Abstract: A device (10) for occluding the fallopian tube comprising a retention member (12) and a mesh material (30) supported by the retention member. The retention member has a first lower profile configuration for delivery and a second expanded configuration for placement within the fallopian tube. The mesh material is configured to block passage of an egg through the tube. The member has a plurality of tube engagement members (14) to secure the retention member to the fallopian tube.
FALLOPIAN TUBE OCCLUSION DEVICE

BACKGROUND

This application claims priority from provisional application serial no. 60/932,405 filed May 31, 2007, the entire contents of which is incorporated herein by reference.

Field of the Invention

This application relates to a minimally invasive device for occluding the fallopian tubes.

Background of Related Art

Tubal ligation is one method of female sterilization. It can be performed laparoscopically by access through the patient's abdomen where the surgeon severs and closes the ends of the fallopian tubes by tying, applying clamps or cauterizaton. These devices achieve occlusion by external application to the tube.

Other methods involve transcervical access. In some techniques, various agents are injected within each of the fallopian tubes to close or block the tubes. In other transcervical procedures, mechanical devices are inserted and anchored within the tube to promote tissue ingrowth and scar tissue formation to occlude the tubes. In other techniques, radiofrequency energy electrodes are inserted and energized to thermally damage the tube, causing scarring to occlude it.

The need exists for an improved device for occluding the fallopian tubes which can be inserted in a minimally invasive fashion.

SUMMARY

The present invention overcomes the problems and deficiencies of the prior art. The present invention provides a device for occluding the fallopian tube comprising a retention member and a mesh material supported by the retention member. The retention member has a first lower profile configuration for delivery and a second expanded configuration for placement within the fallopian tube. The mesh material is configured to block passage of an egg through the fallopian tube. The retention member has a plurality
of fallopian tube engagement members to secure the retention member to the fallopian tube.

In one embodiment, the mesh is attached to an outer surface of the retention member. In another embodiment, the retention member has a plurality of struts defining a space therebetween and the mesh fills a substantial region of the space. In another alternate embodiment, the mesh is a strip of material connected to the retention member and spanning an opening of the retention member. In this embodiment, the mesh is preferably positioned at a region adjacent the tube engagement members. In a preferred embodiment, the retention member is composed of shape memory material.

The tube engagement members may include a plurality of teeth. In a preferred embodiment, the retention member has a plurality of struts and the struts terminate in the tube engagement members. In a preferred embodiment, the retention member is composed of shape memory material.

The present invention also provides a device for occluding the fallopian tube comprising a tube laser cut to form a series of struts, wherein the tube has a first lower profile configuration for delivery and a second expanded configuration for placement. The struts extend outwardly so that a distal region of the struts has a greater dimension and the struts define a space therebetween. A mesh material is supported by the struts and provides a blocking member to block the egg from passage through the fallopian tube.

In one embodiment, the mesh material fills a substantial area of the space between the struts. In another embodiment, the mesh material is in the form of a narrow strip attached to one or more of the struts. In another embodiment, the mesh material is attached to an outer surface of the struts, and extends across a proximal region of the device.

A method for fallopian tube occlusion is also provided comprising the steps of inserting into the fallopian tube a sheath containing a retention member having a plurality of struts in a reduced profile position, exposing the retention member from the sheath to enable it to expand to engage a wall of the fallopian tube, subsequently inserting mesh material within a space between the plurality of struts, and withdrawing the sheath to leave the retention member in the fallopian tube so the mesh material fills the space within the tube to occlude the tube.
In one embodiment, the retention member has a plurality of shape memory struts and the step of exposing the retention member enables the struts to move toward a shape memorized position.

