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DEVICE FOR INTRAVAGINAL, BARRIER-TYPE PREVENTION OF CONCEPTION AND INFECTION

VORRICHUNG ZUR INTRAVAGINALEN VERHÜTUNG VON EMPFÄNGNIS UND INFEKTIONEN MITTELS EINER BARRIERE

DISPOSITIF DE PREVENTION INTRAVAGINALE DE TYPE BARRIERE CONTRE LA CONCEPTION ET LES INFECTIONS

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Description

BACKGROUND OF THE INVENTION

The invention concerns contraception and the prevention of sexually-transmitted diseases by means of a barrier-type device and further concerns a barrier-type device which engages the cervix by extending into the fornices and seals circumferentially against the vagina in the vicinity of the cervix.

The prior art devices for intravaginal, barrier-type prevention of conception and sexually transmitted disease include the diaphragm and the cervical cap. The diaphragm, when seated intravaginally, acts as a barrier to prevent sperm from entering the os of the cervix. A spermicide is used on the inside of the diaphragm to kill sperm traversing over the edge of the diaphragm. Characteristically, the diaphragm is a shallow rubber cup with a rim. A round, coiled metal spring is disposed in the rim. The diaphragm is intended for intravaginal disposition between the posterior aspect of the pubic bone and the posterior fornix. When thus seated, the diaphragm presses against the vaginal wall to form a continuous barrier therewith.

The cervical cap is a small, vaulted or domed device, more rigid than the diaphragm, and dimensioned to fit over a cervix, where it is retained by suction, rather than spring tension as is the diaphragm. The cap includes a dome which rises from a rim. The cap is slipped over the cervical protrusion to block access to the uterus through the cervical os.

None of these barrier-type devices is adapted to fit in close anatomical sealing engagement between the vagina and the cervix. The diaphragm is held against the upper part of the vaginal canal by the tension of the spring in its rim, while the cervical cap typically engages only the tip of the cervix. Both of these devices can be dislodged by sexual arousal, coital activity, or orgasm. As is known, such dislodgement can reduce the effectiveness of the barrier and permit unintended fertilization or unwanted infection, or worse, both.

DE-845832 shows an example of a cervical cap comprising a dome containing spermicidal material, basal ring around the opening of dome, and circumferential basal crown extending from the basal ring. This prior art document forms the basis of the preamble of Claim 1.

In failing to take into account the anatomy and physiology of the vagina and the cervix, the prior art barrier-type devices form imperfect barriers against the penetration of sperm and agents of sexually transmitted diseases into the uterus. As is known, fertilization and infection take place within the uterus, therefore the more sound the barrier to the cervical os, the greater the likelihood of preventing such fertilization or infection. With the emergence of the AIDS (acquired immune deficiency syndrome) epidemic, the risk of life-threatening infection accompanying heterosexual activity is increasing steadily. Therefore, there is a compelling need for intravaginal, barrier-type devices which can substantially decrease the risk of infection by significantly increasing the effectiveness and reliability of the barrier to entry of bodily fluids such as semen, into the cervix.

Accordingly, it is an urgent objective of the invention to provide an intravaginal, barrier-type device which accounts for, and takes advantage of, the anatomy and physiology of the vagina in the vicinity of the cervix to form a more perfect, and a more reliable, seal against the movement of sperm and micro organisms from the vaginal canal through the cervical os into the uterus.

A significant advantage of such a device is the simultaneous reduction of the prospects of conception.

SUMMARY OF THE INVENTION

The invention is based upon the inventor's critical observation that a barrier-type device engaging the cervix and the vagina by seating at the bottom of the fornices and then folding back from the fornices along the interior vaginal walls provides a barrier adapted to the anatomy and physiology of the vagina in the vicinity of the cervix which not only forms a reliable, effective seal, but also resists displacement during arousal, intercourse, and orgasm.

The device conforms to and seamlessly engages the vagina in the vicinity of the cervix. The sealing engagement of the device with the vagina is continuous during all of the expected physiological changes which the vagina and cervix undergo as the result of, for example, sexual activity and menstrual cycle.

