working hours of doctors
- Sizable malpractice premium for indemnity insurances

Gynaecologists are sued mostly for following reasons:

- Failure of diagnosis such as cancer
- Higher qualification and more experienced gynaecologist are more likely to be sued because they are treating high risk patients more often so more adverse outcome possibility
- Medical Care given below the national standard of care(medical malpractice)
- Direct cause and effect relationship (poor outcomes)(maloccurence)

To minimize the risk of law suits, the approach to practice of medicine must have element of risk management which are following:

- Reduce the human errors
- Increase likelihood of desired results
- Documentation and providing medical record
- Investigate incident and adverse outcome in your setting
- Carryout targeted audit
- Having policies and protocols

At individual level following remedial measures to be advised:

- Adopt good surgical technique
- Appropriate knowledge of current developments & therapies
- Adequate documentation of events & procedure
- Good patient communication at offsetting & at indoor care
- Informed consent must for all procedure

This manual about informed consent contains some samples of informed consent sheet of commonest Gynaecologic and Obstetric surgical procedures, Refusal of Treatment form, and consent form for Autopsy of Stillbirth which may be helpful to readers but may be further customized according to individual need. REFERENCES:

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Section – II Consent form for Gynaecologic Surgery Procedures

Section – II Consent form for Gynaecologic Surgery Procedures Generic Informed Consent for surgery

•	Given name(s):	
	Date of birth/Age:	Sex: M /F
Condition and procedure: This condition requires the following proc	cedure. (Doctor to document in patient's ov	un words)
The following will be performed (Nature	& purpose of procedure):	
Risks of the procedure:		
	is procedure. They include but are not limite	ed to the following.
• Infection can occur, requiring antibio	otics and further treatment.	
Allergic reaction can occur from med	licines or by blood transfusion.	
 Bleeding could occur and may requir have been taking blood thinning drug 	re a return to the operating room. Bleeding ges.	is more common if you
 Small areas of the lung can collapse, physiotherapy. 	increasing the risk of chest infection. This m	nay need antibiotics and
 Increased risk in obese people of wo thrombosis. 	und infection, chest infection, heart and lun	g complications, and
Heart attack or stroke could occur du		
 Blood clot in the leg (DVT) causing pa to the lungs and produce damage to 	ain and swelling. In rare cases part of the clo organ	ot may break off and go
•	n or paralysis/paraplegia/quadriplegia.	
• Cardiac arrest & Death as a result of	this procedure is possible.	
Specific Risks:	ch may happened specifically with this type	of currency. They include
but are not limited to the following: (Doc	•	
Significant risks and procedure options:		
•		
The likelihood of success of above proce	aure is: Good /fair / poor	
Patient consent:		

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me
- Other relevant procedure/ treatment options and their associated risks
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.

- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About my procedure
 - Blood & Blood Products Transfusion

On the basis of the above statements, I request to have the procedure

- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	. Signature:	Date:

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent for Laparotomy

Family name:	Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
This condition requires the following procedure	. (Doctor to document in patient's own words)

The following procedure will be performed:

Surgical examination of the inside of the abdomen & the internal organs for any abnormality will be performed. This is done through a 15-30cm cut into the abdomen, depending on the size of the abdomen.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- Heavy bleeding in the abdomen. This may need fluid replacement, blood transfusion or further surgery. This may mean a longer stay in hospital and longer recovery time.
- Damage to other organs, such as bladder or bowel, which may need further surgery. This may mean a longer stay in hospital and longer recovery time.
- Infections such as pus in the abdomen. This may need surgical drainage and antibiotics.
- Bowel blockage after the operation. This may be temporary or in the longer term. Treatment may be a drip to give fluids into the vein and no food or fluids by mouth. If it doesn't get better, bowel surgery may be necessary which may include a colostomy. This can be temporary or permanent.
- Adhesions (bands of scar tissue) which can cause bowel obstruction. This can be a short term or long term complication and may need further surgery.
- The wound may not heal normally. The scar can be thickened and red and may be painful. This is permanent and can be disfiguring.
- Poor wound healing. The wound may burst open which may require long term wound care with
 dressings and antibiotics, or a hernia i.e. rupture can form in the long term. This may need repair by
 further surgery.
- There is increased risk in smokers of wound and chest infections, heart and lung complications and thrombosis.
- There is very low possibility of a fistula (a connecting passage between one area and another) developing.
- There is a possibility that the symptoms/pain you have been experiencing and the reason for this operation, may not resolve or worsen as a complication of the procedure.

Significant risks and procedure options: Risks of not having this procedure: Anaesthetic: The likelihood of success of above procedure is: Good /fair / poor. Patient consent: I request Dr.	 The cause of pain/other symptoms sometimes cannot be found, if you are having an exploratory operation. 	
Anaesthetic:	Significant risks and procedure options:	
The likelihood of success of above procedure is: Good /fair / poor. Patient consent: I request Dr		
Patient consent: request Dr		
I request Dr	The likelihood of success of above procedure is: Good /fair / poor.	
I request Dr	Patient consent:	
 Iacknowledge that the doctor has explained; My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me. The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me. Other relevant procedure/ treatment options and their associated risks. I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery. My prognosis and the risks of not having the procedure. That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care. The procedure may include a blood transfusion. Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital. If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan. A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training. I have been given the following Patient Information Sheet/s or explained verbally to me: About Your Anaesthetic Blood & Blood Products Transfusion About Laprotomy I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction. I understand that image/s or video footage my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor. I understand		
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Name of Patient:	to consider my decision.	
	On the basis of the above statements, I request to have the procedure.	
Circulations of Doubless	Name of Patient:	
Signature of Doctor:	Signature of Doctor: Signature of witness:	

<u>Section – II Consent form for Gynaecologic Surgery Procedures</u>

Informed Consent for Hysterectomy (Vaginal, Laparoscopic, Abdominal)

Family name:	Given name(s):
•	Date of birth/Age: Sex: M /F
Condition and procedure:	
The doctor has explained that you have the	e following condition: (Doctor to document in patient's own words)

The following procedure will be performed:

Removal of the uterus (womb) will be performed in the following way:

Vaginal (through the vagina)
 Laparoscopic ('key hole')
 Abdominal (through cut in abdomen)
 Ovaries will also be removed

Yes /No
Yes /No

If yes, which ovaries
 Left /Right/Both

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
 to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- The infection in the operation site or pelvis or urinary tract, requiring antibiotics and further treatment.
- The injury to other organs such as the ureter(s), bladder or bowel.
- A connection (i e fistula) may develop between the bladder and the vagina, or bowel or peritoneum.
- The bowel blockage after the operation.

Abdominal Hysterectomy:

- Bleeding into the abdominal wound from surrounding blood vessels.
- Poor wound healing.
- The wound scar may become thickened, red and painful.

Vaginal Hysterectomy:

- Risk of conversion to laparotomy (cut required in the abdomen).
- Higher risk of ureteric injury
- Recurrence of prolapse i.e. Vaginal repair may not be successful in the short or long term and may need further corrective surgery.
- Occurrence of pain during sexual intercourse or altered sexual function after vaginal repair.

Laparoscopic assisted Vaginal Hysterectomy:

• Risk of conversion to laparotomy (incision similar to Abdominal Hysterectomy).

- Change in bladder and bowel habits.
- Feelings of depression and anxiety.
- Onset menopause in pre-menopausal women if both ovaries are removed.

Significant risks and procedure options: Risks of not having this procedure: Anaesthetic:
Patient consent:
I request Dr perform upon me the above mentioned procedure.
I acknowledge that the doctor has explained;
 My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me. The anaesthetic required for this procedure. I understand the risks, including the risks that are specific
to me.
 Other relevant procedure/ treatment options and their associated risks.
 I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery. My prognosis and the risks of not having the procedure.
 That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
The procedure may include a blood transfusion.
 Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
• If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
 A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
 I have been given the following Patient Information Sheet/s or explained verbally to me:
About Your Anaesthetic and / or about Epidural & Spinal Anaesthesia
Blood & Blood Products Transfusion Abstract Heaterstein (Mariant Incompanies Abstracts)
About Hysterectomy (Vaginal, Laparoscopic, Abdominal)
• I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
• I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
 I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
• I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
• I certify and acknowledge that I have read this form or had it read to me; that I understand the risks,
alternatives and expected results of the this procedure and that I had ample time to ask questions and
to consider my decision.
On the basis of the above statements, I request to have the procedure

Signature of Doctor: Signature of witness:

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent for Laproscopy

Family name:	.Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
The doctor has explained that you have the following condi	tion: (Doctor to document in patient's own words)

The following will be performed:

A trocar will be put into the abdomen and instruments passed down the trocar to examine the inside of the abdomen and pelvis using a camera and video monitor. Sometimes, Adhesions, bands of fibrous tissue, which are most commonly due to previous abdominal or pelvic surgery or infections such as appendicitis or tubal infection grows around the bowel or other organs. If so, the doctor may need to cut these. The doctor may also need to operate on the pelvic organs.

Risks of this procedure

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific risks:

- Adhesions are difficult to diagnose with certainty without surgery and there is no alternative treatment for adhesions other than surgery. Adhesions can be the result of surgery and it cannot, therefore, be guaranteed that they won't come back after surgery to treat them. Sometimes a single band of tissue that is causing the bowel to twist or kink causes the problem. In that case, successful treatment is very high. Other cases, however, are caused by many filmy adhesions and these are much more difficult to get rid of.
- Accidental injury to the bowel, blood vessels and the urinary tract can occur during the passing of the laparoscope. If a complication happens during the surgery, then repair is usually possible at the time often through the small cuts. However, it may also be necessary to make a larger cut to repair the bowel, blood vessel or urinary tract injuries. In case of bowel injury, it may be necessary for the bowel to be brought out onto the abdomen. This allows waste to drain into a bag worn over the end of the bowel (known as a colostomy) so the injured bowel can heal. This colostomy would normally be closed at a separate operation.
- Heavy bleeding inside the abdomen. His may need fluid replacement, blood transfusion or further surgery. This may mean a longer than expected stay in hospital and longer recovery time.
- Damage to other organs, such as bladder or bowel, which may need further surgery. This may mean a longer than expected stay in hospital and longer recovery time.
- Rarely the gas, which is passed into the abdomen, can cause heart and chest complications.
- Infections such as pus collections in the abdominal cavity. This may need surgical drainage and antibiotics.
- Adhesions (bands of scar tissue) may form and cause a bowel obstruction. This can be a short term or a long term complication and may need further surgery.
- In some people, healing of the wound may be abnormal and the wound can be thickened and red and the scar may be painful.
- A weakness in the wound with the development of a hernia (rupture). This may need further surgery.
- There is increased risk in smokers of wound and chest infections, heart and lung complications and thrombosis.
- There is very low possibility of a fistula (a connecting passage between one area and another) developing.

- There is a possibility that the symptom(s)/pain you have been experiencing and the reason for this operation, may not resolve or worsen as a complication of the procedure.
- The cause of pain/other symptoms sometimes cannot be found, if you are having an exploratory operation.

Significant risks and procedure options:	
Risks of not having this procedure:	
Patient consent:	
I request Dr	to perform upon me the above mentioned procedure.

I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About your Anaesthetic.
 - About Laparoscopy
 - o Blood & Blood Products Transfusion
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements,	I request to have t	the procedure

Name of Patient:	Signature:	Date:
	3 3 3 3	
Signature of Doctor:	Signatı	ure of witness:

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent form for Burch Colpo-Suspension

•	Given name(s):
Condition and procedure:	procedure. ((Doctor to document in patient's own words)
The following will be performed: The cut is made across the upper edi	ge of the pubic hair. This allows the surgeon to get to the bladder neck.

The cut is made across the upper edge of the pubic hair. This allows the surgeon to get to the bladder neck. Stitches are placed in tissues next to the bladder neck. This is to hang it from ligaments on the front of the pelvic bone. These stitches will support the bladder neck. This has a very high success rate (better than 9 in 10 women) of treating genuine stress Incontinence. This falls to a high success rate in the coming years. If required, this is followed by looking into the bladder to make sure no damage has been done (cystoscopy).

