

Introducing a Revolutionary, Non-Surgical Approach to Permanent Contraception

What is FemBloc?

FemBloc is a non-surgical, non-hysteroscopic permanent contraception solution that places a blended polymer to form scar tissue to occlude the fallopian tubes to prevent pregnancy. FemBloc requires no anesthesia or incisions, and is a safe and effective alternative to traditional surgical sterilization methods.



Non-Surgical

No incisions or general anesthesia



Non-Hysteroscopic

Placed similar to intrauterine devices



No Permanent Implant

Blended polymer degrades, leaving no permanent foreign material



Convenient

Simple procedure with no special equipment



Highly Effective

0% pregnancy rate in confirmed occlusion cases¹



Safe

Permanent birth control is achieved with patient's own tissue



Hormone-Free

Preserves the body's natural hormonal balance



Minimal Recovery

Same-day recovery with no downtime

Efficacy & Safety Snapshot¹

0% PF

PREGNANCY RATE

among women confirmed bilaterally occluded at 3 months (n=51).

NO

SERIOUS ADVERSE EVENTS ECTOPIC PREGNANCIES UTERINE PERFORATIONS

MILD & TRANSIENT AES

The most common adverse events were spotting and cramping.

96% of Women

would recommend FemBloc to friends or family.

1. Liu, J. H., Blumenthal, P. D., Castano, P. M., Chudnoff, S. C., Gawron, L. M., Johnstone, E. B., Lee-Sepsick, K. (2025). FemBloc Non-Surgical Permanent Contraception for Occlusion of the Fallopian Tubes. J Gynecol Reprod Med, 9(1), 01-12. doi: 10.33140/JGRM.09.01.05.

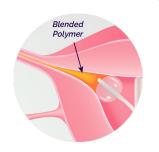




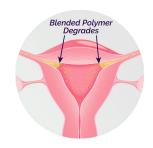
How FemBloc Works



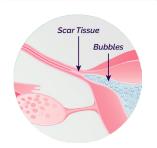
 Delivery system is advanced through the cervix and two balloon catheters are inflated in each uterine cornu.



 A proprietary blended polymer is delivered into each cornu and both fallopian tubes to trigger a natural wound healing response.



 Over ~3 months, the blended polymer degrades, leaving behind the patient's own scar tissue for tubal occlusion.



 An ultrasound-based confirmation test confirms bilateral occlusion prior to patient reliance.

Procedure Overview¹

Attribute	Value
Completed Procedure/Test	99.6% (FemBloc) 99.5% (Confirmation)
FemBloc Procedure Time (mean)	7:36 minutes
Confirmation Test Time (mean)	16:37 minutes
Setting	100% performed in-office
Pain Score (0-10) (average)	4.6 (FemBloc) 3.4 (Confirmation)

Global Momentum

- European CE Mark approval of entire FemBloc system achieved June 2025 under EU MDR.
- UK MHRA approval of entire FemBloc system achieved August 2025.
- · Pivotal trial on-going for U.S. FDA approval.

What Physicians Are Saying

Much easier than surgical tubal ligation or Essure[®].

Highly satisfied with ease of use and patient tolerability.

100% of Investigators would recommend to a colleague.



Learn More

Contact your representative for physician training. www.femasys.com | www.fembloc.com



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