DISPOSABLE DEVICE FOR ANALYSIS OF A BODY FLUID

The present invention relates to the analysis of a body fluid taken directly from an elongate intracorporeal cavity. By way of a nonlimiting example, the present invention will be introduced, defined and explained with reference to the recovery of the cervical mucus in the vaginal cavity of women for detection of the period of fertility.

In order to determine the periods of fertility in women, it is known, and it has been proposed, to detect and monitor the presence and/or concentration of certain biochemical or biological constituents of said mucus, such as a peroxidase or a compound having a peroxidase activity, or such as a mucopolysaccharide or a glycoprotein, by using reagents or colored reagent systems, for example, in the former case, an oxidation-reduction compound, of which at least the oxidized form is colored, for example guaiacol, and in the latter case safranin.

The solutions which have been proposed for the use of such reagents or reactive systems have generally been rudimentary and have in practice been difficult for women to put into use, namely:

- taking a sample of the cervical mucus in situ, with an element capable of recovering the mucus, such as a swab, then bringing said element into contact, outside the vaginal cavity, with the reagent in liquid form;
- introducing into the vaginal cavity an absorbent tampon impregnated or coated with a layer of reagent, and withdrawing said tampon after a certain time, and treating the layer of reagent with a color-developing agent.

These methods are known, for example, from the documents FR-A-2,216,975, EP-A-0,363,196 and EP-A-0,555,109.

The disadvantages of these methods result from the need to treat the extract of cervical mucus

recovered extemporaneously, in order to reveal the possible presence or concentration of certain of its biochemical constituents.

Furthermore, in order to culture microorganisms directly, in particular pathogens, which may be present in an elongate intracorporeal cavity, a disposable device according to the embodiment in Figure 3 of the document US-C-4,257,427 has been proposed, said device comprising:

- a conduit element, made of transparent plastic, surrounded by a sheath of flexible foam; this element has, at its two ends, a relatively narrow neck and a base; it is inherently sufficiently stiff or rigid along its length for it to be pushed via its base and introduced into the intracorporeal cavity;
- a means for transferring the body fluid assumed to contain the microorganisms, said means being mounted at least in part in the interior of the conduit element, extending from a zone for sampling of the body fluid, outside the neck, to an axial collection point for said fluid, situated inside the conduit element, toward its base; this means consists of a wick of braided filaments, the outer unravelled end of which serves for taking the samples, and the inner end of which concentrates the body fluid which has been collected; an intermediate tube ensures sealing between the wick and the neck of the conduit element in such a way that the interior of the latter is substantially isolated from the intracorporeal cavity in which said element has been introduced;
- a culture medium arranged inside the conduit element, against its base.

This device is introduced by the user into the intracorporeal cavity, in which it remains for a period of time sufficient to permit the recovery of the body

fluid, and the culturing of the microorganisms sought. The user then removes the device from the intracorporeal cavity and observes the development, or lack of development, of the culture of the microorganism sought, through the wall of the conduit element.

The subject of the present invention is a disposable device permitting extemporaneous analysis of a body fluid, with results which can be identified directly by the user, and without any intervention by the latter, other than that of introducing and removing said device into and from the intracorporeal cavity.

In accordance with the present invention, the starting point is a disposable device such as is described in the document US-C-4,257,427, which device is then modified specifically in the following manner:

- on the one hand, the sampling zone of the transfer means consists of a peripheral ring, supported in the aperture of the neck of the conduit element, protruding radially and to the outside of said neck, and forming a passage for communication between the intracorporeal cavity and the inside of said conduit element;
- and, on the other hand, a reaction means is mounted or arranged on the conduit element, in a manner visible to the user, and in relation with the axial collection point, in order to receive at least some of the body fluid which has been collected, and it comprises at least one reagent capable of reacting with at least one component of the body fluid, or in the presence of a biological or biochemical state of said fluid, in order to give a reaction product, preferably colored, revealing the presence of said biological or biochemical state, or of said component in said body fluid, and directly identifiable by the user.