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific risks:

- The bladder may be over-active after the operation. You may need to go to the toilet a lot, may have sudden urge to pass urine and may leak urine. These symptoms may be controlled by bladder retraining and drug therapy. The drug therapy is then slowly cut back. Injury to urinary bladder/ ureter or the urethra may require further surgery.
- A rupture (hernia) through the top of the vagina. Sometimes further surgery is needed if the hernia becomes large enough.
- Haemorrhage from large arteries and veins about the bladder, vagina and pelvis. This may require a blood transfusion and further surgery.
- Infection in the operation site, pelvis or urinary tract. This may require treatment with antibiotics.
- Problems with passing urine. This is rare, but may need long term care. If this does happen, you may have to pass a tube (catheter) into your bladder to drain the urine.
- There is a higher risk in smokers. This may cause wound and chest infections, heart and lung problems and blood clots in the veins.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:
the Bhalthand of access of above accessions in Cond. I fair I was a

The likelihood of success of above procedure is: Good / fair / poor

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I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - Blood & Blood Products Transfusion
 - About Burch Colpo-suspension
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	: Signature:	Date:
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Signature of Doctor:	Signatu	re of witness:
- 0	- 0	

<u>Section – II Consent form for Gynaecologic Surgery Procedures</u>

Informed Consent form for Excision of a Bartholins Gland

Famil	ly name:	Given name(s):
Addre	ess:	Date of birth/Age: Sex: M /F
Condi	ition and procedure:	
		((Doctor to document in patient's own words)
	ollowing procedure will be performed:	a of a Double distance or a base of
	procedure is a surgical incision and drainage ral risks of an excision of a Bartholins glan	•
	Infection can occur, requiring antibiotics an	
	Allergic reaction can occur from medicines	•
	have been taking blood thinning drugs.	urn to the operating room. Bleeding is more common if you
	Small areas of the lung can collapse, increase ohysiotherapy.	sing the risk of chest infection. This may need antibiotics and
		ection, chest infection, heart and lung complications, and
	thrombosis.	contony chest infection, heart and rang complications, and
-	Heart attack or stroke could occur due to th	ne strain on the heart.
• E		swelling. In rare cases part of the clot may break off and go
	Lose of function of any limb or organ or par	alvsis/paraplegia/quadriplegia.
	Cardiac arrest & Death as a result of this pr	
	fic Risks:	
•		not return to normal in approximately 20% of cases.
	Recurrence of abscess or cyst is common ar	
	•	ome time. This may require long term wound dressings.
		l and chest infections, heart and lung complications and
	thrombosis.	
Signif	ficant risks and procedure options:	
Risks	of not having this procedure:	
Anaes	sthetic:	
The li	ikelihood of success of above procedure is	: Good / fair / poor
	nt consent:	
		. to perform upon me the above mentioned procedure.
I ackn	nowledge that the doctor has explained;	

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.

- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - Excision of a Bartholins Gland
 - Blood & Blood Products Transfusion
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given has to the results. Understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	Signature:	Date:
Signature of Doctor:		. Signature of witness:

<u>Section – II Consent form for Gynaecologic Surgery Procedures</u> Informed Consent for Cone biopsy of cervix

Family name:						
Address:	Date of birth/Age:	Sex: M /F				
Condition and procedure: This condition requires the following procedure. ((Doctor to document in patient's own words).						
The following procedure will be performed:						
The end of the cervix is surgically removed. The end conditions in the area and further treatment is then g		nologist for any unusual				
Risks of the procedure: There are risks and complications with this procedure.	a. Thou include but are not limit	ad to the following				
General risks:	e. They include but are not innit	ed to the following.				
 Infection can occur, requiring antibiotics and fu 	rther treatment					
 Allergic reaction can occur from medicines or b 						
Bleeding could occur and may require a return have been taking blood thinning drugs.	•	is more common if you				
 Small areas of the lung can collapse, increasing physiotherapy. 	the risk of chest infection. This n	nay need antibiotics and				
 Increased risk in obese people of wound infecti thrombosis. 	on, chest infection, heart and lui	ng complications, and				
 Heart attack or stroke could occur due to the st 						
Blood clot in the leg (DVT) causing pain and sweets to the lungs and produce demand to a great the control of the lungs and produce demand to a great the control of the lungs and the lungs are the control of the lungs and the lungs are the lun	elling. In rare cases part of the clo	ot may break off and go				
to the lungs and produce damage to organ	ic/paraplagia/guadriplagia					
Lose of function of any limb or organ or paralysCardiac arrest & Death as a result of this proces						
Specific Risks:	idie is possible.					
 Damage and narrowing of the cervix may occur 	which can cause painful periods	and difficulty in Jahour				
 The cervix may not be competent and cause pro 		· ·				
 Haemorrhage from the cervix, which may need within weeks of the procedure. 						
 Infection may be introduced into the uterus or antibiotics. 	tubes or abdomen. This may req	uire treatment with				
 There is increased risk in obese people of woun and thrombosis. 	d infection, chest infection, hear	t and lung complications				
 There is increased risk in smokers of wound and thrombosis. 	d chest infections, heart and lung	complications and				
Significant risks and procedure options:						
Risks of not having this procedure:						
Anaesthetic:						
The likelihood of success of above procedure is: Go						

Patient consent:

I request Dr. to perform upon me the above mention procedure. I acknowledge that the doctor has explained;

• My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.

- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - o Blood & Blood Products Transfusion
 - About Cone Biopsy
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	Signature:	Date:
Signature of Doctor:	Signature o	of witness:

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent for Dilatation & Curettage

Family name:	Given name(s):					
	Date of birth/Age: Sex: M /F					
Condition and procedure:						
This condition requires the following procedure. ((Doctor to document in patient's own words)						
The following procedure will be per	rformed:					
After widening / dilating the cervix (the neck of the womb), an instrument attached to suction is passed into					
the uterus (womb). The lining of the	e uterus and any other tissue that looks abnormal inside the uterus is					
hen removed and may be sent to pathology for tests.						

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
 to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- Damage or tearing of the cervix. This may need repair. It can also possibly lead to early pregnancy loss in future pregnancies.
- Damage to the uterus due to a perforation (puncture) and possible bowel damage. This may need more surgery and a longer hospital stay than expected.
- Severe bleeding (haemorrhage) from the uterus. This may need a blood transfusion.
- Infection in the uterus and tubes. This may need antibiotics.
- The tissue inside the uterus may not all be removed. This may need further surgery.
- Rarely air may get into the blood stream. This air embolism can cause the heart to stop. This can be fatal.
- After the procedure is performed, there may be bleeding for up to 10 to 14 days.
- Your first period after the procedure may be late. It may be longer or shorter than usual. There may be more or less than the usual amount of blood loss.
- There is higher risk in smokers of chest infection, heart and lung problems and blood clots in the veins.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:
The likelihood of success of above procedure is: Good / fair / poor

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I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - o About D & C
 - Blood & Blood Products Transfusion
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	: Signature:	Date:
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Signature of Doctor:	Signatuı	re of witness:

<u>Section – II Consent form for Gynaecologic Surgery Procedures</u> Informed Concent for Diagnostic hystoroscopy and D. &.

	informed Consent for Diagnostic hysteroscopy and D & C
Fami	ily name:Given name(s):
Addr	ess: Date of birth/Age: Sex: M /F
Cond	lition and procedure:
	condition requires the following procedure. ((Doctor to document in patient's own words)
	following procedure will be performed:
wom	er an anaesthesia, the cervix is carefully dilated until there is enough room to pass a telescope into the lib. The womb is then filled with fluid/gas, which gives a better view of the inside. A telescope is used to
usua	f there is anything abnormal inside the womb. The fluid is then drained out. The lining of the womb is lly scraped to collect tissues. The lining of the uterus and any other tissue that looks abnormal inside the us is then removed and may be sent to pathology for examination.
	s of the procedure:
Ther	e are risks and complications with this procedure. They include but are not limited to the following.
Gene	eral risks:
•	Infection can occur, requiring antibiotics and further treatment.
•	Allergic reaction can occur from medicines or by blood transfusion.
	Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
	Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
	Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
•	Heart attack or stroke could occur due to the strain on the heart.
	Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
	Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
	Cardiac arrest & Death as a result of this procedure is possible.
-	ific risks:
	Bleeding that can be so heavy that a blood transfusion may be needed. It may also need further surgery. Damage may occur to the uterus with rupture or perforation. This may require a laparoscopy and/or

- laparotomy, and/or longer stay in hospital than expected. In the event of uterine perforation, there is a risk of damage to other organs, such as bowel or bladder, which may require further corrective surgery.
- Rarely, the procedure may not be able to be completed, due to narrowing of the inside of the cervix. If the condition continues, further surgery will be necessary.
- Infection can occur in the uterus. This can cause heavy bleeding or discharge, worsening cramps or high fevers. The infection may affect the fallopian tubes and cause problems with getting pregnant in the future. Antibiotics are used to treat the infection.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:
The likelihood of success of above procedure is: Good / fair / poor

Patient consen	

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About Diagnostic Hysteroscopy & D and C
 - o Blood & Blood Products Transfusion
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	Signature:	Date:	
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Signature of Doctor:	Signature	e of witness:	

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent for Endometrial Resection/ Ablation

illiornied Consent	of Endometrial Resection, Abiation
Family name:	Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
This condition requires the following proced	dure. ((Doctor to document in patient's own words)
The following procedure will be performed	:
The cervix is dilated and an instrument is pa	issed through the cervix into the uterus. The lining of the uterus
is then removed using electric current (diatl	nermy) or other methods.
Risks of the procedure:	
There are risks and complications with this p	procedure. They include but are not limited to the following.

- General risks:Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people & smoker of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific risks:

- The procedure may not be able to be completed, due to narrowing of the cervical canal or problems inside the uterus. This may need further surgery or surgery abandoned.
- Damage to the uterus with perforation by the diathermy instrument, can cause bleeding and may need further surgery to repair the damage. Injuries may not be recognised at time of surgery.
- Damage, by burning, to bowel or bladder. This will need further surgery and a longer than expected stay in hospital. Adhesions may result and a colostomy may be needed. Injuries may not be recognised at time of surgery.
- Infection could be introduced into the uterus or tubes or abdominal cavity. This may need treatment with antibiotics.
- The excessive bleeding (Haemorrhage) from the uterus or blood vessel can occur. A catheter may be passed into the uterus to provide balloon pressure to the wall of the uterus for a few hours to control bleeding. Blood transfusion, further surgery and possibly hysterectomy may be necessary if the bleeding doesn't stop.
- The distending fluid used to stretch the uterus can be absorbed causing coma or death. Both are extremely rare.
- There is risk of failure of procedure in short or long term
- Possibility of pain, and/or bleeding which may get worse after the procedure, and may be long term.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:
The likelihood of success of above procedure is: Good / fair / poor

Patient consent:

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - o Blood & Blood Products Transfusion
 - About Endometrial Resection/ Ablation
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	Signature:	Date:	
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Signature of Doctor:	Signatura	of witness:	
Signature of Doctor		DI WILIIESS	

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent for LLETZ of the Cervix

Family name:	
Condition and procedure: This condition requires the following procedure. (Doctor to document in patient's own words)	
The following procedure will be performed: A large loop excision of the transformation zone (LLETZ) is where a small piece of the cervix is cauterised (burnt) with an electric current.	
 Risks of the procedure: There are risks and complications with this procedure. They include but are not limited to the following. General risks: Infection can occur, requiring antibiotics and further treatment. Allergic reaction can occur from medicines or by blood transfusion. Bleeding could occur and may require a return to the operating room. Bleeding is more common if have been taking blood thinning drugs. Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics physiotherapy. Increased risk in obese people of wound infection, chest infection, heart and lung complications, are thrombosis. Heart attack or stroke could occur due to the strain on the heart. Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and to the lungs and produce damage to organ Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia. Cardiac arrest & Death as a result of this procedure is possible. 	you s and nd
 Specific Risks: Excessive bleeding from the cervix, which may need blood transfusion or further surgery, either init or within weeks of the procedure. Future pregnancies, usually around 20 weeks of pregnancy, may suffer from weakness of the cervix support a growing pregnancy. This is an uncommon risk. Infection may be introduced into the cervix, uterus, tubes or abdomen. This may require treatment antibiotics. Damage and narrowing of the cervix could occur which can cause painful periods and difficulty in labour. There is increased risk in smokers of wound and chest infections, heart and lung complications and thrombosis. Significant risks and procedure options:	to with

Anaesthetic:

The likelihood of success of above procedure is: Good / fair / poor

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I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - o About Your Anaesthetic
 - Blood & Blood Products Transfusion
 - o About LLETZ of the Cervix
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Name of Patient:	Signature:	Date:	
	G		
C'anal and David	61	•	
Signature of Doctor:	Signature of	witness:	.

