

Fibroids in Fertility Patients

Key Document code:	WAHT-TP-027	
Key Documents Owner:	Miss K Brown	Consultant Gynaecologist
Approved by:	Gynaecology Governance Meeting	
Date of Approval:	14 th November 2025	
Medicines Safety Committee Approval	10 th December 202	
Date of review:	14 th November 2028	
This is the most current version and should be used until a revised document is in place		

Key Amendments

Date	Amendment	Approved by
26 th January 2019	Documents extended for 3 years	Mr Hughes
14 th December 2020	Documents approved for 3 years	Miss Blackwell
14 th November 2025	Documents updated and approved for 3 years	Miss Blackwell

Fibroids may be detected at ultrasound scan, hysteroscopy or laparoscopy. They may be sub-mucosal, intramural or sub-serosal. Their presence may reduce fertility by various mechanisms: cavity distortion, tubal occlusion, foreign body, vascularity of the uterus, or inflammation. Fibroids are potentially linked to a 40% reduction in chance of live birth.

Sub-mucosal fibroids

If these significantly distort the uterine cavity, they may be associated with a reduced pregnancy rate during assisted conception. It appears that their removal by transcervical resection of fibroid (TCRF) improves the pregnancy rate.

1. If a sub-mucosal fibroid is suspected on baseline ultrasound scan, then perform a saline scan or organise a hysteroscopy to confirm.
2. If present, then discuss with the patient the advantages and disadvantages of TCRF.

Advantages include a probable improvement in pregnancy rate.

Disadvantages include risks of anaesthesia and procedural risks including infection, adhesions, bleeding, uterine perforation and fluid overload.

1. If you wish to proceed then discuss with the Consultant and add to waiting list.
2. The patient may benefit from at least one injection of Zoladex (3.6 mg) sub-cutaneously 3-5 weeks prior to procedure to shrink the fibroid and ensure a thin endometrium (and therefore a visible fibroid) at surgery.
 - a. If the largest diameter of the fibroid is ≥ 3 cm then two Zoladex injections 4 weeks apart could be given to shrink to fibroid further and reduce operative blood loss.
 - b. An alternative would be a course of Ryeqo
 - c. Discuss this with the operating surgeon to confirm preference

Intra-mural fibroids

These are fibroids within the wall of the uterus. They usually do not distort the uterine cavity and traditionally have been considered not to reduce the pregnancy rate. However, controlled data suggests a lower pregnancy rate in the presence of intra-mural fibroids during IVF treatment.

WAHT-TP-027

As yet, we do not know whether there are any benefits in terms of increased pregnancy rate in removing intra-mural fibroids not distorting the uterine cavity. Occasionally they may cause tubal blockage by external pressure on the tubal lumen at the cornua. If necessary, discuss with a senior doctor in the unit.

Sub-serosal fibroids

These are fibroids on the external wall of the uterus. They do not appear to affect the pregnancy rate. However, if they are significant in size, or causing symptoms; management should be discussed in the Fertility MDT.

Size

<3cm	Limited evidence this will improve fertility
>5cm	Advise surgical removal prior to IVF
<5cm, but multiple	Surgery may be warranted

Management

1. Surgery is the preferred option for fertility preservation – TCRF or myomectomy
 - a. No benefit of UFE prior to surgery
 - b. Advantage of using vasopressin prior to myomectomy for blood loss
 - c. Prior to surgery: consider fibroid MRI mapping, as more accurate than USS

Other options must be discussed in the Fertility MDT:

2. Uterine Artery Embolisation (UAE), which can block the blood supply and shrink the fibroid
3. MR guided focused ultrasound on the uterine leiomyomas

References

Management of Subfertility and Assisted Conception, Joint RCOG/BFS meeting April 2023

Monitoring

This section should identify how the Trust plans to monitor compliance with, and the effectiveness of, this policy. It should include auditable standards and/or key performance indicators (KPIs) and details on the methods for monitoring compliance.

The NHSLA requirements are:

Organisations should measure, monitor and evaluate compliance with the minimum requirements within the NHSLA Risk Management Standards. This should include the use of audits and data related to the minimum requirements. The organisation should define the frequency and detail of the measurement, monitoring and evaluation processes.

Monitoring demonstrates whether the process for managing risk, as described in the approved documentation, is working across the entire organisation. Where failings have been identified, action plans must have been drawn up and changes made to reduce the risks. Monitoring is normally proactive - designed to highlight issues before an incident occurs - and should consider both positive and negative aspects of a process.

The table below should help to detail the 'Who, What, Where and How' for the monitoring of this policy.

Page/ Section of Key Document	Key control:	Checks to be carried out to confirm compliance with the Policy:	How often the check will be carried out:	Responsible for carrying out the check:	Results of check reported to: (Responsible for also ensuring actions are developed to address any areas of non-compliance)	Frequency of reporting:
	WHAT?	HOW?	WHEN?	WHO?	WHERE?	WHEN?
	These are the 'key' parts of the process that we are relying on to manage risk. We may not be able to monitor every part of the process, but we MUST monitor the key elements, otherwise we won't know whether we are keeping patients, visitors and/or staff safe.	What are we going to do to make sure the key parts of the process we have identified are being followed? (Sometimes a year) techniques to consider are; audits, spot-checks, analysis of incident trends, monitoring of attendance at training.)	Be realistic. Set achievable frequencies. Use terms such as '10 times a year' instead of 'monthly'.	Who is responsible for the check? Is it listed in the 'duties' section of the Policy? Is it in the job description?	Who will receive the monitoring results? Where is this a committee the committee's responsibility for monitoring the process must be described within its terms of reference.	Use terms such as '10 times a year' instead of 'monthly'.
	Accurate reporting of Fibroids on USS and discussion of	Analysis of an incidents and surgical complications	Whenever necessary	Fertility Lead and Lead Fertility Nurse	Fertility MDT	Whenever necessary following incidents

	management in Fertility MDT if in question					
--	---	--	--	--	--	--

WAHT-TP-027

Supporting Document 1 - Equality Impact Assessment Tool

Equality and Health Inequalities Impact Assessment (EHIA) Tool

Herefordshire & Worcestershire STP - Equality and Health Inequalities Impact Assessment (HEIA) Form
 Please read HEIA guidelines when completing this form