The invention provides a barrier tee device for a barrier-type device for preventing conception and infection by sexually-transmitted diseases, comprising:

- a cervical dome fabricated from a flexible physiologically non-reactive material having an opening and a curved surface extending from said opening to a dome top; a continuous fornical rim on said dome, said rim defining said opening; and a continuous annular vaginal brim circumscripting said dome, being formed as a backward fold from said rim and extending downwardly and outwardly with respect to said opening; characterised in that the vaginal brim has an outward concave bias or curve; and said vaginal brim exhibits lateral symmetry and has a vertical dimension between its outside edge and said rim which increases continuously with circumferential symmetry from an anterior extension to a posterior extension of said brim, said anterior extension being diametrically opposite said posterior extension.

Preferably the device includes an annular lip on said rim which extend inwardly from the rim toward a centreline of the dome, and an annular groove being formed by an outward fold of said vaginal brim at said annular lip.

The invention also provides a kit for intravaginal,
barrier-type prevention of conception and sexually-transmitted infection, comprising, in combination:

a barrier-type device according to the invention; and

an applicator including:

a lower body with a main member and a projection attached at an angle to said main member, said projection including a concave recess and an edge for engaging said barrier-type device between said vaginal brim and said dome; and an inserter, slidably disposed in said main member for being moved to a first, retracted position for engaging said vaginal brim to a second, extended position for disengaging said vaginal brim.

Preferably the fornical rim of the device forms an inwardly extending lip and a fluid trapping, inwardly extending groove between said brim and said lip.

The radial distance between said dome and said brim may decrease upwardly with respect to the dome peak and towards the dome opening to a minimum distance just below said rim.

In a preferred embodiment said fornical rim, which is annular and folded, defines fluid trapping means extending radially inwardly of the dome wall, said rim connecting said dome means with said brim means to form a portion of said barrier-type device such that said fluid trapping means is a generally annular groove-shaped cavity in communication with said brim means acting as a scoop and being for trapping fluids entering said brim means acting as a scoop.

The dome means and said brim means may mutually define an annular passageway between said brim means acting as a scoop and said annular groove-shaped cavity, said passageway having a radial width less than the greatest radial width of said cavity. The fluid trapping means preferably extends radially inwardly of said brim means acting as a scoop.

One will appreciate how the summarized invention achieves the above-stated objectives when the detailed description is read with reference to the below-described drawings, in which:

Figure 1 illustrates a perspective view of the barrier-type device of the invention;

Figure 2 is a plan view of the device;

Figure 3 is a sectional view of the device taken along lines 3-3 of Figure 2;

Figure 4 is a sectional view, rotated 90° from Figure 3, and taken along lines 4-4 of Figure 3;

Figure 5 is a partial, sagittal section of female sexual anatomy showing, in schematic representation, the posterior aspect of the vaginal tract and the barrier-type device in place covering the cervix, and sealing to the posterior of the vaginal canal;

Figure 6 illustrates a schematic of the posterior of the vaginal canal from a frontal aspect showing the barrier-type device in place;

Figure 7-10 illustrate the structure and operation of the applicator for insertion of the barrier-type device of Figures 1-6 into the vagina; and

Figures 11 and 12 illustrate an extractor for removal of the barrier-type device from intravaginal placement.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

An intravaginal, barrier-type device for preventing conception and infection by sexually-transmitted diseases is illustrated structurally in Figures 1-4. Preferably, the device is a single piece formed by molding a flexible, physiologically non-reactive material, such as latex rubber. The device 8 includes a cervical dome 10. The dome preferably has a hemi-spherical or hemi-ovoid shape which forms a cervical enclosure. At the opening of the enclosure, the dome 10 transitions to a fornical rim 12, which in turn, transitions by a backward fold to a vaginal brim 14. As best seen in Figures 3 and 4, the dome has a peak, or top, 16.