By virtue of the present invention, the recovery in particular of the body fluid remains

independent of the differential pressure conditions between the intracorporeal cavity and the inside of the conduit element, this fact facilitating the transfer of the body fluid toward the reaction means. This advantage is decisive when the body fluid, for example the cervical mucus, is more or less liquid, and/or more or less abundant.

The conduit element 2 is preferably adapted in shape and dimensions to be fitted and held directly, by simple construction [sic], in the intracorporeal cavity, and its outer surface is chosen to be biocompatible with the inner wall of the intracorporeal cavity.

The term "biocompatible" refers to the fact that the contact between the outer surface of the conduit element and the wall of the intracorporeal cavity does not generate any adverse biological reaction, for example of the toxic or allergic type.

The means for transferring the body fluid advantageously comprises a bundle of filiform elements which are gathered together at one end at the axial collection point and are spread out at the other end about the peripheral sampling ring, each filiform element being capable of conveying the body fluid from an outer end to an inner end, for example by absorption or capillarity.

The filiform elements are preferably hollow filaments, or else comprise fibers of at least one biocompatible polymer, in particular a polymer chosen from among the hydrophilic natural or synthetic polymers, such as polyesters, polyacrylonitriles, polycarbonates, polyethylenes and polypropylenes, silicones, alginates, polyurethane foams, or else cellulose or its esters, optionally impregnated with an agent liquefying the body fluid.

The reaction means is preferably arranged inside the conduit element, at its closed base, facing the axial collection point of the transfer means.

In this case, the base of the conduit element is transparent, and the reagent is arranged against this base, and in fact faces the axial collection point of the transfer means, in such a way that when the reaction product is colored, this color or absence of color can be seen or visualized by the user.

In a variant according to the invention, the base is at least partially open, and a stem for transfer of the body fluid, from the axial collection point to a distal element for recovery of said fluid, passes through the base via its aperture and emerges from the conduit element.

In this case, the transfer stem preferably consists of the same filiform elements as for the transfer means, which may optionally continue through the base of the conduit element, and the reaction means can be the same or different, depending on whether the aim is to demonstrate a biological component or biological state, or two different biological components or states.

This variant at [sic] the advantage of indicating the presence or absence of a biological state or component of the body fluid without the user being obliged to withdraw the device from the intracorporeal cavity.

The analysis device according to the invention, as it has been described, can therefore be used directly, in particular without subsequent addition of color reagent or developer by the user, and in this latter case the color reaction is observed immediately after the removal of the device.

The present invention is now described with reference to the attached drawing, in which:

- Figure 1 represents a cross section of a disposable analysis device according to the invention;
- Figure 2 represents an assembly ready for use, comprising a device according to Figure 1 together

with an applicator system;

- Figure 3 represents a cross section of a disposable device according to another variant of the invention.

In a general manner, and as illustrated in Figure 1, a disposable analysis device 1 for recovery of body fluid, for example cervical mucus, directly from an elongate intracorporeal cavity, for example the vaginal cavity, comprises:

- a conduit element 2 adapted in shape and dimensions to be fitted directly in, and held by simple constriction of, the intracorporeal cavity; this element has at its two ends, respectively, a relatively wide neck 2a and a base 2b having the same cross section; this element is inherently sufficiently stiff or rigid along its length to be pushed via its base 2b, and introduced via its neck 2a, into the intracorporeal cavity; and its outer surface is chosen to be biocompatible with the inner wall of the intracorporeal cavity;
- a means 3 for transferring the body fluid, said means being mounted at least in part in the interior of the conduit element 2, and extending from a peripheral ring or zone 3a for sampling of the body fluid, protruding radially and to the outside of the neck 2a of said conduit element 2, to an axial collection point 3b for said fluid, situated inside said conduit element 2, toward the base 2b thereof. The ring 3a is supported in the aperture of the neck 2 [sic] of the conduit element 2 and forms a passage, for example axial, for communication between the intracorporeal cavity and the inside of the conduit element.