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent for Ovarian Cystectomy/ Oophorectomy

•	procedure. ((Doctor to document in patient 5 0 mi moras)
Condition and procedure: This condition requires the follow	ving procedure. ((Doctor to document in patient's own words)
	Date of birth/Age: Sex: M /F
•	Given name(s):

- Laparotomy
- Laparoscopy
- o Removal of ovary (include side).....

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
 to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- Severe bleeding can occur therefore blood transfusion may be required to replace blood loss.
- Infection in the operation site or pelvis or urinary tract can occur which may be treated by wound dressings and / or antibiotics.
- Injury to other organs such as the ureter(s) (tube leading from kidney to bladder) bladder or bowel may occur. This is an uncommon risk. Further surgery will be needed to repair the injuries. For bladder injuries, a catheter (plastic tube) may be put into the bladder to drain the urine away until the bladder is healed. For ureteric injury, a plastic tube (stent) is placed in the ureter for 6 weeks and then removed by cystoscopy. If the bowel is injured, bowel resection and a possibility of a temporary or permanent colostomy. A damaged kidney may require removal.
- Bowel blockage after the operation. This may be temporary or in the longer term. Initial treatment may be a drip to give fluids into the vein and no food or fluids by mouth. If it doesn't get better, bowel surgery may be necessary which may include a colostomy. This can be temporary or permanent.
- The poor early wound healing and the wound may burst open which may require long term wound care with dressings and antibiotics, or a hernia can form in the long term which may need repair by further surgery.
- Wound may heal normally with a thickened scar which may be red and painful. This is called a "keloid" and may be permanent and can be disfiguring.
- Sometimes a small part of the ovary may be left behind and could cause further problems, like pain. & occurrence of other cysts, which may need future surgery.
- Adhesions may form at the site of cyst removal.

consider my decision.

- If both ovaries are removed before onset of menopause, risk of osteoporosis and may require therapy eg. hormone replacement therapy.
- Need for oophorectomy when consented for a cystectomy if bleeding excessive or ovary badly damaged.
- There is increased risk in smokers of wound and chest infections, heart and lung complications and thrombosis.

Significant risks and procedure options: Risks of not having this procedure: Anaesthetic: The likelihood of success of above procedure is: Good / fair / poor
Patient consent:
I request Dr to perform upon me the above mentioned procedure.
I acknowledge that the doctor has explained;
My medical condition and the proposed procedure, including additional treatment if the doctor finds
something unexpected. I understand the risks, including the risks that are specific to me.
• The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
Other relevant procedure/ treatment options and their associated risks.
 I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery. My prognosis and the risks of not having the procedure.
• That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
The procedure may include a blood transfusion.
• Tissues and blood may be removed and could be used for diagnosis or management of my condition,
stored and disposed of sensitively by the hospital.
• If immediate life-threatening events happen during the procedure, they will be treated based on my
discussions with the doctor or my Acute Resuscitation Plan.
A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor
undergoing further training.
 I have been given the following Patient Information Sheet/s or explained verbally to me:
About Your Anaesthetic
o Blood & Blood Products Transfusion
 Information leaflet about Ovarian Cystectomy/ Oophorectomy
I was able to ask questions and raise concerns with the doctor about my condition, the proposed
procedure and its risks, and my treatment options. My questions and concerns have been discussed and
answered to my satisfaction.
• I understand I have the right to change my mind at any time, including after I have signed this form but,
preferably following a discussion with my doctor.
 I understand that image/s or video footage may be recorded as part of and during my procedure and that
these image/s or video/s will assist the doctor to provide appropriate treatment.
I accept that medicine is not an exact science and understand that no guarantees can be given to the
results and understanding these limitations.
• I certify and acknowledge that I have read this form or had it read to me; that I understand the risks,
alternatives and expected results of the this procedure and that I had ample time to ask questions and to

Signature of Doctor: Signature of witness:

<u>Section – II Consent form for Gynaecologic Surgery Procedures</u>

Informed Consent for Insertion of Tension Free sub or mid urethral Vaginal / Trans- obturator Tape with / without Cystoscopy

Family name:	Given name(s):
Address:	
Condition and procedure:	
This condition requires the following procedure. (Docto	or to document in patient's own words)
	, , , , , , , , , , , , , , , , , , ,

The following procedure will be performed:

The damaged ligaments are replaced by a 1cm wide tape of synthetic mesh. This tape returns the support for the urethra to the surrounding tissues. This is routinely followed by looking into the bladder to make sure no damage has been done (cystoscopy).

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

The success rate is very high (approximately 90%). The long-term success rate is not yet known.

- The bladder may be over-active after the operation. You may need to go to the toilet a lot, may have sudden urges to pass urine and may leak urine. These symptoms are usually managed by bladder retraining and drug therapy. A small proportion of patients will continue to have long standing bladder symptoms despite treatment.
- Problems with passing urine (retention) are uncommon. This rarely needs long term management. If this happens, the tape may be divided through the vaginal cut. There is a small risk of the urinary incontinence returning.
- Infection requiring antibiotics and further treatment.
- Excessive bleeding is very rare.
- There is increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis
- There is a higher risk in smokers. This may cause wound and chest infections, heart and lung problems and blood clots in the veins.
- The urethra or the bladder can sometimes be damaged & require repair.
- The tape may erode through the urethra in the years after the operation. This would need repair of the urethra and a urethral catheter for 2 weeks.

Circuitionat viales and averaged up autions.	
Significant risks and procedure options:	
Anaesthetic:	
The likelihood of success of above procedure is: Good / fair / poor	
Patient consent:	
I request Dr to perform up	on me the above said procedure.
I acknowledge that the doctor has explained;	
My medical condition and the proposed procedure, including additions and the proposed procedure, including the risks the proposed procedure.	
something unexpected. I understand the risks, including the risks tha	-
The anaesthetic required for this procedure. I understand the risks, in to me.	icidaing the risks that are specific
to me. Other relevant precedure/treatment entions and their associated rich	-kc
 Other relevant procedure/ treatment options and their associated ris I consent to tackle/operate/remove any additional abnormal patholo 	
 My prognosis and the risks of not having the procedure. 	ngy encountered during surgery.
 That no guarantee has been made that the procedure will improve m 	ny condition even though it has
been carried out with due professional care.	ny contantion even though it has
The procedure may include a blood transfusion.	
Tissues and blood may be removed and could be used for diagnosis of the could be used for diagnosis.	or management of my condition,
stored and disposed of sensitively by the hospital.	,
• If immediate life-threatening events happen during the procedure, the	ney will be treated based on my
discussions with the doctor or my Acute Resuscitation Plan.	
A doctor other than the Consultant may conduct the procedure. I und	derstand this could be a doctor
undergoing further training.	
 I have been given the following Patient Information Sheet/s or explain 	ned verbally to me:
About Your Anaesthetic	
Blood & Blood Products Transfusion	
 About Insertion of Tension Free sub or mid urethral Vaginal Tape 	
I was able to ask questions and raise concerns with the doctor about	· · · · · · · · · · · · · · · · · · ·
procedure and its risks, and my treatment options. My questions and	concerns have been discussed and
answered to my satisfaction.	
I understand I have the right to change my mind at any time, including a discussion with my destar.	ig after i have signed this form but,
preferably following a discussion with my doctor.	of and during my procedure and
 I understand that image/s or video footage may be recorded as part of that these image/s or video/s will assist the doctor to provide appropriate 	- · ·
 I accept that medicine is not an exact science and understand that no 	
results and understanding these limitations.	guarantees can be given to the
 I certify and acknowledge that I have read this form or had it read to 	me: that Lunderstand the risks
alternatives and expected results of the this procedure and that I had	
to consider my decision.	dumple time to ask questions and
On the basis of the above statements, I request to have the procedure	
Name of Patient: Signature:	Date:
Signature of Doctors	ure of witness:
Signature of Doctor: Signature	ire or withess:

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent form for Electric or Cryo cautery to cervix

Family name:	Given name(s):
•	Date of birth/Age: Sex: M /F
Condition and procedure:	, ,
,	dure. ((Doctor to document in patient's own words)
The fellowing was adversarill be weatherness	

The following procedure will be performed:

The mouth of the cervix having lesion is cauterised with an electrical current/cryo-cautery.

Risks of a electric cautery or cryo cautery to cervix:

There are risks and complications with this procedure. They include but are not limited to the following.

General risk:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese & smoker people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- Damage & narrowing of the cervix could occur which can cause painful periods and difficulty in labour.
- In future pregnancies, there is a risk of losing the baby between 16 and 20 weeks of pregnancy. This is an uncommon risk.
- Haemorrhage from the cervix, which may need blood transfusion or further surgery, either initially or within weeks of the procedure.
- There is increased vaginal discharge following cautery, may require some medications & vaginal douching.

Significant risks and procedure options:							
·							
Anaesthetic:							
The likelihood of success of above procedure is:	Good / fair / poor						
Patient consent:							
I request Dr	to perform upon me the above mentioned procedure.						
I acknowledge that the doctor has explained;							

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.

- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About Electric or cryo Cautery of cervix
 - o Blood & Blood Products Transfusion
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I i	request to	have the	procedure
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Name of Patient:	Signature:	Date:	
	8 3 3 3		
Signature of Doctor:	Signature of v	vitness:	

Section – II Consent form for Gynaecologic Surgery Procedures

Informed cons	ent for Vaginal Hysterectomy & Repair	
Family name:	Given name(s):	
Address:	Date of birth/Age:	Sex: M /F
Condition and procedure: This condition requires the following procedure:	rocedure. (<i>(Doctor to document in patient's own w</i> o	ords)
The following procedure will be perform	rmed:	•••••
- •	n the vagina and repair of any utero- vaginal prolaps	se.
 Ovaries will also be removed 		
	Left /Right / Both	
Risks of the procedure:	, 6 .	
There are risks and complications with	this procedure. They include but are not limited to	the following.
General risks:		
 Infection can occur, requiring anti- 	biotics and further treatment.	
Allergic reaction can occur from m	nedicines or by blood transfusion.	
	uire a return to the operating room. Bleeding is mo	re common if you
have been taking blood thinning d	_	
	se, increasing the risk of chest infection. This may n	eed antibiotics and
physiotherapy.		
	s people of wound infection, chest infection, heart a	and lung
complications, and thrombosis.	al a ta tha atasta a a tha haa a	
Heart attack or stroke could occur Plead slat in the log (D)/T) causing		wheat off and an
	pain and swelling. In rare cases part of the clot ma	y break on and go
to the lungs and produce damage	to organ gan or paralysis/paraplegia/quadriplegia.	
Lose of function of any limb or orgCardiac arrest & Death as a result		
Specific risks:	or this procedure is possible.	
-	m large blood vessels about the uterus.	
 Collection of blood clot at top of v 	_	
•	ration site or pelvis or urinary tract requiring antibi	otics and further
treatment.	ration site of pervis of utiliary tract requiring antibio	otics and further
	s the ureter(s) (tube leading from kidney to bladder	·) bladder or bowel.
The repair of these organs may be	, , ,	,
	vagina and other organs (bladder, bowel).	
 Bowel blockage can occur after the 		
 Pain in the perineum, which can la 	•	
•	nal repair may not be successful, in the short or lon	ig term and may
	results in recurrence of the prolapse.	,
	during sexual intercourse or altered sexual function	after vaginal
repair surgery.	6	
 Change in bladder and bowel habi 	ts can occur occasionally.	
Onset of menopause in pre- meno	pausal women if both ovaries are removed.	
	s:	
The likelihood of success of above pro	cedure is: Good / fair / poor	

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I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - o About Your Anaesthetic (Epidural & Spinal Anaesthesia)
 - o Blood & Blood Products Transfusion
 - About Vaginal Hysterectomy with repair
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given has to the results. Understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:	
	G		
Signature of Doctor:	Sig	gnature of witness:	

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent for Vaginal Repair

Family name:	Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
This condition requires the following proc	edure. (Doctor to document in patient's own words)

The following procedure will be performed:

Repair of any prolapse through the vagina. This involves a cut in the vaginal area to repair the prolapse of the bladder and / or the rectum (lower bowel) and/or vaginal entrance. It may be necessary to pass a catheter into the bladder after the operation to drain the urine until healing has taken place. Use of mesh or special stich to Sacro-Spinous ligament may be required for vaginal repair.