As Figures 1-4 reveal, the vaginal brim circumscribes, or surrounds, the dome and extends upwardly from the rim 12. The brim exhibits lateral symmetry, as illustrated in Figure 3, where the lateral extensions 14a and 14b of the brim 14 are substantially equal in form and dimension. The width of the brim increases continuously from the anterior to the posterior of the device 8. This increase is revealed in Figures 1 and 4. In Figure 4, the anterior extension 14c of the brim 14 has a smaller dimension 22 than the dimension 23 at the posterior extension 14d of the brim 14. The brim also has an outward bias, or curve, to it, as is seen in Figures 1, 3, and 4.

The width of the brim increases with circumferential symmetry from an anterior vaginal location 18 on the anterior extension 14c of the brim to a posterior vaginal location 20 on the posterior portion 14d of the brim. The circumferential symmetry of the brim 14 is illustrated in Figure 2. In Figure 2, point A over the lateral extension 14a is displaced in a counterclockwise direction on the rim 12 from the anterior vaginal location by a rim segment equal to the rim segment by which the point B over the lateral extension 14b is displaced on the rim 12 clockwise from the anterior vaginal location 18. As shown in Figure 3, the dimension 26 reflecting the width of the brim at extension 14a is equal to the dimension 28 reflecting the width of the brim at extension 14b.

An annular lip 24 is formed in the device 8 by an inward extension of the rim 12 along the periphery of the opening into the dome 10. The lip 24 juts into the opening toward the center line 25 of the dome 10. The backward fold of the rim 12 at the lip 24 by which the rim 12 transitions to the brim 14 forms an annular groove 27 between the brim 14 and dome 10.

The significance of the structure of the barrier-type
device 8 is illustrated in Figures 5 and 6. In Figures 5 and 6, female anatomy is conventionally illustrated as including a cervix 30 through which the cervical os 32 opens to the interior of the uterus 34. The vagina 36 includes an inner surface 37 which transitions to the cervix 30 by way of a fornix, an anatomical fold or recess. As is known, the fornix increases in depth from its anterior to its posterior aspects, which are usually referred to as the anterior fornix 38 and the posterior fornix 39, respectively. Further, at its rear, the vagina 36 transitions, by way of the anterior vaginal wall 40 to the anterior fornix 38, and by way of the posterior vaginal wall 42 to the posterior fornix 39. As shown in the frontal cross section of Figure 6, the fornix is laterally symmetrical.

Figures 5 and 6 illustrate how the barrier-type device of Figures 1-4 is adapted to the anatomy and physiology of the vagina in the vicinity of the cervix. As shown in Figure 5, the proper orientation of the device 8 finds the cervical dome 10 engaging the cervix, which extends through the opening defined by the rim 12. The rim is seated at the bottom of the fornices. The annular lip 24 grips the bottom of the cervix at the fornices, thereby forming a circumferential seal. The rim 14 extends from the bottom of the fornices in close sealing engagement with the walls of the vagina, thereby continuing the circumferential seal along the vagina forwardly from the fornices. The barrier-type device is oriented with respect to the cervix to place the anterior extension 14c against the anterior vaginal wall 40, in the vicinity of the anterior fornix. Similarly, the posterior extension 14d is oriented to engage the posterior vaginal wall 42 in the vicinity of the posterior fornix. As shown in Figure 6, the orientation of Figure 5 disposes the lateral rim extensions 14a and 14b against the lateral vaginal walls in the vicinity of the lateral fornices.

As shown in Figures 5 and 6, not only does the device seal to the vagina and cervix, it orients the annular groove 27 toward the opening of the vagina. Thus oriented, the groove 27 will trap fluids traveling along the inside of the brim 14 toward the dome 10.