The conduit element 2 can have a composite structure and can comprise a relatively rigid tube 5, for example made of transparent plastic material, with a neck 5a and a base 5b on which is fitted, if appropriate, a biocompatible sleeve 6 surrounding at

least the side wall of the tube, consisting for example of an outer tube 6a made of compounds of natural origin, such as cellulose or cotton, and a rigid inner matrix 6b made of polymer as a support for said second tube, whose anterior edge (in the direction of introduction of the device) is rounded, and whose posterior edge is straight.

The transfer means 3 is mounted at least in part in the conduit element 2; preferably however, and as is illustrated in Figure 1, the transfer means 3 is mounted on a support means 4 which is removable or fixed in relation to the conduit element 2. More specifically, the support means 4 has a funnel shape and is mounted on the rigid tube 5 by means of a screw pitch 5e and a shoulder 5d which are provided on the tube 5 at the level of its neck 5a.

The transfer means 3 comprises a bundle of filiform elements 3c, preferably hollow and hydrophilic filaments, which are gathered together at one end at the axial collection point 3b, and are spread out at the other end about the peripheral sampling ring 3a, each filiform element 3c ensuring the transfer of the body fluid, which is to be recovered, from the outside of the device toward a reaction means, which will be described hereinafter, for example by absorption or capillarity.

These filiform elements 3c can be replaced by other elements ensuring a capillary migration, or a migration by surface tension, of the fluid collected. However, the particular advantage of using the filaments or fibers is that they bend and rub against the intracorporeal wall of the cavity via the peripheral ring 3a of the transfer means 3 upon insertion of the device into said cavity, thereby recovering more fluid than is possible by simple gravitational flow.

Hydrophilic fibers are preferably chosen because these convey the cervical mucus, which consists

for the most part of water, more rapidly toward the reaction means, and in particular fibers made of hydrophilic natural or synthetic biocompatible polymers, such as polyesters, polyacrylonitriles, polycarbonates, polyethylenes and polypropylenes, silicones, alginates, polyurethane foams, or else cellulose or its esters. These fibers can be impregnated with a fluid liquefying, or aiding the transport of, the body fluid.

A removal line 15 in the form of a string or cord is fixed to the conduit element 2, on the side of its base 2b, by way of a groove 5c provided in the tube 5.

A reaction means or system 9 is arranged inside the conduit element 2, at its base 2b, facing or in relation with the axial collection point 3b of the transfer means 3, in order to receive some of the body fluid which has been collected. This reaction means incorporates at least one reagent, and if appropriate a liquefying agent for the cervical mucus. This reaction means or system is deposited for example either directly or on an absorbent layer 12. It is capable of reacting with at least one component of the body fluid which has been collected, in order to give at least one color reaction product, which can be directly identified by the user, revealing the presence of said component in the body fluid. As has already been mentioned, the color reagent on its absorbent layer 12 is arranged against the base 2b, facing the axial collection point 3b of the transfer means 3, and more precisely against the transparent base wall 10 of the tube 5. The layer 12 impregnated with the reagent 9 is covered by a web 11 which is semipermeable, in the sense that it is permeable with respect to the body fluid in the direction of transfer, while it holds back this same fluid in the other direction. This web can also be impregnated with a liquefying agent for the cervical mucus, promoting the passage of the compounds

which are to be detected and being, for example, a surface-active agent.

In a general manner, although not represented in Figure 1, the shape and the dimensions of the conduit element 2 are adapted to those of the vaginal cavity.

In accordance with Figure 2, the device described above can form part of a ready-to-use assembly comprising this same device 1 and an applicator system 16. The latter comprises a guide tube 13, inside which the recovery device 1 is inserted. A pusher 14 is housed inside the guide tube, abutting against the analysis device. Such an assembly can be introduced directly into the intracorporeal cavity, for example the vaginal cavity, and by pushing the pusher 14 the analysis device 1 is released from the applicator system, in order to position said device in this same cavity.

Figure 3 shows a variant of the invention, in accordance with which the device is essentially the same as that described above, but with the difference that the base 2b is equipped with an aperture, and a stem 17 for transfer of the body fluid passes through said aperture and emerges from the conduit element 2 toward a distal element 20 for recovery of said fluid, this element 20 being equipped with a reaction means 19.

This reaction means can be the same as, or different from, that which has already been described, depending on whether the aim is to reveal the presence of one or more biological components or biological states.