Section 1 - Name of Organisation (please tick)

Herefordshire & Worcestershire STP		Herefordshire Council	
Worcestershire Acute Hospitals NHS Trust	<input checked="" type="checkbox"/>	Worcestershire County Council	
Worcestershire Health and Care NHS Trust	<input type="checkbox"/>	Wye Valley NHS Trust	
Other (please state)	<input type="checkbox"/>		

Name of Lead for Activity	
----------------------------------	--

Details of individuals completing this assessment	Name	Job title	e-mail contact
	K Brown	O&G Consultant	Kiritea.brown@nhs.net
Date assessment completed			

Section 2

Activity being assessed (e.g. policy/procedure, document, service redesign, policy, strategy etc.)	Title: Fibroids in Fertility Patients
What is the aim, purpose and/or intended outcomes of this Activity?	To ensure the correct management of fertility patients with fibroids
Who will be affected by the development & implementation of this activity?	<input type="checkbox"/> Service User <input checked="" type="checkbox"/> Patient <input type="checkbox"/> Carers <input type="checkbox"/> Visitors <input checked="" type="checkbox"/> Staff <input type="checkbox"/> Communities <input type="checkbox"/> Other _____
Is this:	<input checked="" type="checkbox"/> Review of an existing activity <input type="checkbox"/> New activity <input type="checkbox"/> Planning to withdraw or reduce a service, activity or presence?
What information and evidence have you reviewed to help inform this assessment? (Please name sources, eg demographic information for patients / services / staff groups affected, complaints etc.)	Management of Subfertility and Assisted Conception, Joint RCOG/BFS meeting April 2023 NICE CG156. Fertility problems: assessment and treatment. Sept 2017.
Summary of engagement or consultation undertaken (e.g. who and how have you engaged with, or why do you believe this is not required)	First agreed in Fertility MDT, then MSC, and Gynae Governance
Summary of relevant findings	As per GL

--	--

Section 3

Please consider the potential impact of this activity (during development & implementation) on each of the equality groups outlined below. **Please tick one or more impact box below for each Equality Group and explain your rationale.** Please note it is possible for the potential impact to be both positive and negative within the same equality group and this should be recorded. Remember to consider the impact on e.g. staff, public, patients, carers etc. in these equality groups.

Equality Group	Potential <u>positive</u> impact	Potential <u>neutral</u> impact	Potential <u>negative</u> impact	Please explain your reasons for any potential positive, neutral or negative impact identified
Age				
Disability				
Gender Reassignment				
Marriage & Civil Partnerships				
Pregnancy & Maternity	y			First agreed in Fertility MDT, MSC then Gynae Governance
Race including Traveling Communities				
Religion & Belief				
Sex				
Sexual Orientation				
Other Vulnerable and Disadvantaged Groups (e.g. carers; care leavers; homeless; Social/Economic deprivation, travelling communities etc.)				
Health Inequalities (any preventable, unfair & unjust differences in health status between groups, populations or individuals that arise from the unequal distribution of social, environmental & economic conditions within societies)				

Section 4

What actions will you take to mitigate any potential negative impacts?	Risk identified	Actions required to reduce / eliminate negative impact	Who will lead on the action?	Timeframe
How will you monitor these actions?	Review and investigate as per any concerns via datix and complaints			
When will you review this HEIA? (e.g in a service redesign, this HEIA should be revisited regularly throughout the design & implementation)	GL will be reviewed every 3 years, or earlier if new guidance available			

Section 5 - Please read and agree to the following Equality Statement**1. Equality Statement**

1.1. All public bodies have a statutory duty under the Equality Act 2010 to set out arrangements to assess and consult on how their policies and functions impact on the 9 protected characteristics: Age; Disability; Gender Reassignment; Marriage & Civil Partnership; Pregnancy & Maternity; Race; Religion & Belief; Sex; Sexual Orientation

1.2. Our Organisations will challenge discrimination, promote equality, respect human rights, and aims to design and implement services, policies and measures that meet the diverse needs of our service, and population, ensuring that none are placed at a disadvantage over others.

1.3. All staff are expected to deliver services and provide services and care in a manner which respects the individuality of service users, patients, carer's etc, and as such treat them and members of the workforce respectfully, paying due regard to the 9 protected characteristics.

Signature of person completing HEIA	
Date signed	25.1.26
Comments:	
Signature of person the Leader Person for this activity	
Date signed	25.1.26
Comments:	



Supporting Document 2 – Financial Impact Assessment

To be completed by the key document author and attached to key document when submitted to the appropriate committee for consideration and approval.

	Title of document:	Yes/No
1.	Does the implementation of this document require any additional Capital resources	N
2.	Does the implementation of this document require additional revenue	N
3.	Does the implementation of this document require additional manpower	N
4.	Does the implementation of this document release any manpower costs through a change in practice	N
5.	Are there additional staff training costs associated with implementing this document which cannot be delivered through current training programmes or allocated training times for staff	N
	Other comments:	

If the response to any of the above is yes, please complete a business case and which is signed by your Finance Manager and Directorate Manager for consideration by the Accountable Director before progressing to the relevant committee for approval.