In use, a spermicide such as NONOXYNOL-9, is applied to the total surface of the device 8, and the device is inserted into the vagina to engage the cervix with the orientation illustrated in Figures 5 and 6. Thus, seated, the barrier-type device of the invention will seal from the base of the cervix, in the fornices and continuously and circumferentially along the vagina. This seal is superior to the prior art barrier devices. As is known, the diaphragm would engage only the posterior fornix, extending across the top of the vaginal wall forward of the anterior of the cervix. The cervical cap engages only the top of the cervix, and, generally, does not extend fully into the posterior, anterior, and lateral fornices. However, the device 8 fits precisely to the anatomy and physiology of the vagina in the vicinity of the cervix. Such an anatomically adapted form will not only seal effectively when the device is seated, but will also seal reliably by resisting unintentional displacement or dislodgment. The device is retained in place by suction exerted in the cervix by the dome 10, by the grip exerted on the base of the cervix by the annular lip 24, by a snug fitting between the rim 12 of the fornices, and by the outward curve of the brim, which flattens against the vagina. All of these mechanisms help prevent dislodgment of the device and contribute to the seal which the device makes.

An applicator for intravaginal placement of the barrier-type device 8 is illustrated in Figures 7-10. The applicator 50 consists of a speculum-type portion with a main body, or a handle, 52 which transitions into a projection 54 having an angle of, preferably, 40° with the main body 52. As seen in Figure 7, the angled projection 54 has a concave recess 55. A rounded forward edge 56 is provided on the distal end of the projection 54. A movable inserter 57 is slidably disposed in the main portion 52 and moved in the main portion between a first position, indicated by the solid lines in Figure 7, and a second, release, position denoted in Figure 7 by the interrupted lines. The inserter 57 consists of two equivalent elongate pieces 57a and 57b having serrated forward faces 60a and 60b and rear flanges 61a and 61b, respectively. The applicator 50 is formed from a physiologically inert material, such as a relatively rigid plastic, and has all of its edges and ends smoothed and rounded to reduce the prospect of injuring the vagina during placement of the device 8.

Figures 8-10 illustrate how the applicator is used to place the barrier-type device 8. As shown in Figure 8, the barrier-type device is carried on the applicator by engagement of the forward end 56 in the groove 27 of the device 8 formed between the dome 10, the rim 12, and the posterior vaginal extension 14d. The brim 14 is grasped between the serrated surfaces 60a and 60b at the anterior vaginal extension 14c. The inserter 57 is retracted to the first position. In this position, the serrated faces 60a and 60b are forced together with the anterior vaginal extension 14c grasped between them. As the inserter 57 is drawn backward to the first position, the barrier-type device is stretched and retained on the applicator 50 by the tension of the flexible material from which the device 8 is formed. Thus arranged, the barrier-type device on the applicator 50 is placed intravaginally, with the device oriented as in Figure 5.

After traversing the vaginal canal, the end 56 of the applicator 50 will place the posterior vaginal extension 14d of the brim against the posterior vaginal wall and seat the rim 12 in the posterior fornix 39. When this occurs, the anterior vaginal extension 14c will be forward of the anterior fornix 38. Next, the inserter 57 is slid forwardly in the main body 52 to the second position, which will move the anterior vaginal extension 14c into engagement with the anterior fornix 38. When this occurs, the barrier-type device 8 will be retained in the engagement of Figure 5 by the mechanisms described above.

Removal of the applicator can be understood with reference to Figures 9 and 10. When the barrier-type
device is seated by forward movement of the inserter 57, the flexibility of the members 57a and 57b will cause the serrated faces 60a and 60b to spring apart slightly. Next, with the barrier-type device retained in position by suction, the upper member 57b is moved rearwardly toward the first position, which disengages the inserter from the anterior vaginal extension 14c of the brim. The lower member 57a is then pulled rearwardly. This completely disengages the applicator 52 from the anterior fornical extension 14c of the brim, and permits the applicator 52 to be removed.

As shown in Figures 11 and 12, an extractor 64 has an elongate portion 66 with a cross section substantially flattened in a plane. The distal tip of the elongate portion 66 transitions to a hook 68. The hook curves backwardly over the elongate portion and is flattened in the same plane. An operating handle 69 is attached to the proximal end of the portion 66, to be substantially perpendicular to the plane of flattening of the portion and hooks.