The stem 17, which can be a sheath of rigid or flexible polyethylene or similar, also comprises filiform elements 18 on the inside, these either being the same as for the device illustrated in Figure 1, or else different, the proviso being that they guarantee the transfer of the body fluid from the axial

collection point 3b toward the reaction means 19 of the distal element 20.

The analysis devices described above can therefore be used not only for the detection of a component in a body fluid, but also for detecting any chemical, biochemical or biological state of this same body fluid, for diagnostic, prophylactic or therapeutic purposes. This means that this analysis device can incorporate very different reagents or reaction systems, of a purely chemical type, for example enzymatic, or biological, for example an antigen or an antibody, or else immunoenzymatic.

Consequently, an analysis device according to the invention can have very broad applications, among which there may be mentioned:

- the detection of a hormone, and in particular of a hormonal peak;
- the sampling and the histological and/or cyto-chemical analysis of a body fluid, in particular for detecting pathological conditions, or for demonstrating certain physiological phases of a natural cycle, for example of a hormonal cycle.

The following reagents will be mentioned in particular, by way of example, for the detection of a glycoprotein or of a mucopolysaccharide, and consequently of the fertile period in women: safranin, toluidine blue O, Alcian blue, trypan blue, a tolonium salt, PAS (periodic acid-Schiff), alkaline phosphatase or a mixture of these, if appropriate combined with an agent promoting the color reaction, such as polyvinylpyrrolidone.

CLAIMS

1. A disposable device (1) for analysis of a body fluid present in an elongate intracorporeal cavity, comprising:

- a conduit element (2) having respectively, at its two ends, a neck (2a) and a base (2b), inherently sufficiently stiff or rigid along its length for it to be pushed via its base (2b) and introduced into the intracorporeal cavity;
- a means (3) for transferring the body fluid, said means being mounted at least in part in the interior of the conduit element, extending in [sic] a zone (3a) for sampling of the body fluid, outside the neck (2a), to an axial collection point (3b) for said fluid, situated inside said conduit element (2), toward its base (2b);

wherein, on the one hand, the sampling zone (3a) of the transfer means (3) consists of a peripheral ring, supported in the aperture of the neck (2) [sic] of the conduit element (2), protruding radially and to the outside of said neck, and forming a passage for communication between the intracorporeal cavity and the interior of said conduit element, and, on the other hand, a reaction means (9, 19) is mounted on the conduit element, in a manner visible to the user, and in relation with the axial collection point (3b), in order to receive at least some of the body fluid which has been collected, and it comprises at least one reagent capable of reacting with at least one component of the body fluid, or in the presence of a biological or biochemical state of said fluid, in order to give at least one reaction product, preferably colored, revealing the presence of said biological or biochemical state, or of said component in said body fluid, and directly identifiable by the user.

2. The device as claimed in claim 1, wherein the conduit element (2) is adapted in shape and dimensions to be fitted and held directly, by simple construction

[sic], in the intracorporeal cavity, and its outer surface is chosen to be biocompatible with the inner wall of the intracorporeal cavity.

3. The device as claimed in claim 1, wherein the transfer means (3) comprises a bundle of filiform elements (3c) which are gathered together at one end at the axial collection point (3b) and are spread out at the other end about the peripheral sampling ring (3a), each filiform element being capable of conveying the body fluid from an outer end to an inner end, for example by absorption or capillarity.

4. The device as claimed in claim 3, wherein the filiform elements (3c) are hollow filaments.

5. The device as claimed in claim 3, wherein the filiform elements (3c) comprise fibers of at least one biocompatible polymer, in particular a polymer chosen from among the hydrophilic natural or synthetic polymers, in particular polyesters, polyacrylonitriles, alginates, polycarbonates, polyethylenes and polypropylenes, silicones, polyurethane foams, or else cellulose or its esters, optionally impregnated with an agent liquefying the body fluid.

6. The device as claimed in claim 1, wherein the reaction means (9) is arranged inside the conduit element (2), at its closed base (2b), facing the axial collection point (3b) of the transfer means (3).