Risks of a vaginal repair along with mesh and/or with Sacrospinous colpopexy:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific risks:

- Infection in the operation site or urinary tract requires antibiotics and further treatment.
- Injury to other organs such as the ureter(s) (tube leading from kidney to bladder) bladder or bowel.
- Difficulty passing urine immediately following surgery which is usually temporary but which may require a catheter to be reinserted into the bladder, or you may be taught to pass your own catheter until you are able to pass urine without assistance.
- Stress incontinence of urine following surgery. Stress incontinence is a common condition where urine leaks when you cough, sneeze or perform various other activities involving abdominal straining. In this case, whilst no problem existed before surgery, often there is an unknown weakness of the bladder which leads to this problem when surgery is carried out.
- A connection (fistula) may develop between the bladder and the vagina.
- A connection (fistula) may develop between the rectum and the vagina leading to leakage of faeces through the vagina (recto vaginal fistula).
- Pain in the perineum(area between vagina and rectum), which can last up to six weeks after surgery.
- Change in bladder and bowel habits.
- Narrowing or shortening of the vagina and pain during sexual intercourse.
- Reoccurrence of the original complaint (prolapse) with the passage of time.
- There is increased risk in smokers of wound and chest infections, heart and lung complications and thrombosis.
- Mesh in the form of a permanent or semi permanent artificial support may be used in vaginal repairs to support tissues in this area that are weak. Specific complications with the use of "mesh" are:
 - Infection this may require removal of the mesh.

- Mesh protrusion or erosion part of the mesh wears through a gap that develops in the vaginal skin so that it pokes out. This will usually require surgery to trim the loose portion and to close the gap in the skin.
- Rejection loosening of the whole mesh. The mesh may partially or completely protrude through the vaginal skin causing discharge or pain to your partner during sex. This may need another operation to divide the mesh that is protruding out or to remove the entire mesh if there is infection present.

Significant risks and procedure option	ons:
Risks of not having this procedure: .	
The likelihood of success of above p	procedure is: Good / fair / poor
Patient consent:	

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor/ my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - o About Your Anaesthetic (Epidural & Spinal Anaesthesia)
 - o Blood & Blood Products Transfusion
 - About Vaginal repair anterior colporraphy
 - o About Vaginal repair posterior colpo-perineorraphy
 - About Vaginal Repair with Mesh
 - About Vaginal Repair- with Sacrospinous Colpopexy
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure				
	Signature:	Date:		
Signature of Doctor:	Signatuı	re of witness:		

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent for Removal of Genital Warts

	ng procedure. (<i>(Doctor to document in patient's own v</i>	
The following procedure will be pe	erformed: wths on the genitals caused by a viral skin disease. Th	
Risks of removal of genital warts: There are risks and complications wards. Infection can occur, requiring an allergic reaction can occur from Bleeding could occur and may have been taking blood thinning. Small areas of the lung can coll physiotherapy. Increased risk in obese people thrombosis. Heart attack or stroke could occur and may have been taking blood thinning. Increased risk in obese people thrombosis. Blood clot in the leg (DVT) cause to the lungs and produce damage. Lose of function of any limb or	with this procedure. They include but are not limited to antibiotics and further treatment. In medicines or by blood transfusion. In require a return to the operating room. Bleeding is ming drugs. Illapse, increasing the risk of chest infection. This may of wound infection, chest infection, heart and lung contact the strain on the heart. In sing pain and swelling. In rare cases part of the clot meant and swelling.	nore common if you need antibiotics and complications, and
 further warts. The area where the wart was r The area of the wart may be the theorem of the wart may be the wart was represented in the wart may be the war	currence) and they are usually due to a virus infection removed is an open wound and will take time to heal hickened and some discolouring and pain in the scar. e people of wound infection, chest infection, heart ares. errs of wound and chest infections, heart and lung contions:	nd lung mplications and

Anaesthetic:

The likelihood of success of above procedure is: Good / fair / poor

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I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - Blood & Blood Products Transfusion
 - o About Removal of Genital Warts
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:
	0 1 11	
_		_
Signature of Doctor:	Signatur	re of witness:

Section – II Consent form for Gynaecologic Surgery Procedures Informed consent for Insertion of Intra Uterine Device (Cu-T / Progresterone Releasing)

The doctor has explained that you have the following condition:(Doctor to document in patient's own words)

The following procedure will be performed:

A progesterone releasing intra-uterine device or Cu-T IUD will be put inside the uterus through the vagina. This will provide long term birth control.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you
 have been taking blood thinning drugs
- Death as a result of this procedure is possible, but very rare.

Specific risks:

- The infection can be passed into the uterus and spread into the pelvis. This may need treatment with antibiotics, and can cause infertility (unable to get pregnant).
- Abnormal bleeding in the first three months may occur. This usually settles its own or some medication can be given to stop bleeding. If not the device may have to be removed.
- Puncture of the wall of the uterus when the intrauterine device is put inside. The device will not work, so pregnancy is possible. There may be infection, which will need antibiotics. The device may need to be removed and a Laparo-Hystroscopy may be necessary to do this.
- The uterus may push out the intra-uterine device in 1 in 20 women. If this happens, pregnancy is a possibility.
- Pain may be felt during and shortly after insertion.
- Ovarian cysts in 1 in 8 women, which may cause pain and painful sex. They usually disappear in 2-3 months. If they persist, further monitoring is required.
- The intra-uterine device may not provide complete relief from the symptoms for which it is being used.
- Pregnancy in 1 in 600 women. A third of these will have an ectopic (tubal) pregnancy. In which case, surgery will be required to remove the foetus and possibly the tube to prevent the tube from rupturing.
 If the tube ruptures, it can be life threatening and emergency surgery will be required.

Significant risks and procedure options:	
Risks of not having this procedure:	
Anaesthetic:	
The likelihood of success of above procedure is: Good / fair / poor	

Patient consent:

I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About insertion of IUD
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:
	G	
Signature of Doctor:	Signature	of witness:

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent form for Female Sterilisation

Family name:	Given name(s):	
Address:	Date of birth/Age:	Sex: M /F
Condition and procedure:		
The doctor has explained that you have the fo	ollowing condition: (Doctor to document in patie	nt's own words)

The following procedure will be performed:

The operation is usually done Laparoscopically – which is commonly known as keyhole surgery. You will be given anaesthesia during the operation which may be a general anaesthetic given by needle into a vein or under spinal anaesthesia given in the back in spinal cord or under local anaesthesia with sedation. One or two cuts will be made into your abdominal wall and a gas/air piped into the abdomen in order to lift up the abdominal wall. A telescope will be put through one of the cuts and sterilising instrument through another. The cuts will be closed, usually with a dissolvable stitch or sticky tape. Most women go home on the day of surgery.

Sometime this operation is carried out through small incision in abdomen known as mini-laprotomy.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risk:**

- Infection can occur, requiring antibiotics and further treatment.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs.
- Rarely death as a result of this procedure is possible.

Specific risks:

- This sterilisation operation is intended to make you sterile. You should not have the operation if you are uncertain about whether you will want children in the future. It should be assumed that this operation cannot be reversed. In some cases, it is possible to re-open the tubes by micro-surgery. Discuss the success rate with your doctor. All contraceptive techniques, including sterilisation, have a failure rate. Pregnancies have even been reported after hysterectomy (removal of the womb). If the tubes are cut, the removed pieces of tubes will be examined under the microscope to prove sterilisation. There are risks and complications with this procedure. They include but are not limited to the following.
- Accidental injury to the bowel, blood vessels and the urinary tract. Repair is usually possible at the time

 often through the small cuts. It may also be necessary to make a larger cut to repair the bowel, blood vessel or urinary tract injuries.
- In case of bowel injury, it may be necessary for a temporary colostomy to allow the injured bowel to heal. This colostomy would normally be closed at a separate operation a few weeks later.
- Rarely gas, used to inflate the abdomen, can cause heart and breathing problems in 1 in 60,000 women. Death is a very rare risk.
- About future pregnancy: The failure rate of the two commonest laparoscopic sterilisations (Filshie clip and Fallope ring) is about 1 in 170 to 1 in 250 women who will become pregnant after female

- sterilisation. Pregnancy may also happen outside the womb (ectopic pregnancy) and may require emergency surgery. This is rare.
- If the operation cannot be completed through the laparoscope, then open surgery may have to be done. This will mean a larger cut above the pubis about 5-8 cm, a longer stay in hospital and a longer recovery rate.
- Burns on the skin due to use of electrical equipment in less than 1 in 100 women. These may take a few days to appear.

Significant risks and procedure options:	
Risks of not having this procedure:	
Anaesthetic:	
The likelihood of success of above proce	
Patient consent:	
I request Dr	to perform upon me the above mentioned procedure
I acknowledge that the doctor has explain	ned;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About Female Sterilisation
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

to consider my decision.		
On the basis of the above statements	, I request to have the procedure	
Name of Patient:	Signature:	Date:
Signature of Doctor:	Signature	e of witness:

MUNICIPAL CORPORATION GREATER MUMBAI (MCGM) STATUARY REQUISITE FOR STERILISATION OPERATION

Annexure-1

CHECKLIST TO BE FILLED BEFORE STERILISATION OPERATION OF A MALE OR FEMALE BY THE DOCTOR CONCERNED

01112	REIST TO BE FIELD BEFORE STERILISATION OF ERATION OF A WALL OR FLIMALE BY THE DOCTOR	· contermit
1	Whether the age of the client is within laid down norms (Male clients should be below the age	Yes/No
	of 60 years, for female should be below the age of 45 yrs and above 22 yrs)	
2	whether information relating to marital status, No. of living children and age of the youngest	Yes/No
	child obtained	
3	Whether the client has been counseled regarding sterilization so as to help the clients make	Yes/No
	informed and voluntary decision	
4	Consent form, Whether the client has understood the consent for and following relative	Yes/No
	contraindications	
5	Whether the client has been examined for excluding medical contraindication i e psychiatric	Yes/No
	disorder and physical illness. The surgeon / doctor should examine for the following relative	
	contraindications	
	a) Psychiatric disorder	
	b) Physical illness	
	I)Acute febrile illness	
	II)Jaundice or other chronic liver disease	
	III)Anaemia (haemoglobin less than 8 gm.%)	
	IV)Chronic systemic disease, including tuberculosis, brochial asthema, blood	
	dyscrasias, heart disease, uncontrolled diabetes, hypertention and	
	thyrotoxicosis	
	V)Malignancy	
	VI)Skin conditions, including infection involving operation site	
	VII)Pelvic infection, adhesions or mass	
	VIII)Severe nutritional deficiency such as generalized oedema, anaemia, and	
	vitamin deficiency c) Allergy to local anesthesia	
	(alternative anaesthesia or procedure must be provided)	
	d) Gross obesity	
	e) The following conditions in post-partum clients	
	I)Puerperal fever	
	II)Prolonged rupture of membranes (24 hrs)	
	III)Pre-eclampsia or eclampsia	
	IV)Ante partum or post-partum haemorrhage resulting in Hb < 8 gm.%	
	V)Trauma to genital tract	
	VI)History of post-partum psychosis	
6	Whether assessment and screening of client has been done as follows:	
6.1	Whether the client has been physically examined for pulse, Blood pressure, respiratory rate,	Yes/No
	temperature, body weight, general condition, and nutritional status, auscultation of heart and	
	lung, examination og abdomen, pelvic exam., and other exam. As indicated by clients medical	
	history or general physical exam.	
6.2	Laboratory examination: blood test for haemoglobin, urine analysis for sugar, and albumin and	Yes/No
	other laboratory examination	,
6.3	Final medical assessment of the operating surgeons: whether surgeon has verified of the client	Yes/No
	including abdominal/pelvic examination before conduction of the surgery	/
7	Whether instruction relating to prevention of infection has been followed?	Yes/No
7.1	Whether cleaning and fumigation of the OT has been done	Yes/No
7.2	Proper arrangement for decontamination of articles after surgery is available for items that	Yes/No
- · · -	come in contact with blood or other body fluids by placing in the solution of disinfectant for 10	. 55/146
	minutes (surgical instruments, gloves, needles, syringes, cotton gauze etc.	
7.3	Sterlisation procedure of equipments/ instruments required for surgery has been carried out	Yes/No
_	as laid down in the guidelines	,
	<u>. </u>	1