**BRIEF DESCRIPTION OF THE DRAWINGS**

Preferred embodiment(s) of the present disclosure are described herein with reference to the drawings wherein:

Figure 1 is a perspective view of a the mesh retention member of the fallopian tube occlusion device of the present invention shown in the collapsed position for delivery;

Figure 2 is a transverse cross-sectional view taken along line 2-2 of Figure 1;

Figure 3 is a cross-sectional view taking along lines 3-3 of Figure 1 showing a portion of the retention member within a delivery sheath;

Figure 4 is a perspective view showing the occlusion device in the expanded condition without the mesh therein;

Figure 5 is a view similar to Figure 4 showing the mesh positioned in the device;

Figures 6 and 7 are side and front views of the device of Figure 5;

Figure 8 is an anatomical view showing insertion of the delivery device through the uterus of a patient to access the fallopian tubes;

Figure 9 shows the delivery device being inserted into one of the fallopian tubes;

Figure 10 is a close up view showing the delivery device within the fallopian tube;

Figure 11 is a close up view showing the occlusion device deployed in the fallopian tube;

Figures 12 and 13 are views similar to Figures 10 and 11 except showing placement of the occlusion device in the other fallopian tube;

Figure 14 is an anatomical view illustrating placement of the occlusion device in both fallopian tubes;

Figures 15 and 16 are perspective and side views, respectively, of an alternate embodiment of the present invention showing the occlusion device having a strip of mesh supported therein;
Figure 17 is a side view of another alternate embodiment of the occlusion device having a mesh filling substantially the entire space between the struts;

Figure 18 is a side view of another alternate embodiment of the occlusion device having a mesh positioned on the outside of the retention member; and

Figure 19 is a perspective view of another alternate embodiment of the occlusion device having a mesh supported by a wound wire.

**DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS**

Referring now in detail to the drawings where like reference numerals identify similar or like components throughout, the several views, an occlusion device for placement in the fallopian tubes for contraception is disclosed. The device can be inserted minimally invasively and preferably in an office procedure. A hysteroscope can be used for direct visualization. The occlusion device includes a securement member and mesh material. The securement member provides for attachment to the fallopian tube as well as a support or retention for the various embodiments of the mesh described below.

With initial reference to Figures 1-3 which shows the occlusion device 10 in the low profile collapsed delivery configuration, and Figure 4, which shows the occlusion device 10 in the expanded placement configuration, the occlusion device 10 includes a securement or retention component (member) 12. Securement member 12 has engagement hooks 14 for engaging the fallopian tube wall to retain the securement member 12 in the tube to prevent migration as described below. The mesh can be advanced into the member 12 in situ or alternatively can be positioned in the securement member 12 in the delivery position and then advanced together with the securement member 12 for placement in the fallopian tube. The occlusion device 10 is preferably formed from a laser cut tube, although other ways of forming the device are also contemplated. The mesh is not shown in Figure 1-4 for clarity.

Turning to Figures 4-7 illustrating the device 10 in the expanded (deployed) position, the retention component 12 is in the form of a bell shaped device with struts as described in detail with respect to the vessel filter disclosed in patent application serial no. 10/899,429, filed July 13, 2004, (the '429 application), the entire contents of which is incorporated herein by reference. The device 10 has a proximal end 11a and distal end 11b. Securement member 12 is preferably composed of shape memory material, such as...
Nitinol, with an austenitic shape memorized position illustrated in Figure 4 and has a plurality of struts 13 emerging from apex 18 at proximal end 11a and terminating in engaging or retention hooks 14 at distal end 11b. In this embodiment, six struts are provided although a different number of struts is also contemplated. A retrieval hook 16 is positioned on the proximal end 11a to enable the device 10 to be grasped by a snare or other retrieval device and removed if desired. The struts 13 can be interconnected by interconnecting struts 17 which join adjacent struts. More specifically, the struts preferably divide at region 19 into two connecting struts 17, angling away from each other, and then join at region 21 to form extending strut portions 23 terminating in hooks 14. The interconnecting struts 17 stiffen the device to enhance retention and increase the radial force. They also provide a more symmetric and uniform deployment. The hooks are configured to engage the wall of the fallopian tube for maintaining the position of the occlusion device 10. Thus struts are preferably flared and create a distal opening and a space between the struts. For clarity, not all the identical parts are labeled throughout the drawings. It should be appreciated that materials other than Nitinol or shape memory are also contemplated.