The barrier-type device 8 is extracted by insertion of the elongate portion 66, hook first, flat into the vagina in the direction of the anterior fornix. When the anterior fornix is encountered, the extractor is rotated 90° by the handle, to bring the hook 68 into engagement with the back of the brim 12 at the anterior vaginal location. The extractor is pulled out, bringing the device 8 with it.

Obviously, many modifications and variations of the present invention are possible in light of the above teachings, and it is therefore understood that within the scope of the disclosed inventive concept, the invention may be practiced other than as specifically described.

Claims

1. A barrier-type device (8) for preventing conception and infection by sexually-transmitted diseases, comprising:

   a cervical dome (10) fabricated from a flexible, physiologically non-reactive material, having an opening and a curved surface extending from said opening to a dome top (16); a continuous fornical rim (12) on said dome, said rim defining said opening; and a continuous annular vaginal brim (14) circumscribing said dome (10), being formed as a backward fold from said rim (12) and extending downwardly and outwardly with respect to said opening; characterised in that

   the vaginal brim (14) has an outward concave bias or curve; and said vaginal brim (14) exhibits lateral symmetry and has a vertical dimension between its outside edge and said rim (12) which increases continuously with circumferential symmetry from an anterior extension (14c) to a posterior extension (14d) of said brim, said anterior extension (14c) being diametrically opposite said posterior extension (14d).

2. The barrier-type device of Claim 1 further including an annular lip (24) on said rim (12), said annular lip (24) extending inwardly from said rim (12) toward a centerline of said dome, and an annular groove (27) between said vaginal brim (12) and said dome, said annular groove (27) being formed by an outward fold of said vaginal brim (12) at said annular lip (24).

3. A kit for intravaginal, barrier-type prevention of conception and sexually-transmitted infection, comprising, in combination:

   a barrier-type device (8) according to Claim 1; and

   an applicator (50) including:

   a lower body with a main member (52) and a projection (54) attached at an angle to said main member, said projection including a concave recess (55) and an edge (56) for engaging said barrier-type device between said vaginal brim (12) and said dome (10); and an inserter (57), slidably disposed in said main member for being moved to a first, retracted position for engaging said vaginal brim (12) to a second, extended position for disengaging said vaginal brim (12).

4. The barrier-type device of Claim 1, wherein said forncial rim (12) forms an inwardly extending lip (24) and a fluid trapping, inwardly extending groove (27) between said brim (14) and said lip (24).

5. The barrier-type device of Claim 4, wherein the radial distance between said dome (10) and said brim (14) decreases upwardly with respect to the dome peak and towards the dome opening to a minimum distance just below said rim (12).

6. A barrier-type device according to Claim 1, wherein said fornical rim (12), which is annular and folded, defines fluid trapping means extending radially inwardly of the dome wall, said rim (12) connecting said dome means (10) with said brim means (14) to form a portion of said barrier-type device such that said fluid trapping means is a generally annular groove-shaped cavity in communication with said brim means (14) acting as a scoop and being for trapping fluids entering said brim means (14) acting as a scoop.

7. The barrier-type device of Claim 6, wherein said dome means (10) and said brim means (14) mutually define an annular passageway between said brim means acting as a scoop and said annular groove-shaped cavity (27), said passageway having a radial width less than the greatest radial width of said cavity.
8. The barrier-type device of Claim 7, wherein said fluid trapping means extends radially inwardly of said brim means (14) acting as a scoop.