7. The device as claimed in claim 6, wherein the reaction product is colored, the base (2b) is transparent (10), and the reagent (9) is arranged in a relatively thin layer against the base (2b), facing the axial collection point (3b) of the transfer means (3).

8. The device as claimed in claim 7, wherein the layer of reagent (9) is covered by a web (11) which is semipermeable, in the sense that it is permeable with respect to the body fluid in the direction of transfer, while it holds back said fluid in the other direction, and wherein the web or the layer or both are optionally impregnated with a liquefying agent promoting the

passage of the components which are to be detected.

9. The device as claimed in any one of claims 1 to 5, wherein the base (2b) is at least partially open, and wherein a stem (17) for transfer of the body fluid, from the axial collection point (3b) to a distal element (20) for recovery of said fluid, passes through the base via its aperture and emerges from the conduit element (2).

10. The device as claimed in claim 9, wherein the reaction means (19) is associated with the distal element (20) for recovery of the body fluid.

11. The device as claimed in claim 1, wherein the conduit element (2) comprises a relatively rigid tube (5) with a neck (5a) and a base (5b), and if appropriate a biocompatible sleeve (6) surrounding at least the side wall of said tube.

12. The device as claimed in claim 1, wherein the transfer means (3) is mounted on a support means (4) which is removable or fixed in relation to the conduit element (2).

13. An assembly ready for use, comprising a device (1) as claimed in claim 1 and an applicator system (16).

SG9701094-6

PCT ORGANISATION MONDIALE DE LA PROPRIETE INTELLECTUELLE
Bureau international

DEMANDE INTERNATIONALE PUBLIEE EN VERTU DU TRAITE DE COOPERATION EN MATIERE DE BREVETS (PCT)

(51) Classification internationale des brevets 6 : G01N 33/52, A61B 10/00, C12Q 1/28 — A1

(11) Numéro de publication internationale: WO 96/09544

(43) Date de publication internationale: 28 mars 1996 (28.03.96)

(21) Numéro de la demande internationale: PCT/FR95/01199

(22) Date de dépôt international: 19 septembre 1995 (19.09.95)

(30) Données relatives à la priorité:
94/11349 19 septembre 1994 (19.09.94) FR

(71)(72) Déposant et inventeur: CHAFFRINGEON, Bernard [FR/FR]; La Logère, F-69480 Anse (FR).

(74) Mandataire: CABINET GERMAIN & MAUREAU; Boîte postale 3011, F-69392 Lyon Cédex 03 (FR).

(81) Etats désignés: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TT, UA, UG, US, UZ, VN, brevet européen (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), brevet OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), brevet ARIPO (KE, MW, SD, SZ, UG).

Publiée

Avec rapport de recherche internationale.
Avant l'expiration du délai prévu pour la modification des revendications, sera republiée si de telles modifications sont reçues.