Signature and name of the surgeon Date

Annexure-2

A. Application for sterilization operation and consent form

Name of Patient: shri/smt
Husband / wife's name
Father's name and address
Operation centre
Dear sir / Madam,
Kindly make arrangements for my sterilization operation. My age is years and my
husband/wife's age isyears.
I am married and husband/ wife is alive. We have Male and Female living
children. The age of my youngest child is years. I have decided to undergo sterilization operation
independently and on my own without any outside pressure, inducement or force. I am aware that other
methods of contraceptions are available to me. I know that for all practical purposes this operation is
permanent and that after the operation will be unable to have any more children. I also know that there are
still some chances of failure of the operation for which the hospital / institution and operating doctor will
not be held responsible by me or my relatives or any other person whomsoever. My wife / husband has not
been sterilized previously. I am aware that I am undergoing operation, which carries an element of risk. I
have been explained the eligibility criteria for the operation and I affirm that I am eligible to undergo
operation according to criteria. I agree to undergo the operation under any type of anesthesia which the
doctor thinks suitable for me. After sterilization operation if I get pregnant, then I shall report within four
weeks to the doctor / hospitaland will get abortion done free of cost. Under such circumstances, the state
government will pay a compensation of RS. 5000/- to me which will be acceptable to me. I know that if I am
unable to get the pregnancy aborted within four weeks of pregnancy, then I will not be entitled to claim any
compensation from any court of law in this regard. I agree to come for follow-up to the centre / doctor as
instructed, failing which I shall be responsible for the consequences, if any.
I have read the above mentioned facts / information* in my own language.
Thave read the above mentioned facts / information in my own language.
Religion:
Age:
Business / occupation:
Signature of applicant/acceptor:
Signature of witness:
Full Name and address:
Tuli Name and address.
Shri / smt have been explained other
methods of contraception available and the failure associated with other methods have been explained fully.
The thous of contraception available and the failure associated with other methods have been explained rang.
Signature of counselors:
B. <u>Certificate of medical officer</u>
I certify that I have satisfied myself that shri / smt is within the
eligible age group and is mentially and medically fit for a sterilization operation. There is no evidence that
he / she has undergone a sterilization operation previously. I have explained all clauses to the client and that
this form has the authority of a legal document.
, ,
Signature of operating doctor:
Name and address of doctor:
Signature of medical officer:
Name and address of medical officer:
Name and address of medical officer:

C. <u>Denial of sterilization</u>

I certify that shri / smt is not suitable client for sterilization / resterilisation for the following reasons.
1
2
He / she has been provided the following alternative methods of contraception.
Signature of counsellor or doctor making decision:
Name and address of counselor or doctor:* *Counselor can be any health personnel including doctor
D. <u>For official use only</u>
To be filled by examining doctor
Note: if the surgeon is himself health examiner, the certificate may be given by him.
Age of client according to appearance:
Blood pressure:
Whether client has under gone sterilization earlier or not
As per examination by doctor, the client is mentally and medically fit for sterilization operation.
I have confirmed from client regarding his/her marital status and number of living children. I have explained pros and cons of the sterilization operation to the client and he/she him/her self is mentally ready for operation.
Signature of the client:
Signature of the surgeon:
Name of doctor in capital letters:
Present place of posting:
E. <u>Certificate of the surgeon</u>
I have performed the sterilization operation. During the operation there was no visible signs of earlier sterilization and as per appearance he/she was within the age limit for sterilization. If it is female sterilization the type of operation performed: Abdominal / Vaginal / Laproscopic / Minilap General / Local anesthesia used
Generally Local anestricsia asca
Signature of the surgeon:

Economical, social and demographic details of the client undergoing sterilization operation

(IVIOI	iting report of the district family werrare bureau should be accompanied by the following performal
1.	Name of client
2.	Name of head of family shri/smt
3.	Name of father / husband
4.	Mohalla House no
5.	PHC / Urban centre
6.	Ward
7. o	Religion Caste: General / SC / ST / NT / VJNT / OBC
	Whether married Yes / NO
	Age of applicant (completed years)
	Age of husband / wife (completed years)
	No. of alive children a) Sons
	Age of marriage
	Educational qualification:
	a. Husband: Illiterate /literate/primary/junior school/high school/graduate and above
	b. Wife: Illiterate /literate/primary/junior school/high school/graduate and above
15.	Difference from the last termination of pregnancy (delivery or abortion) years months
	nt particulars:
•	t given to applicant Rupees paise paise
Name .	re of applicant
	Follow up
	concerned with the service of applicant
	post
	f appointment
	Promoters
	Health worker
	Medical officer PHC
	Surgeon
	ctomy method adopted: Abdominal / vaginal / Laproscopic /Laprotomy
Type of	anesthesia: General / Local / Spinal
Full nar	ne of person going to give follow-up
Present	address
	Other Information
1.	Whether any contraception method has been adopted earlier: Yes / No
	If yes, 1) Name of method
2.	Whether promoter of applicant is regional worker of family welfare programme: Yes / No
	If yes, whether applicant is inhabitant of the jurisdiction of that worker: Yes / No
3.	Reason for application of sterilization: Limited family / diseasers / financial / other
	rtify that above mentioned particular is correct.
	ce
Sigi	nature & Name and address

Section – II Consent form for Gynaecologic Surgery Procedures Informed Consent for Artificial Insemination (AI/CI/IUI/SIFT/SPF)

- "	
	Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
,	he following condition: (Doctor to document in patient's own words)
The following procedure will be perform	ned:
The insemination of the wife artificially w	vith the semen / sperms for achieving conception .
Site of insemination: (tick which is appop	iate)

- Vaginal insemination
- o Intra-cervical insemination
- o Intrauterine insemination
- Sperm intra fallopian transfer
- Sperm perfusion of fallopian tubes

With sperms of: (tick which is applicable)

- Sperms of the Husband
- Sperms of the Donor

Risks of the AIH or IUI:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs .

Specific Risks:

There are some risks/ complications, which may happened specifically with this type of procedure. They Include but are not limited to the following:

- We understand that even though the insemination may be repeated as often as recommended by the doctor, there is no guarantee or assurance that pregnancy or a live birth will result.
- We have also been told that the outcome of pregnancy may not be the same as those of the general pregnant population, for example in respect of abortion, multiple pregnancies, anomalies or complications of pregnancy or delivery.
- The procedure carried out does not ensure a positive result, nor does it guarantee a mentally and physically normal child.
- Although serological screening tests are done but there is rare possibility of sexually transmittable infections to women.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:
The likelihood of success of above procedure is: Good / fair / poor

Patient consent:

I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care. There is no guarantee or assurance that pregnancy or a live birth will result.
- The outcome of pregnancy may not be the same as those of the general pregnant population, for example in respect of abortion, multiple pregnancies, anomalies or complications of pregnancy or delivery.
- The procedure carried out does not ensure a positive result, nor does it guarantee a mentally and physically normal child. This consent holds good for all the cycles performed at the clinic.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - About Artificial (Intrauterine) insemination (by Husband or Donor sperms)
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On	t	he	basis	of	the a	bove sta	itements,	l request	to r	าave tl	ne proced	lure.
----	---	----	-------	----	-------	----------	-----------	-----------	------	---------	-----------	-------

Signature of wife:	
Signature of Husband:	
Signature of Doctor:	Signature of witness:

Section III Informed consent form Obstetric surgery

Section III Informed consent form – Obstetric surgery Informed consent form for caesarean section

Family name:	Given name(s):
•	Date of birth/Age: Sex: M /F
Condition and procedure:	
This condition requires the following pro	ocedure: (Doctor to document in patient's own word)

The following procedure will be performed:

This operation is to remove the baby from the uterus, which is done by a cut in the lower abdomen and cut in the uterus. The doctor then brings out the baby and occasionally forceps may be needed to help the birth of the baby through the abdominal wound out of the uterus. The placenta will also be removed and the cord cut between the placenta and the baby.

Risks of a caesarean section:

There are risks and complications with this procedure. They include but are not limited to the following. **Common risks and complications (>5%) include:**

- o Infection in the operation site or pelvis or urinary tract. Treatment may be wound dressings and/ or antibiotics.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you
 have been taking blood thinning drugs.
- Treatment is with antibiotics and a drain into the wound for a few days.
- The uterus may not contract properly after the operation. This can lead to excess vaginal bleeding, treated with hormone injection/s to contract the uterus. In severe cases, it may be necessary to remove the uterus, preventing future pregnancies.
- Adhesions (bands of scar tissue) may form and cause bowel obstruction. This can be a short term or a long term complication and may need further surgery.
- Increased risk in obese people & smokers of wound infection, chest infection, heart and lung complications, and thrombosis.

Uncommon risks and complications (1-5%) include:

- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Minor skin cut/s to the baby, more common in breech births when the baby's bottom is against the wall
 of the uterus. The baby's bottom, face or body may be cut when the uterus is cut. This usually heals
 quickly, treated with a band-aid.
- Injury to other organs such as the ureter/s (tube leading from kidney to bladder) bladder or bowel.
 Further surgery will be needed to repair the injuries.
- The wound may not heal normally. The scar can be thickened and red and may be painful. This is permanent and can be disfiguring.
- o Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off (thrombo-embolism)and go to the lungs which could be fatal.
- The scar may rupture (burst) in future pregnancies or during labour. The risk is highest if the cut is made down the uterus rather than across the lower part of the uterus. Scar rupture can be fatal or lead to hysterectomy as a life saving measure.
- o Fertility may be reduced after a caesarean section.

Rare risks and complications (<1%) include:

For future pregnancies a slightly higher risk of placenta previa. (The afterbirth lies across the lower part
of the womb). This can cause major blood loss and the placenta may grow into surrounding organs such
as the bladder. A blood transfusion may be needed. Removal of the uterus and repair of the bladder
and other organs may be required.

- Severe bleeding from large blood vessels about the uterus, which will need emergency surgery to repair the damaged blood vessels. A blood transfusion may be required to replace blood loss.
- o Rarely, in severe cases, the uterus may have to be removed, stopping future pregnancies.
- Bowel blockage after the operation. This may be temporary or in the longer term. If it doesn't get better
 with initial treatment, bowel surgery may be necessary which may include a colostomy. This can be
 temporary or permanent.
- Poor wound healing and the wound may burst open which may require long term wound care with dressings and antibiotics, or a hernia can form in the long term. This may need repair by further surgery.
- Death as a result of this procedure is possible.