**Patentansprüche**

1. Barriereartige Vorrichtung (8) zur Verhütung von Empfängnis und Infektionen durch sexuell übertragbare Krankheiten, aufweisend eine zervikale Kappe (10), die hergestellt ist aus einem flexiblen, physiologisch nicht reaktiven Material mit einer Öffnung und einer gekrümmten Oberfläche, die sich von der Öffnung bis zum Scheitel (16) der Kappe erstreckt; eine durchgehende fornikale Randzone (12) der Kappe, die die Öffnung definiert; und einen durchgehenden ringförmigen vaginalen Rand (14), der die Kappe (10) umgibt und der als Zurückfal tung der Randzone (12) geformt ist und sich bezüglich der Öffnung abwärts und nach außen gerichtet erstreckt, **dadurch gekennzeichnet**, daß der vagonale Rand (14) eine nach außen gerichtete kon kave Steigung oder Krümmung aufweist, daß der vaginale Rand (14) eine laterale Symmetrie auf weist und eine vertikale Ausdehnung zwischen der außenliegenden Kante und dem Rand (14) auf weist, welche kontinuierlich mit der kreisumfänglichen Symmetrie von einer vorderen Ausdehnung (14c) zu einer rückwärtigen Ausdehnung (14d) des Randes zunimmt und daß die vordere Ausdehnung (14c) diametral entgegengesetzt zu der rückwär tigen Ausdehnung (14d) angeordnet ist.

2. Barriereartige Vorrichtungen nach Anspruch 1, **dadurch gekennzeichnet**, daß des weiteren eine ringförmige Lippe (24) an der Randzone (12) vorge sehen ist, daß die ringförmige Lippe (24) sich von der Randzone (12) nach innen in Richtung auf eine Mittellinie der Kappe erstreckt, und daß eine ringförmige Ausdehnung (27) zwischen dem vaginalen Rand (14) und der Kappe vorgesehen ist, wobei die ringförmige Ausdehnung (27) durch eine Auswechs elfung des vaginalen Randes (14) an der ringförmigen Lippe (24) geformt ist.

3. Ausrüstung zur intravaginalen barriereartigen Ver hütung von Empfängnis und sexuell übertragbaren Infektionen, enthaltend in Kombination eine bar riereartige Vorrichtung (8) nach Anspruch 1 und einen Applikator (50) aufweisend, einen unteren Körper mit einem Hauptteil (52) und einen in einem Winkel zu dem Hauptteil angeordneten Vorsprung (54), wobei der Vorsprung eine konkave Ausneh mung (55) und eine Kante (56) zum Angriff an die barriereartige Vorrichtung zwischen dem vagina len Rand (14) und der Kappe (10) aufweist, und eine Einführenrichtung (57), die gleitbar in dem Hauptteil angeordnet ist, um in eine erste zurückge zogene Position zum Angriff auf den vaginalen Rand (14) und in eine zweite vorgestreckte Position zur Entkoppelung mit dem vaginalen Rand (14) bewegbar ist.

4. Barriereartige Vorrichtung nach Anspruch 1, **dadurch gekennzeichnet**, daß die fornikale Rand zone (12) eine sich nach innen hin ausdehnende Lippe (24) und eine flüssigkeitsauffangende, sich einräts erstreckende Ausdehnung (27), die zwischen dem Rand (14) und der Lippe (24) angeord net ist, bildet.


6. Barriereartige Vorrichtung nach Anspruch 1, **dadurch gekennzeichnet**, daß die fornikale Rand zone (12), welche ringförmig und gefaltet ausgebil det ist, flüssigkeitsauffangende Mittel definiert, die sich radial einräts von der Kappenwandung erstrecken, wobei die Randzone (12) die Kappe (10) mit dem Rand (14) verbindet, um einen Teil der barriereartigen Vorrichtung zu bilden, so daß das flüssigkeitsauffangende Mittel eine im wesentlichen ringförmige und nutenartig geformte Kavität dar stellt, die in Verbindung mit dem Rand (14) steht und als Aufnehmer wirkt und für den Rand (14) erreichen aufgefangene Flüssigkeiten als Auf nehmer wirkt.

7. Barriereartige Vorrichtung nach Anspruch 6, **dadurch gekennzeichnet**, daß die Kappe (10) und der Rand (14) gegenseitig einen ringförmigen Durchgang zwischen dem als Aufnehmer wirken den Rand und der ringförmigen, nutenartig geform ten Kavität (27) definieren, wobei der Durchgang eine radiale Weite aufweist, die kleiner ist als die größte radiale Weite der Kavität.