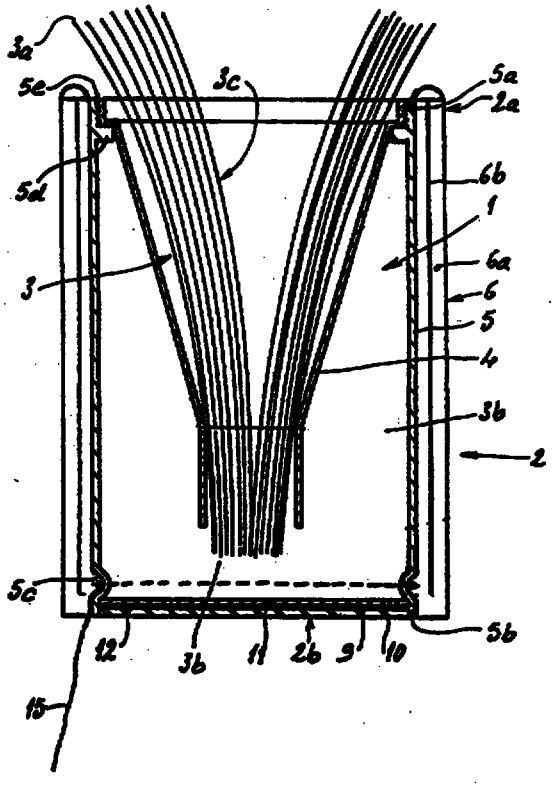
RTMP copy

(54) Title: SINGLE-USE BODY FLUID ANALYSIS DEVICE

(54) Titre: DISPOSITIF A USAGE UNIQUE D'ANALYSE D'UN LIQUIDE CORPOREL

(57) Abstract

A single-use device (1) for analysing a body fluid in an elongate body cavity, including a pipe element (2) with a neck (2a) and a bottom wall (2b), and a liquid transfer assembly (3) partially located within the element (2) and extending through a sampling region (3a) outside the neck (2a) to an axial collection point (3b) located within said element (2) in the region of the bottom wall (2b) thereof. The sampling region (3a) of the transfer assembly (3) consists of a peripheral ring held in the neck opening (2) and projecting radially outwards therefrom to define a communicating passage between the cavity and the inside of said element, and a user-visible reaction portion (9, 19) fitted on the element (2) and linked to the axial collection point (3b) for receiving the collected body fluid includes at least one reagent capable of reacting with one component of the fluid to provide a directly user-identifiable tinted reaction product that indicates the presence of said component.



(57) Abrégé

Dispositif (1) à usage unique d'analyse d'un liquide corporel présent dans une cavité intracorporelle allongée, comprenant: un élément (2) de conduit, présentant un col (2a) et un fond (2b); un moyen de transfert (3) du liquide, monté pour partie à l'intérieur de l'élément (2) s'étendant dans une zone (3a) de prélèvement, à l'extérieur du col (2a), à un point (3b) de collecte axiale, situé à l'intérieur dudit élément (2), du côté de son fond (2b). La zone (3a) de prélèvement du moyen de transfert (3) consiste en une couronne périphérique, supportée dans l'ouverture du col (2), saillant radialement et à l'extérieur dudit col, et ménageant un passage de communication entre la cavité et l'intérieur dudit élément, et d'autre part un moyen réactionnel (9, 19) est monté sur l'élément (2) de manière visible pour l'utilisateur, en relation avec le point (3b) de collecte axiale, pour recevoir le liquide corporel collecté, et comprend au moins un réactif susceptible de réagir avec un composant du liquide, pour donner un produit de réaction coloré, révélant la présence dudit composant, et identifiable directement par l'utilisateur.

UNIQUEMENT A TITRE D'INFORMATION

Codes utilisés pour identifier les Etats parties au PCT, sur les pages de couverture des brochures publiant des demandes internationales en vertu du PCT.

AT	Autriche	GB	Royaume-Uni	MR	Mauritanie
AU	Australie	GE	Géorgie	MW	Malawi
BB	Barbade	GN	Guinée	NE	Niger
BE	Belgique	GR	Grèce	NL	Pays-Bas
BF	Burkina Faso	HU	Hongrie	NO	Norvège
BG	Bulgarie	IE	Irlande	NZ	Nouvelle-Zélande
BJ	Bénin	IT	Italie	PL	Pologne
BR	Brésil	JP	Japon	PT	Portugal
BY	Bélarus	KE	Kenya	RO	Roumanie
CA	Canada	KG	Kirghizistan	RU	Fédération de Russie
CF	République centrafricaine	KP	République populaire démocratique de Corée	SD	Soudan
CG	Congo			SE	Suède
CH	Suisse	KR	République de Corée	SI	Slovénie
CI	Côte d'Ivoire	KZ	Kazakhstan	SK	Slovaquie
CM	Cameroun	LI	Liechtenstein	SN	Sénégal
CN	Chine	LK	Sri Lanka	TD	Tchad
CS	Tchécoslovaquie	LU	Luxembourg	TG	Togo
CZ	République tchèque	LV	Lettonie	TJ	Tadjikistan
DE	Allemagne	MC	Monaco	TT	Trinité-et-Tobago
DK	Danemark	MD	République de Moldova	UA	Ukraine
ES	Espagne	MG	Madagascar	US	Etats-Unis d'Amérique
FI	Finlande	ML	Mali	UZ	Ouzbékistan
FR	France	MN	Mongolie	VN	Viet Nam
GA	Gabon				