Significant risks and procedure options:	
Risks of not having this procedure:	
Anaesthetic:	
The likelihood of success of above procedure i	s: Good / fair / poor
Patient consent:	
I request Dr	to perform upon me the above mentioned procedure
I acknowledge that the doctor has explained;	

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic (Epidural / spinal anaesthesia)
 - o Blood & Blood Products Transfusion
 - About Caesarean section
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure			
Name of Patient:	Signature:	Date:	
	Signatu		

<u>Section III Informed consent form – Obstetric surgery</u>

Informed Consent for Removal of Ectopic Pregnancy by Laparoscopy / Laparotomy

Family name:Given nar	ne(s):
Address:	
Condition and procedure:	
This condition requires the following procedure. (Doctor to a	,

The following procedure will be performed:

An ectopic pregnancy is where the pregnancy takes place outside the uterus, usually in the fallopian tube sometime in abdomen or ovarian. This may need to be removed by either a laparoscope (key hole) approach or through a large incision know as laparotomy.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
 to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

- There is severe internal bleeding in the abdomen. This may need blood transfusion or further surgery.
- Damage to bladder, bowel or ureter (drainage tube from kidney to bladder). A larger cut to repair the
 damage may be necessary. In case of bowel injury, it may be necessary for a temporary colostomy to
 allow the injured bowel to heal. This colostomy is usually closed by surgery a few weeks later.
- Damage to the womb due to the instruments used to move the womb. A perforation (small hole) can happen in 1 in 100 women and will usually heal without further problem. Very rarely, the uterus is removed if bleeding is life threatening.
- Rarely gas, used to inflate the abdomen, can cause heart and breathing problems in 1 in 60,000 women. Death is a very rare risk.
- Removal of only the pregnancy and not the tube may result in some placenta (afterbirth) being left behind and continue to grow. This is treated with further surgery or medication.
- Ovarian pregnancy may require removal of that ovary completely.
- Future pregnancy: There may be recurrence of ectopic pregnancy.
- A minor wound infection or womb infection, which is treated with antibiotics. More serious infections
 such as pus collections inside the abdomen are treated in hospital with antibiotics and sometimes
 further surgery.
- Adhesions (bands of scar tissue) may cause blockage of the bowel and/ or difficulty getting pregnant. This can be a short term or a long-term complication and may need further surgery.
- Abnormal wound healing. The wound can be thickened and red and may be painful.
- Hernia may form where the cuts were made and cause pain and swelling.
- Very low possibility of a fistula (a connecting passage between one area and another) developing.

Anaesthetic:		
Patient consent:		
l request Dr	to perform upon me the	above mentioned procedure.
l acknowledge that the doctor has explained	:	
 My medical condition and the proposed something unexpected. I understand th The anaesthetic required for this proced 	e risks, including the risks tha	t are specific to me.
to me.		
• Other relevant procedure/ treatment o	ptions and their associated ris	sks.
I consent to tackle/operate/remove anyMy prognosis and the risks of not having	g the procedure.	
 That no guarantee has been made that been carried out with due professional 	care.	ny condition even though it has
 The procedure may include a blood trans 		
 Tissues and blood may be removed and stored and disposed of sensitively by th 	e hospital.	
 If immediate life-threatening events had discussions with the doctor or my Acute 	· ·	ney will be treated based on my
 A doctor other than the Consultant may undergoing further training. 	conduct the procedure. I und	derstand this could be a doctor
 I have been given the following Patient About Your Anaesthetic 		ined verbally to me:
o Blood & Blood Products Transfusion		
 About Removal of Ectopic Pregnancy 		
 I was able to ask questions and raise co procedure and its risks, and my treatme answered to my satisfaction. 		
 I understand I have the right to change preferably following a discussion with n 	•	ng after I have signed this form but,
 I understand that image/s or video foot that these image/s or video/s will assist 	age may be recorded as part	
 I accept that medicine is not an exact so the results. Understanding these limitat 	cience and understand that no	
 I certify and acknowledge that I have re alternatives and expected results of the to consider my decision. 	ad this form or had it read to	· · · · · · · · · · · · · · · · · · ·
On the basis of the above statements, I req	uest to have the procedure	
Name of Patient:	Signature:	Date:
Signature of Doctor:	Signatu	ure of witness:

Section III Informed consent form – Obstetric surgery Request and Consent form for MTP (MTP ACT Regulation)

Request Form for M.T.P.

I, the undersigned, Mrs/Miss		(Name)
Husband's /father's Name		request
Dr	to tern	ninate my pregnancy:
	my physical or mental health.	
	k that if the child were born it would suffer from	such physical or mental
abnormalities as to be seriou		
 As this pregnancy has resulted 	, , ,	
	ed as a result of failure of the contraceptive tech	-
	coitus interruptus /periodic abstinences / tubect	tomy or tubal ligation /
vasectomy/ (a		
	injury to my physical or mental health by reason	of my actual /
reasonably foreseeable envi	ronment.	
Signed in my presence	Signature	
Name & signature & address of the v	witness	
· ·		
<u>Cor</u>	nsent Form (C-form) For M.T.P.	
I, the undersigned, Mrs. / Miss		(Name)
Aged completed year	rs and residing at	
Give my free consent for the operati	ion of Medical Termination of my pregnancy by	
Dr	under	anesthesia to be
administered by Dr	I have beer	n explained and I have
understood the procedures propose	d to be employed for terminating the pregnancy	and for anaesthetizing
me along with the likely risks and co	mplication of both these, including the remote p	ossibility of this
pregnancy continuing in-spite of the	procedures employed for terminating it. It also	understand that
·	dones or in addition to them for terminating the	
- · · · · · · · · · · · · · · · · · · ·	ary or desirable and I consent for them, if the su	rgeon and /or the
anesthetist think them essential and	beneficial to me.	
Signed in my presence	Signature	
Signature & Name and address of th	ne witness	

(MTP certification as per regulation)

(Secret)

Form - I (See Regulation - 3 of MTP act)

, see,	.guidion 5 or iviii det/
(Name and qualification of th	ne Registered Medical Practitioner in Block Letters)
(Full address of	of Registered Medical Practitioner)
(Name and qualification of th	ne Registered Medical Practitioner in Block Letters)
(Full address of	of Registered Medical Practitioner)
bear the serial No	minated the pregnancy of the women referred to above who in the admission Register of the hospital/ approved place (here, mention time in weeks) and this ination.
Date:	Signature (s) of registered Medical Practitioner(s)
Place:	
Strike out whichever is not applicable Of the reasons specified in item (i) to (v) wi	rite the one which is appropriate:-

- (i) In order to save the life of the pregnant woman.
- (ii) In order to prevent grave injury to the physical or mental health of pregnant women
- (iii) In view of the substantial risk that if the child was born it would suffer from such physical or mental abnormalities as to be serious handicapped.
- (iv) As the pregnancy is alleged by pregnant women to have been caused by rape.
- As the pregnancy has occurred as a result of failure of any contraceptive device or method (v) used by married woman or her husband for the purpose of limiting the number of children.

Note:- account may be taken of the pregnant women's actual or reasonably foreseeable environment in determining whether the continuance of a pregnancy would involve a grave injury to her physical or mental health

Informed consent for MTP-1st trimester

Family name:	Given name(s):	
Address:		Sex: M /F
Condition and procedure:		·
The doctor has explained that you have the follo	owing condition: (Doctor to document in pa	itient's own words)
· · · · · · · · · · · · · · · · · · ·		

The following procedure will be performed:

Dilatation (stretching) of the cervix and removal of the foetus and placenta by curettage using an instrument with suction attached inserted into the uterine cavity.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific risks:

- The haemorrhage could occur from the uterus, which may require treatment with blood transfusion.
- Infection may occur in the uterus and tubes, which will require treatment with antibiotics.
- Removal of the lining of the womb (endometrium) can lead to scarring inside the uterus and may cause difficulty with future fertility.
- Incomplete removal of tissue within the uterus is possible which might lead to the necessity of further surgery.
- Damage or tearing of the cervix which may require repair and possibly lead to early pregnancy loss in future pregnancies.
- Damage to the uterus due to a perforation and possible bowel damage. This may require further surgery including the possibility of resection of bowel which may include a colostomy.
- Damage to fallopian tubes is possible, which will affect fertility and there is a small possibility of damage to bladder and blood vessels which could require further surgery. This may include laparoscopy, laparotomy or hysterectomy, and a longer hospital stay than expected.
- Rarely, air may be introduced into the circulation leading to cardiac arrest.

Significant risks and procedure options:
•
 Continue with the pregnancy and keep the baby
 Continue with the pregnancy and have the baby adopted after the delivery.
Risks of not having this procedure:
Anaesthetic:
The likelihood of success of above procedure is: Good / fair / poor

Patient consent:

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - o About Your Anaesthetic and/Epidural & Spinal /Local Anaesthesia
 - MTP first trimester
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:	
	Ü		
Signature of Doctor:	Cigno	ture of witness:	
Signature of Doctor		ture or withess	.

Informed consent for MTP-2nd trimester

Family name:Given name(s):
Address: Date of birth/Age: Sex: M /F
Condition and procedure:
The doctor has explained that you have the following condition: (Doctor to document in patient's own words)
The following procedure will be performed:
Medications and/ or catheter will be inserted into the vagina/uterus that will induce labour. The foetus
(baby) and placenta will be delivered vaginally. The delivery of the foetus and placenta may take up to 48
hours and rarely, 72 hours.
Risks of the procedure:
There are risks and complications with this procedure. They include but are not limited to the following.
General risks:
 Infection can occur, requiring antibiotics and further treatment.
 Allergic reaction can occur from medicines or by blood transfusion.
• Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
• Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
 Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
Heart attack or stroke could occur due to the strain on the heart.
• Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
to the lungs and produce damage to organ
 Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
 Cardiac arrest & Death as a result of this procedure is possible.
Specific risks:
 Incomplete passing of placenta, which may need surgery.
 Excessive bleeding from the uterus, which may require treatment with blood transfusion.
 Infection in the uterus and tubes, which will require treatment with antibiotics.
• Damage or tearing of the cervix, which may require repair and possibly lead to early pregnancy loss in future pregnancies.
 Rupture of the uterus, which may need surgery to either repair it or to remove the uterus and a longer
hospital stay than expected.
 Failure of procedure: To bring about the desired termination, a further attempt may then be made, or
alternatively, surgery including laparotomy and hysterotomy, where a cut is made in the abdomen and
then into the uterus.
Significant risks and procedure options:
Relevant treatment options include:
Continue with the pregnancy and keep the baby
 Continue with the pregnancy and have the baby adopted after the delivery.
Risks of not having this procedure:
Anaesthetic:

The likelihood of success of above procedure is: Good / fair / poor

-				
Pati	ient	can	COL	۱+۰

I request Dr. to perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s:
 - About Your Anaesthetic
 - MTP second trimester
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure			
Name of Patient:	Signature:	Date:	
	3		
Signature of Doctor:	Signature	of witness:	
Signature of Doctor	3igilatul	= UI WILIIE33	

Section III Informed consent form – Obstetric surgery Informed Consent for Epidural Pain Relief for Labour Pain

Family name:	Given name(s):	
•	Date of birth/Age:	
Condition and procedure:		
The doctor has explained that you ha	ve the following condition: (Doctor to document	in patient's own words)

The following procedure will be performed:

An epidural is given into 'the epidural space' of your back by means of a very fine plastic tube which is inserted through an epidural needle and the needle is removed after the tubing is in place. Low strength local anaesthetic and other pain relieving drugs are given through the tubing to decrease pain. It works by blocking the pain signals from reaching your brain. The fine plastic tube is taped to your back and drugs can be given through this fine tube until your baby is born. This analgesia or anaesthetic takes 15 – 30 minutes to work.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

Most Common side effects and complications of epidural analgesia or anaesthetic

- Nausea, vomiting, itching and shivering can occur
- Blood pressure could fall & Headache can occur
- Pain, backache and/or bruising at injection site can occur
- Sometimes the epidural anaesthetic works only partially.
- Problems in passing urine, which is usually temporary.
- Haematoma or bleeding, If you take blood thinning medicines, you are more likely to get a haematoma as it may affect your blood clotting. Your anaesthetist will discuss this with you.