8. Barriereartige Vorrichtung nach Anspruch 7, **dadurch gekennzeichnet**, daß die flüssigkeitsauf fangenden Mittel sich von dem Rand (14), der als Aufnehmer dient, radial einräts erstrecken.

**Revendications**

1. Dispositif contraceptif du type barrière (8) et de pré vention contre l'infection par les maladies sexuelle ment transmissibles, comprenant : un dôme cervical (10) fabriqué dans un maté riau flexible, biocompatible présentant une ouverture et une surface curviligne s'étendant de cette ouverture jusqu'au sommet (16) du dôme ; un rebord de fornix (12) continu et replié
sur ce dôme, ce rebord définissant l’ouverture ; et un bord vaginal (14) continu et annulaire entourant le dôme (10), conformé selon une pliure arrière dans le rebord replié (12) et s’étendant vers le bas et vers l’extérieur par rapport à l’ouverture ; caractérisé en ce que : le bord vaginal (14) affecte une forme concave inclinée ou incurvée vers l’extérieur et présente une symétrie latérale et une dimension verticale entre son bord extérieur et le rebord replié (12) en augmentation continue avec symétrie périmétrique à partir d’une extension antérieure (14c) jusqu’à une extension postérieure (14d) du bord, l’extension antérieure (14c) étant diamétralement opposée à l’extension postérieure (14d).

2. Dispositif de type barrière selon la revendication 1, comprenant de plus une lèvre annulaire (24) sur le rebord replié (12), cette lèvre annulaire (24) s’étendant vers l’intérieur à partir du rebord (12) vers un axe médian du dôme, et une rainure (27) entre le bord vaginal (14) et le dôme, cette rainure annulaire (27) étant formée selon une pliure s’étendant vers l’extérieur du bord (12) au niveau de la lèvre annulaire (24).

3. Ensemble de prévention intravaginale de type barrière contre la conception et les infections sexuellement transmissibles, comprenant en combinaison :

   un dispositif du type barrière (8) selon la revendication 1 ; et
   un applicateur (50) comprenant :

   un corps inférieur comportant une pièce principale (52) et une prolongation (54) rapportée sur cette pièce selon un certain angle, la prolongation comportant un évidement concave (55) et une arête (56) pour l’engagement du dispositif de type barrière entre le bord replié vaginal (14) et le dôme (10) ; et une pièce d’insertion (57) montée à coulissement dans la pièce principale pour être déplacée d’une première position rétractée pour l’engagement du bord vaginal (14) dans une deuxième position en extension pour le désengagement du bord vaginal (14).

4. Dispositif du type barrière selon la revendication 1, dans lequel le bord replié de fornix (12) forme une lèvre (24) s’étendant vers l’intérieur et une rainure (27) de piégeage du fluide s’étendant vers l’extérieur entre le bord (14) et la lèvre (24).

5. Dispositif du type barrière selon la revendication 4, dans lequel la distance radiale entre le dôme (10) et le bord (14) diminue vers le haut par rapport au sommet du dôme et vers l’ouverture du dôme pour atteindre une distance minimale juste en-dessous du bord replié (12).

6. Dispositif du type barrière selon la revendication 1, dans lequel le bord de fornix (12), annulaire et replié, détermine un moyen de piégeage de fluide s’étendant radialement vers l’extérieur de la paroi du dôme, le bord replié (12) reliant le dôme (10) au bord (14) pour former une partie du dispositif du type barrière telle que le moyen de piégeage de fluide est généralement une cavité annulaire formée en rainure communiquant avec le bord (14) faisant office de collecteur aux fins de piéger les liquides entrant dans le bord (14) à fonction de collecteur.

7. Dispositif du type barrière selon la revendication 6, dans lequel le dôme (10) et le bord (14) définissent mutuellement un passage annulaire entre le bord collecteur et la cavité (27) annulaire en forme de rainure, ce passage ayant une largeur radiale inférieure à la plus grande largeur radiale de la cavité.

8. Dispositif de type barrière selon la revendication 7, dans lequel le moyen de piégeage de liquide s’étend radialement vers l’intérieur du bord collecteur (14).