9701094-6

FIG 1

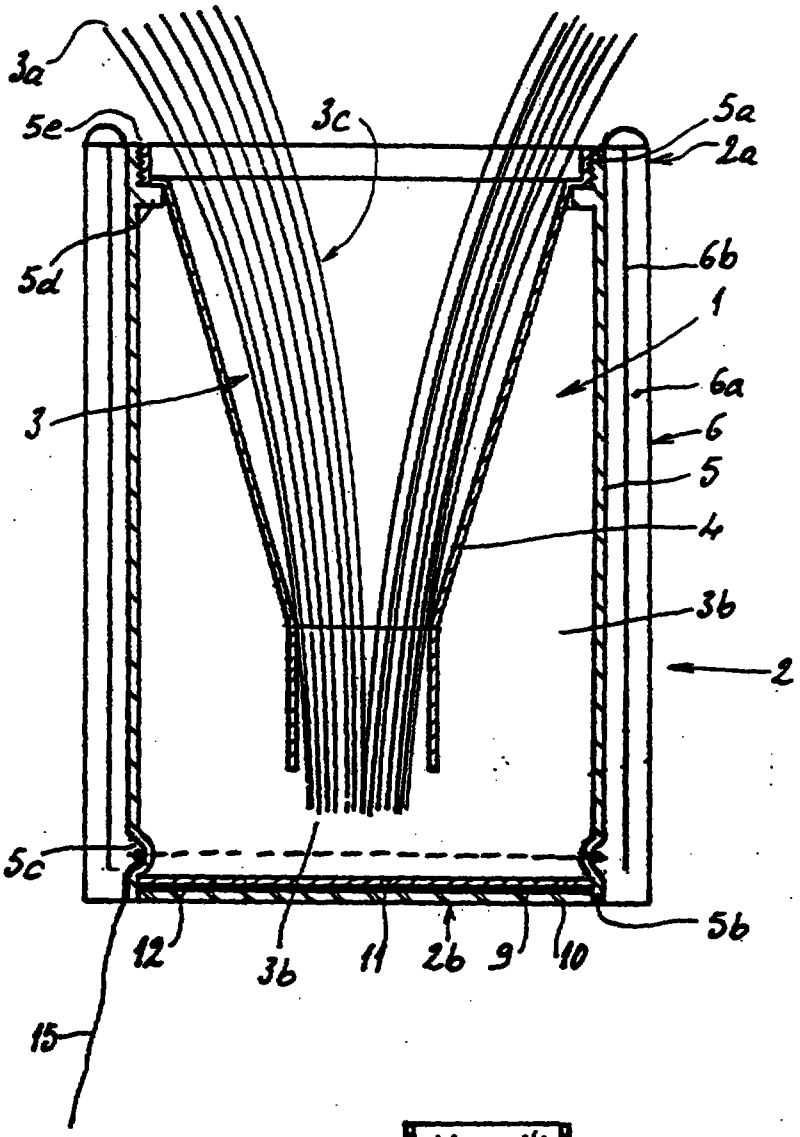


FIG 2

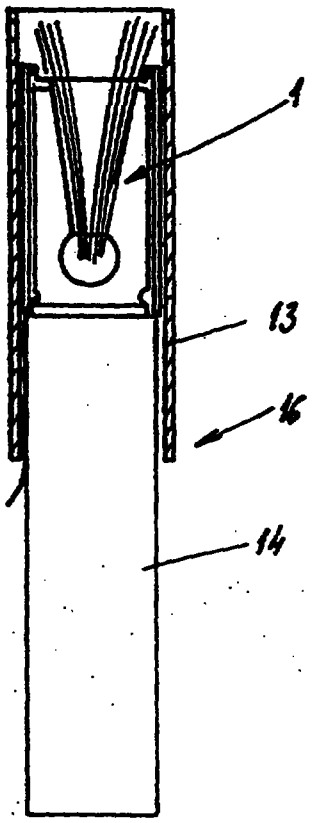


FIG 3