Less common side effects and complications of epidural analgesia or anaesthetic

- Severe headache If this happens you may need to have bed rest for several days. Sometimes a 'blood patch' is needed to be done to take away this headache.
- A change to a general anaesthetic for Caesarean Section maybe necessary if the epidural/spinal is not adequate
- Intense itching or rash can be due to drug allergy.
- Temporary nerve damage is remote possibility.

Uncommon side effects and complications from epidural analgesia or anaesthetic

- Infection around the injection spot.
- Nerve damage due to the needle when doing a block.
- Overdose of drugs.
- Cardiac arrest.
- An existing medical condition getting worse.

Very rare risks

- Permanent nerve damage with possible paralysis.
- Blood clot with spinal cord damage.
- The block may go higher than planned and affect breathing by paralysing the breathing muscles.
- Breakage of needles, catheters etc possibly requiring surgery to remove them.
- Epidural abscess & Meningitis
- Death

Disadvantages of an epidural anaesthetic

- It can slow down the second stage of your labour.
- You are more likely to need forceps or a vacuum extraction to help the baby out.
- Occasionally, your legs may feel very heavy and numb, this makes walking around difficult.

Risks to your baby

Some drugs that are given to	you during labour w	ill cross the placenta	and appear to	o have little or n	o effect
on the baby.					

Significant risks and procedure options:	
Risks of not having this procedure:	
The likelihood of success of above procedu	ure is: Good / fair / poor
Patient consent:	
I request Dr	to perform upon me the above mentioned procedure.
I acknowledge that the doctor has explaine	rd;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your epidural pain relief for labour pain
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:
	0	
Signature of Doctor/Anesthetist:		
Signature of witness:		

Section III Informed	<u>d consent form – Obstetric surgery</u>	
Inform	ned consent for use of Mesopristol m	edicine in MTP
	Given name(s):	
Address:	Date of l	birth/Age: Sex: M /F
Condition and Treatment	t:	
The doctor has explained	I that you have the following condition:	
(Doctor to document in po		
This condition red	quires the use of drug Misoprostol for ripening o	of cervix & induction of labour for
the purpose of medical te	ermination of pregnancy. Misoprostol tablets ma	ay be administered to you orally or
intra-vaginally.		
I have read the in	nformation leaflet regarding using this drug. My	signature indicates my
acknowledgement of rece	eiving and understanding this information and c	consent for the use of this
medication for the above	e purpose.	
-	g is a synthetic prostaglandin. The Prostaglandins	
	vomb and induce uterine contractions, mainly fo	
•	in induce miscarriage for the purpose of termina	
_	20 weeks of pregnancy. It is also used to treat ha	emorrhage (bleeding) after normal
delivery.		
	ently is not licensed for the purpose of inducing	
· ·	censed for the treatment of gastrointestinal ulco	ers in men and non- pregnant
women.		
-	nal and local trials and research have demonstra	
	our for the purpose of medical termination of pr	egnancy or stillbirth and for
	to surgical termination of pregnancy.	
•	ntly widely used around the world for these purp	ooses. It has been shown to be as
effective as other license	d medications, with fewer side effects.	

Risks and side effects of Misoprostol

There are risks and side effects with taking Misoprostol drug. They include but are not limited to the following.

- Common side effects of the medication such as nausea, vomiting, diarrhoea, chills, abdominal cramps, dizziness and low grade fever.
- Strong, sustained uterine contractions may occur after repeated intra-vaginal doses of Misoprostol.
- In women who have previously had a caesarean section, there have been rare reports of rupture of the uterine scar associated with Misoprostol induction of labour (risk of 1-4%). This is not unique to Misoprostol, and can occur whenever labour is induced or promoted in women with a uterine scar. A modified lower dose is used to reduce this potential risk. If uterine rupture occurs then an operation is required to control any bleeding, repair the uterus and deliver the baby but In severe cases, hysterectomy (removal of the womb) may be needed if bleeding cannot be controlled. As mentioned above, this can occur with the use of other licensed prostaglandin medications as well.
- If drug is used for the purpose of medical termination of pregnancy there is occasional chance of abortion failure and the need for surgical intervention for any reason may be aroused.

Significant risks with this medication:
Risks of not having this medication:

Patient consent:

I acknowledge that the doctor has explained;

- The risks of this medication being used, including the risks that are specific to me, and the likely outcomes.
- Other relevant procedure/treatment options and their associated risks.
- My prognosis and the risks of not having the medication.
- That no guarantee has been made that the medication/treatment will improve my condition even though it has been carried out with due professional care.
- If immediate life-threatening events happen during the treatment, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- Misoprostal is not licensed for the purpose of induction (starting) of labour but has been found to be as effective and safe as similar, licensed medications in numerous trials (research) and in clinical practice.
- A doctor other than the Consultant may conduct the treatment. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - Misoprostol Information sheet
- I was able to ask questions and raise concerns with the doctor about the medication, the proposed treatment and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my treatment and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this drug and that I had ample time to ask questions and to consider my decision.
- I have my doctor's name, address and phone number and know that I can call if I have any questions or concerns.

On the basis of the above statements,

I agree to the use of Misoprostal drug to induce labour for the purpose of medical termination of pregnancy.

Name of Patient:	Signature:	Date:	
	3 3 3 3 3		
Signature of Doctor:	Signat	ure of witness:	
Signature of Doctor		uie di williess	· • • • • • • • • • • • • • • • • • • •

<u>Section III Informed consent form – Obstetric surgery</u> Informed Consent for Cervical Cerclage operation

Family name:	Given name(s):
•	
Condition and procedure:	
	rocedure. (Doctor to document in patient's own words)
••••••	

The following will be performed:

The nature of the procedure is to suture the cervix for closure of cervical canal in the operating room under anesthesia, usually an epidural anesthetic. The purpose of the procedure is to prevent early pregnancy loss or miscarriage that could result from a known or suspected incompetent cervix.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go
 to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

There are some risks/ complications, which may happened specifically with this type of surgery. They Include but are not limited to the following:

- There is possibility of premature rupture of the fetal membranes (amniotic sac) or loss of fluid that may result in premature labor or infection.
- There is possibility premature labor that may result in the need for hospitalization and medication to attempt to stop labor, or the birth of an immature infant.
- There is possibility of injury to bowel, bladder, ureter or other pelvic or abdominal structures and need for immediate surgery or other additional surgery.
- There is possibility of blood loss necessitating transfusion which carries the risk of exposure to AIDS, hepatitis, and other infectious diseases.
- There is possibility of need for prolonged bed rest and abstention from sexual intercourse for the remainder of the pregnancy.
- There is possibility of difficulty in removal of cervical circlage suture and which may be carried under an anesthetic at the hospital.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

Significant risks and procedure options:
Risks of not having this procedure:
Anaesthetic:

The likelihood of success of above procedure is: Good /fair / poor

Patient consent:

I request Dr. perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- I consent to tackle/operate/remove any additional abnormal pathology encountered during surgery.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - o Blood & Blood Products Transfusion
 - About cervical cerclage operation
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:
	5 6 7 7 7	
Signature of Doctor:	Signature o	f witness:

Section III Informed consent form – Obstetric surgery Informed consent for Amniocentesis/ Chorionic Villus Sampling

Family name:	Given name(s):
Address:	Date of birth/Age: Sex: M /F
Condition and procedure:	
The doctor has explained that you have th	ne following condition: (Doctor to document in patient's own words)

The following will be performed:

- Chorionic villus sampling (Abdominal route): A needle is passed through the abdominal wall and through the wall of the uterus (womb) and into the placenta. This is performed with ultrasound guidance. A small piece of placenta is removed and the needle is withdrawn.
- Chorionic villus sampling (vaginal route): A cannula is passed through cervical canal and through the cavity of the uterus (womb) and into the placenta. This is performed with ultrasound guidance. A small piece of placenta is removed and the cannula is withdrawn.
- Amniocentesis: A needle is passed through the abdominal wall and through the wall of the uterus (womb) and into the sac of fluid (amniotic cavity) around the baby. This is performed with ultrasound guidance. A small amount (10-20mls) of fluid is removed and the needle is withdrawn.

Risks of the procedure:

There are risks and complications with this procedure. They include but are not limited to the following. **General risks:**

- Infection can occur, requiring antibiotics and further treatment.
- Bleeding could occur and is more common if you have been taking blood thinning drugs.
- Death as a result of this procedure is possible.

Specific risks:

Miscarriage / loss of foetus due to chorionic villus sampling in 1 in 100 women and miscarriage / loss of foetus due to amniocentesis in 1 in 200 women if performed before 20 weeks pregnancy. This may be due to:

- Infection may introduced into the abdominal wall or amniotic fluid, which can cause foetal death / miscarriage.
- Bleeding from the uterine wall or around the foetus (baby), which can cause foetal death / miscarriage.
- Rupture / leakage of fluid from amniotic sac, which can cause foetal death / miscarriage.
- Possible bleeding from foetal (baby's) circulation into maternal circulation. This could lead to loss of foetus (miscarriage) or development of antibodies against the foetus' blood. The "Anti D" is given to Rhesus negative women to prevent antibody development.
- Premature labour and delivery may be induced if performed after 20 weeks gestation.
- If there is not enough specimen obtained for testing then second procedure (CVS or Amniocentesis) will be offered.
- There is a 1 or 2 in 100 chance that the result will show that the chromosomes of the placenta are different to that of the foetus. If this happens, a second procedure (Amniocentesis) will be offered to clarify the result.
- Occasionally, the test fails due to unknown reasons, in this situation the test may fail to give a "good enough sample" for chromosome and /or DNA analysis.

Risks of not having this procedure:	ignificant risks and procedure options:
	isks of not having this procedure:
Andestnetic.	naesthetic:

Patient consent:

I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/ treatment options and their associated risks.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - Local Anaesthetic for your procedure
 - o Amniocentesis/Chorionic Villus Sampling
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements,	I request to have the r	procedure
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Name of Patient: Signature: Signature:	Date:
Signature of Doctor: Signature of witness:	

FORM OF CONSENT FOR INVESIVE PRENATAL DIAGNOSTIC TECHNIQUES

(Form G of PCPNDT act)

l,w/o
Age years residing at
Hereby state that I have been explained fully the probable side effects and after effects of the prenatal diagnostic procedures.
I wish to undergo the pre-implantation / prenatal diagnostic techniques/test/procedures in my own
interest to find out the possibility of any abnormalities disease/deformity/disorder in my child I am carrying.
I under take not to terminate the pregnancy if the prenatal procedure/techniques/test conducted
show the absence of disease / deformity / disorder.
I understate that the sex of the foetus will not be disclosed to me.
I understand that breach of this undertaking will make me liable to penalty as prescribed in pre natal diagnostic techniques (regulation and prevention of misuse) Act, 1994 (57 Of 1994) and rules framed thereunder.
Date & Place
Signature of pregnant women
I have explained the contents of above to the patient and her companion
(Name address
relationship) in a language she / they understand.
Signature of doctor with Registration no.
Name/ Address/ Registration no. of genetic clinic
Date

"Perinatal (stillborn baby) Autopsy" Consent Form

, and the same of
Family name:Given name(s):
Address: Date of birth/Age: Sex: M /F
Consent by parent(s) or Substitute Decision-maker:
I am the
(Enter baby's name or "baby of mother's name") and
I am his / her parent / substitute decision-maker.
 I consent to an autopsy (Post Mortem examination) to find out the reasons why my baby died, and understand that there is a possibility that a cause of death may not be found. I understand that autopsy may, also, be considered for confirmation of ultrasound findings, investigation of possible chromosomal / metabolic abnormalities, obtain tissue for storage for future investigations, or identify a syndrome which may be related to, but not represent, the final event causing death. I have been given the opportunity to have a Social Support Person e.g. partner or parent or close friend
present during these discussions.
 I have been given and have read and understood the leaflet "Information about perinatal autopsies". I
 understand that the more thorough the examination, the more information the final report will contain I consent to a pathologist performing: A full post mortem examination
 A limited post mortem examination, which only involves the following organs or regions of the body External examination only, which may include x-rays and placental examination
• The practitioner who is obtaining this consent may / may not (please strike out, as appropriate) request further consultation with me in the event that there are unexpected findings in the limited post morten examination.
• I consent to the pathologist keeping and using tissue samples that have been removed as part of the
autopsy for:
 Medical teaching Yes/ No
 Research - review of microscope slides to aid future research Yes/ No
 Quality control (e.g. small tissue samples are useful to show that routine laboratory tests have worked successfully) Yes / No
 I would like the following additional limits put on the autopsy (e.g. "do not examine the head", "no
organs to be kept"):
I consent to the baby's and mother's medical record (tick as appropriate):
 Being provided to the pathologist to enable him / her to do the autopsy
 Being used to review the treatment which he / she received during life
 Being used for medical education provided that mother's and baby's identity is not revealed.
• I consent to further copies of the completed autopsy report to be sent to the following medical
practitioner(s):
• I have been able to ask questions and raise concerns about the autopsy. My questions and concerns have been discussed and answered to my satisfaction.
• I am aware that the autopsy may commence immediately after I give my consent and so it may not be
possible to withdraw my consent should I change my mind.
There is a possibility that a cause of death may not be found. Therefore, parents who want to know why
their baby has died and consent to an autopsy to find out the reasons may be disappointed and regret their
decision about autopsy if a cause of death is not found.
On the basis of the above statements, I give consent to the autopsy.
Name of parent / substitute decision-maker:
Signature: Date: Signature: Date:
Name of Doctor: Designation: Signature:

Blood and Blood Products Transfusion Consent

Family name:	.Given name(s):	
Address:		

This consent primarily includes intravenous or central venous line infusion of fresh blood and blood products, red cells, platelets and plasma (e.g. fresh frozen plasma and cryoprecipitate).