The hooks 14 preferably extend substantially perpendicular from the strut, and are preferably formed by torquing the struts so the hooks bend out of the plane. Preferably, a first set of hooks is larger than a second set of hooks. Preferably, when formed from a laser cut tube, the larger hooks are formed so that they occupy a region equivalent to the transverse dimension of two adjacent struts. Preferably, three smaller hooks and three larger hooks are provided in alternating arrangement in the embodiment utilizing six struts. The smaller hooks are preferably spaced axially with respect to each other and axially inwardly with respect to the larger hooks as in the filter hooks of the '429 application to minimize the collapsed profile (transverse dimension) of the filter when collapsed for insertion. The penetrating tips 14a (Figure 3) preferably point toward the proximal end 11a of the device 10 and penetrate the tissue to retain the occlusion device.

Each of the hooks 14 can have a series of teeth 14c to engage the fallopian tube wall to provide additional retention to prevent movement of the device 10. A heel 14d is provided which extends past the hook to function as a stop to prevent the occlusion device from going through the fallopian tube wall. The angle of the heel 14d in the
smaller hooks is less than the angle in the larger hooks to provide room for nesting of the hooks as shown in Figure 3. For clarity, not all of the hooks are fully labeled.

The securement (retention) member 12 is maintained in a substantially straightened softer martensitic configuration within the delivery catheter or sheath 50 for delivery as shown in Figure 3. The smaller hooks preferably nest within the larger hooks. Cold saline can be injected during delivery to maintain the device 10 in this martensitic condition to facilitate exit from the distal opening 52 at the distal end portion 54 (Fig. 3) of catheter 50. When the struts 13 exit the delivery sheath (tube 50), they are warmed by body temperature and move toward their illustrated memorized expanded position as shown in Figures 4-7.

As shown in Figures 8-11 and 14, the device 10 is preferably inserted through the uterus within delivery catheter 50 and into the fallopian tube. It is positioned in this embodiment with the struts opening in a distal direction. When positioned in the fallopian tube, the hooks 14 engage the tube wall to retain the device.

The device 10 in the embodiment of Figures 1-7 has mesh material positioned within the retention member 12, filling substantially the entire region of the member 12. A small gap 24 can be left at the proximal region (see e.g. Fig. 6). However, in an alternate embodiment, the gap is filled in with mesh so the mesh fills more of the area between the struts as shown for example in Figure 17 wherein mesh 61 fills substantially the entire space between the struts 63. Retention member 60 is otherwise identical to retention member 12 of Figure 1. The mesh is preferably in the form a tightly woven material to provide sufficiently small spaces to effectively block the eggs from the ovaries traveling through the fallopian tube.

The mesh can be delivered within the retention member 12 such that in the collapsed position the mesh is contained and compressed therein. After delivery, it would expand within the space of the retention member 12, i.e. within the space between the struts.

In an alternate embodiment, the retention member 12 would be placed within the fallopian tube first, and then once in place, the mesh would be delivered through the opening in the distal end 11b of the device and within the space between the struts 13 and/or 17. This in situ delivery could occur in embodiments wherein the device 10 is
implanted in an orientation opposite to that of Figure 11, i.e. the opening between the struts would face in the other direction such that the distal limb would be closer to the uterus than proximal end 11a.

In an alternate embodiment, instead of the mesh filling the space between the struts, the mesh is in the form of a narrow strip. The mesh in this embodiment functions as a screen type blocking member, as shown in Figures 15 and 16 and is substantially circular. The retention (securement) member 72 is otherwise identical to securement member 12 of Figure 1, e.g. struts 73 divide at region 79 into interconnecting struts 77, join at region 81, and terminate in engaging hooks 74. The strip 85 of mesh would preferably be positioned slightly proximal of the hooks 74, e.g. at the region where the strut twists out of the plane so as not to interfere with the hooks. However, the mesh could be placed at other locations along the struts as long as it functions as a cover or blocking member to occlude the fallopian tube. Although a thin strip of mesh is shown, other size blocking strips could also be provided. The mesh is shown attached to an inner surface of the struts but could alternatively be attached to the outer surface. It could be attached to one or more of the struts.

As noted above, although the retention member is shown inserted with the engaging hooks 14 facing in the distal direction, it is also contemplated that the retention member could be oriented in the opposite direction. In this version (not shown), the mesh can be inserted along with the retention member or alternatively if desired could be delivered in situ within the distal opening between the struts.

In the alternate embodiment of Figure 18, the mesh 92 is positioned on the outside of the retention member 90. In all other respects, retention member 90 is similar to member 12. In this embodiment, the mesh 92 is placed on an outer region, covering the outer surfaces of the struts and apex region and interposed between the struts and the fallopian tube when placed. Thus, the mesh functions as a sleeve which prevents passage of the eggs as they would be captured within the sleeve or net-like device.

The mesh in the foregoing embodiments can be attached to the retention member by various methods such as bonding, clamping and suturing.

The method of placement of the occlusion device of the present invention will now be described for occluding a fallopian tube and with reference to Figures 8-14 which
illustrate by way of example the device of Figure 5 with the mesh delivered in conjunction with the securement member (the other embodiments are inserted in a similar fashion). A delivery catheter 50 is inserted through an introducer sheath 100 through the vagina and cervix into the uterus U. The catheter 50 is then advanced into fallopian tube F1. For insertion, the retention member 12 (and mesh) is in the collapsed position.

A pusher (not shown) is advanced distally to advance the occlusion device 10 from the catheter 50. As the struts of device 10 are exposed, they are warmed by body temperature and return toward their shape memorized deployed position as shown in Figure 11 to engage the wall W1 of the fallopian tube. The extent they return to their fully memorized position will depend on the size of the fallopian tube. The catheter 50 is then withdrawn into the uterus and inserted into fallopian tube F2 (Figure 12) (or another catheter is used) wherein another occlusion device 10 is exposed from the catheter 50 so the struts move to the expanded position to engage the wall W2 of tube F2 as illustrated in Figure 13. The expanded mesh functions to block the travel of eggs through the tube.

As can be appreciated, although described for closing the fallopian tube, the occlusion device can be used in other areas of the body.

Note, the material inside or outside the retention member could be non-porous or porous. It could alternatively be made of pericardium, SIS, PET, PTFE, or other materials.

In the alternate embodiment of Figure 19, a wound wire 100 provides a support member for mesh 110. The wire as shown has a substantially conical mesh so the diameter (transverse dimension) at region 112 exceeds the diameter (transverse dimension) at region 114. The mesh 110 is provided on an outer surface as shown. Alternatively, the mesh could be positioned inside or a strip of mesh spanning the opening could be provided. The wire could have hooks, barbs or other surfaces to enhance retention in addition to the outward radial force against the fallopian tube wall.

While the above description contains many specifics, those specifics should not be construed as limitations on the scope of the disclosure, but merely as exemplifications of preferred embodiments thereof. For example other materials can be contained or mounted to the retention member to function to block eggs from passage through the
fallopian tubes. Those skilled in the art will envision many other possible variations that are within the scope and spirit of the disclosure as defined by the claims appended hereto.
WHAT IS CLAIMED IS:

1. A device for occluding the fallopian tube comprising a retention member and a mesh material supported by the retention member, the retention member having a first lower profile configuration for delivery and a second expanded configuration for placement within the fallopian tube, the mesh material configured to block passage of an egg through the fallopian tube, the retention member having at least one fallopian tube engagement member to secure the retention member to the fallopian tube.

2. The device of claim 1, wherein the mesh is attached to an outer surface of the retention member.

3. The device of claim 1, wherein the mesh is positioned within the retention member.

4. The device of claim 1, wherein the retention member has a plurality of struts defining a space therebetween and the mesh fills a substantial region of the space.

5. The device of claim 1, wherein the mesh is a strip of material connected to the retention member and spanning an opening of the retention member.

6. The device of claim 5, wherein the mesh is positioned at a region adjacent the tube engagement members.

7. The device of claim 1, wherein the retention member has a plurality of struts forming a space therebetween and the mesh is positioned to span the space.

8. The device of claim 1, wherein the tube engagement members include a plurality of teeth.

9. The device of claim 1, wherein the retention member has a plurality of struts and the struts terminate in the tube engagement members.
10. The device for claim 1, wherein the retention member comprises a wound wire.

11. The device of claim 1, wherein the retention member is composed of shape memory material.

12. A device for occluding the fallopian tube comprising a tube laser cut to form a series of struts, the tube having a first lower profile configuration for delivery and a second configuration for placement, in the second configuration the tube having an expanded configuration, the struts extending outwardly so that a distal region of the struts has a greater dimension and the struts defining a space therebetween, a mesh material supported by the struts and providing a blocking member to block the egg from passage through the fallopian tube.

13. The device of claim 12, wherein the mesh material fills a substantial area of the space between the struts.

14. The device of claim 12, wherein the mesh material is in the form of a narrow strip attached to one or more of the struts.

15. The device of claim 12, wherein the mesh material is attached to an outer surface of the struts.

16. The device of claim 15, wherein the mesh extends across a proximal region of the device.

17. The device of claim 12, wherein the retention member is composed of shape memory material.

18. A method for fallopian tube occlusion comprising the steps of:
   inserting into the fallopian tube a sheath containing a retention member having a plurality of struts in a reduced profile position;
exposing the retention member from the sheath to enable it to expand to engage a wall of the fallopian tube;

subsequently inserting mesh material within a space between the plurality of struts; and

withdrawing the sheath to leave the retention member in the fallopian tube so the mesh material fills the space within the struts to block passage of an egg through the fallopian tube.

19. The method of claim 18, wherein the retention member has a plurality of shape memory struts and the step of exposing the retention member enables the struts to move toward a shape memorized position.
A. CLASSIFICATION OF SUBJECT MATTER

INV. A61F6/22

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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<th>Category*</th>
<th>Citation of document, with indication, where appropriate, of the relevant passages</th>
<th>Relevant to claim No.</th>
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D Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:
  * A*: document defining the general state of the art which is not considered to be of particular relevance
  * E*: earlier document but published on or after the international filing date
  * L*: document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
  * O*: document referring to an oral disclosure, use, exhibition or other means
  * P*: document published prior to the international filing date but later than the priority date claimed
  * T*: later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
  * X*: document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
  * Y*: document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
  * S*: document member of the same patent family

Date of the actual completion of the international search: 11 August 2008

Date of mailing of the international search report: 10/11/2008

Name and mailing address of the ISA:

European Patent Office, P.B. 5816 Patentlaan 2 NL- 2280 HV Rijswijk
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Authorized officer

Lickel, Andreas

Form PCT/ISA/210 (second sheet) (April 2005)
INTERNATIONAL SEARCH REPORT

Box No. II  Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: 18-19
    because they relate to subject matter not required to be searched by this Authority, namely:
    Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery

2.  Claims Nos.:  
    because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3.  Claims Nos.:  
    because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III  Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable.

2.  As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.

3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-11

Remark on Protest

- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.
This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1-11

   A device for occluding the fallopian tube comprising an expandable retention member and a mesh material, whereby the retention member having at least one fallopian tube engagement member to secure the retention member to the fallopian tube.

2. claims: 12-17

   A device for occluding the fallopian tube comprising an expandable tube laser cut to form a series of struts and a mesh material, whereby the struts extending outwardly so that a distal region of the struts has a greater dimension.
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