You have a transfusion of blood or blood products, which are from volunteer donors. The blood is collected and screened by Blood Bank having licence from govt. A transfusion is necessary to replace a part of your blood and is given to either;

- To replace red blood cells to treat or prevent anaemia, improve oxygen transport and relieve symptoms of dizziness, tiredness or shortness of breath or
- To give you platelets to help stop or prevent bleeding or
- To give a fresh plasma product to stop, treat or prevent bleeding.

Transfusions are given via needle in your vein or via a central line into your vein. During transfusion you will be closely watched for any possible reactions and will also be regularly checked as to whether you may need another blood transfusion.

The doctor has explained that I have the following medical condition for which I need a transfusion:
(Doctor to document in patient's own words)

Your medical condition requires the following blood product/s.

- o Red Cells
- o Platelets
- o Fresh Frozen Plasma
- Cryoprecipitate

Frequency of	of the treatments: (Doctor can specify that the frequency may vary during the course o
treatment) .	
•	
Start Date of	f Transfusion

Risks of blood and blood products transfusion:

Approximate End Date of Transfusion _

Most common reactions to fresh blood or blood products that are being transfused are:

- High temperature
- Rash, itching and hives
- · Feeling unwell

Rare risks are:

- Having too much blood/fluids giving you shortness of breath.
- Haemolysis, the abnormal breakdown of red blood cells.
- The development of antibodies which may complicate future transfusions and/or organ or tissue transplants. If these complications develop in women they can potentially cause problems for all current and future unborn babies.
- Lung injury causing shortness of breath.
- The spread of viral or other infectious germs from the blood of the donors.
- Very rarely, these above reactions can cause severe harm or possibly death.
- There are specific problems for long term multiple transfusions that may be relevant to your medical Condition which shall be discussed by your doctor.

Other relevant treatment options:

In some situations there may be other choices to a blood transfusion and these include – fluid replacement with saline or other artificial compounds and/or iron supplements. Your Doctor will discuss these with you as some choices are not suitable for everybody.

Risks of not having the blood and/or blood products transfusion:

(Doctor to document in space provided)

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Patient consent:

I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- Other relevant procedure/treatment options and their associated risks.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - o About Blood Transfusion & Blood components
- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the this procedure and that I had ample time to ask questions and to consider my decision.

On the basis of the above statements, I request to have the procedure

Name of Patient:	Signature:	Date:
	_	
Signature of Doctor:	Signature of	witness:

<u>Section – IV Miscellaneous Consent form</u>

Informed Consent for Anaesthesia

Family name:	(s):Given name(s):	
•	Date of birth/Age:S	

Condition and procedure:

I, acknowledge that my doctor has explained to me that I will have an operation, diagnostic, or treatment procedure. My doctor has explained the risks of the procedure, advised me of alternative treatments, and told me about the expected outcome and what could happen if my condition remains untreated. I also understand that anesthesia services are needed so that my doctor can perform the procedure.

Risks of the Anaesthesia procedure:

I understand that these risks apply to all forms of anesthesia. They include but are not limited to the following.

General risks:

- Infection can occur, requiring antibiotics and further treatment.
- Allergic reaction can occur from medicines or by blood transfusion.
- Bleeding could occur and may require a return to the operating room. Bleeding is more common if you have been taking blood thinning drugs.
- Small areas of the lung can collapse, increasing the risk of chest infection. This may need antibiotics and physiotherapy.
- Increased risk in obese people of wound infection, chest infection, heart and lung complications, and thrombosis.
- Heart attack or stroke could occur due to the strain on the heart.
- Blood clot in the leg (DVT) causing pain and swelling. In rare cases part of the clot may break off and go to the lungs and produce damage to organ
- Lose of function of any limb or organ or paralysis/paraplegia/quadriplegia.
- Cardiac arrest & Death as a result of this procedure is possible.

Specific Risks:

I understand that the type(s) of anesthesia service checked below will be used for my procedure and that the anesthetic technique to be used is determined by many factors including my physical condition, the type of procedure my doctor is to do, my doctor's preference, and my own preference. It has been explained to me that sometimes an anesthesia technique which involves the use of local anesthetics, with or without sedation, may not succeed completely and therefore another technique may have to be used including general anesthesia. There are some risks/ complications, which may happened specifically with this type of anaesthesia and surgery. They Include but are not limited to the following: (please tick appropriate box)

☐ General Anesthesia	Expected Result	Total unconscious state, possible placement of a tube into
		the windpipe
	Technique used	Drug injected into the bloodstream, breathed into the
		lungs, or administered by other routes
	Specific Risks	Mouth or throat pain, hoarseness, injury to mouth or
		teeth, awareness under anesthesia,
		injury to blood vessels, aspiration, pneumonia
□ Spinal or Epidural	Expected Result	Temporary decrease or loss of feeling and/or movement to
Analgesia/ Anesthesia		lower part of body
□ With sedation	Technique used	Drug injected through a needle/catheter placed either
□ Without sedation		directly into the spinal canal or immediately outside the
		spinal canal
	Specific Risks	Headache, backache, buzzing in the ears, convulsions,
		infection, persistent weakness, numbness, residual pain,

		injury to blood vessels, "total spinal"	
☐ Major / Minor	Expected Result	Temporary loss of feeling and/or movement of a specific	
Nerve Block		limb or area of the body	
□ With sedation	Technique used	Drug injected near nerves providing loss of sensation to	
□ Without sedation		the area of the operation	
	Specific Risks	Infection, convulsions, weakness, persistent numbness,	
		residual pain, injury to blood vessels	
□ Intravenous	Expected Result	Temporary loss of feeling and/or movement of a limb	
Regional Anesthesia	Technique used	Drug injected into veins of arm or leg while using a	
□ With sedation		tourniquet	
□ Without sedation	Specific Risks	Infection, convulsions, persistent numbness, residual pain,	
		injury to blood vessels	
□ Monitored	Expected Result	Reduced anxiety and pain, partial or total amnesia	
Anesthesia Care	Technique used	Drug injected into the bloodstream, breathed into the	
(with sedation)		lungs, or administered by other routes producing a semi- conscious state	
	Specific Risks	An unconscious state, depressed breathing, injury to blood vessels	
□ Monitored	Expected Result	Measurement of vital signs, availability of anesthesia	
Anesthesia Care		provider for further intervention	
(without sedation)	Technique used	None	
	Specific Risks	Increased awareness, anxiety and/or discomfort	

Risks of not having this	procedure:
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The likelihood of success of above procedure is: Good /fair / poor

Patient consent:

I request Dr. perform upon me the above mentioned procedure. I acknowledge that the doctor has explained;

- My medical condition and the proposed procedure, including additional treatment if the doctor finds something unexpected. I understand the risks, including the risks that are specific to me.
- The anaesthetic required for this procedure. I understand the risks, including the risks that are specific to me.
- I also consent to an alternative type of anesthesia, if necessary, as deemed appropriate by them.
- Other relevant procedure/ treatment options and their associated risks.
- My prognosis and the risks of not having the procedure.
- That no guarantee has been made that the procedure will improve my condition even though it has been carried out with due professional care.
- The procedure may include a blood transfusion.
- Tissues and blood may be removed and could be used for diagnosis or management of my condition, stored and disposed of sensitively by the hospital.
- If immediate life-threatening events happen during the procedure, they will be treated based on my discussions with the doctor or my Acute Resuscitation Plan.
- A doctor other than the Consultant may conduct the procedure. I understand this could be a doctor undergoing further training.
- I have been given the following Patient Information Sheet/s or explained verbally to me:
 - About Your Anaesthetic
 - Blood & Blood Products Transfusion

- I was able to ask questions and raise concerns with the doctor about my condition, the proposed procedure and its risks, and my treatment options. My questions and concerns have been discussed and answered to my satisfaction.
- I understand I have the right to change my mind at any time, including after I have signed this form but, preferably following a discussion with my doctor.
- I understand that image/s or video footage may be recorded as part of and during my procedure and that these image/s or video/s will assist the doctor to provide appropriate treatment.
- I accept that medicine is not an exact science and understand that no guarantees can be given to the results and understanding these limitations.

BLOOD TRANSFUSIONS during surgery
The likelihood of needing a blood transfusion for this procedure is: (please tick appropriate box)
□ Highly unlikely
□ Possible
□ Probable
I understand that there are potential risks from blood transfusions, though rare, and that some of these
include transfusion reaction, hepatitis, and AIDS (Acquired Immune Deficiency Syndrome).
Please tick appropriate box:
 □ I give consent to receive blood or blood products as determined by my anesthetist and doctor to be necessary for my well-being.
 □ I give consent to receive blood or blood products only as an emergency life-saving measure. □ I do not want to receive blood or blood products under any circumstance.
I certify and acknowledge that I have read this form or had it read to me; that I understand the risks, alternatives and expected results of the anesthesia service; and that I had ample time to ask questions and to consider my decision.
On the basis of the above statements, I request to have the procedure
Name of Patient:
Signature of Doctor: Signature of witness:

Discussion and Refusal of Treatment

Patient's Name:	
I am being provided information and refusal form so I may fully understan	d the
procedure	
recommended for me and the consequences of my refusal.	
Risks of Not Having the Recommended Treatment:	
I understand that complications to my Reproductive tract & its function, and/or general health may occur if I	
do not proceed with the recommended treatment.	
These complications include:	
I have had an opportunity to ask questions about these risks and any othe	r risks I have heard or thought
about.	
Acknowledgement:	have week and the all ave
	_ have received the above
information about the proposed procedures. I have discussed my treatme	
Dr and have been given the c	opportunity to ask questions and
have them fully answered.	the need for procedures
Dr has informed me of alternate treatment options, risks associated with not taking procedures, a	and my refusal to take procedures.
I personally assume the risks and consequences of my refusal, and release administrators, or personal representatives, those doctors who have been and all liability for ill effects which may result from my refusal to consent to proposed treatment. I acknowledge that I have read this document in its entirety, that I fully un spaces have been completed or crossed off prior to my signing.	consulted in my case from any to the performance of the
I do not wish to proceed with the recommended treatment.	
I also understand that Dr will/may	refuse to treat me if I refuse
necessary diagnostic or therapeutic proposed procedures.	
Patient or Guardian Signature:	Date:
Treating Doctor Signature:	Date:
Witness Signature:	Date: