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628563

NOTICE OF ENTITLEMENT

We, **Toray Industries, Inc.** the applicant/nominated person in respect of Application No. 55669/90, state the following:-

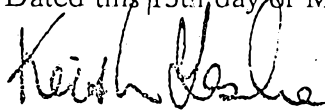
Noriyasu NOGUCHI, Yoshiharu YAMAZAKI, Mitsuyuki HAGIO and Yasuhiko FUTAMI are the inventors.

The Nominated person is entitled to the grant of the patent because the Nominated person derives title to the invention from the inventors.

The Nominated person is entitled to claim priority from the basic application listed on the patent request because the Nominated person made the basic application.

The basic application is the first application made in a Convention country in respect of the invention.

Dated this 13th day of March 1992



(A member of the firm of DAVIES COLLISON CAVE for
and on behalf of the applicant)

PATENT, TRADE MARKS
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13 MAR 1992
MELBOURNE

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(57) The term free length
refers to the length of the fibre when in the form of a straight line under no
tension.

CLAIM

1. A catheter with a balloon, said balloon being reinforced with a composite yarn adhered to or embedded in the balloon, said composite yarn comprising an elastic yarn and a non-elastic yarn with a larger free length than that of said elastic yarn.

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(81) 指定国

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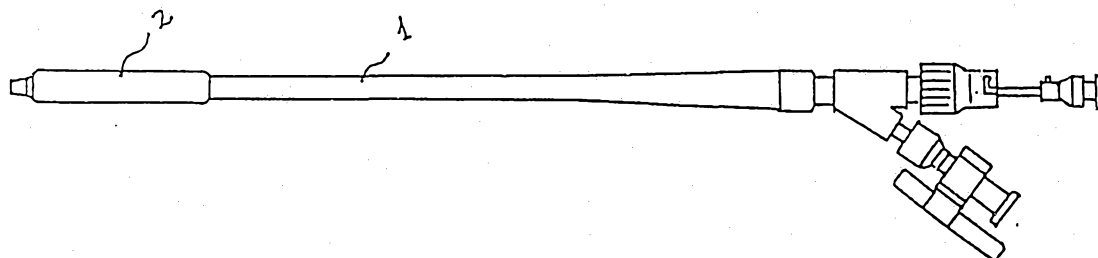
NL(欧州特許), SE(欧州特許), US.

(54) Title: ~~CATHETER-CUM-BALLOON~~

A CATHETER WITH A BALLOON

(54) 発明の名称

バルーン付カテーテル



(57) Abstract

This invention relates to a catheter-cum-balloon characterized in that the balloon is reinforced with composite yarns composed of elastic yarns and non-elastic ones which are each larger than the elastic one in free length. In the catheter-cum-balloon of the present invention, expandability of the balloon is high and thus, when inserted into or pulled out from the blood vessel for the purpose of therapy, the balloon is capable of not only decreasing in diameter without development of creases but also positively expanding to the same size and the same shape at any time. Accordingly, provided are such advantages that: reliability in therapy is increased; an opening incised for therapy, even if small, satisfies the purpose; treatment and care after operation are simple, requiring only a few length of time until cure; and the burden inflicted on the patient and nurse daily is light.



TECHNICAL FIELD

The present invention relates to a catheter with a balloon for expanding a constricted valve of the heart or
5 a constricted or occluded blood vessel and normalizing thereby blood flow in medical treatment.

BACKGROUND TECHNOLOGY

Previously, catheter balloons have been made of synthetic resins such as polyethylene, polyvinyl chloride
10 or polyester. These materials have no stretchability but do resist pressure and thereby prevent blood vessels from being damaged because of excessive expansion. However, when balloons of these materials are expanded, the surfaces thereof become slippery and cause the balloons
15 to slip out, or when they are shrunk, a lot of wrinkles are produced in their surface and they tend to rub the inner walls of the patient's blood vessels. This causes damage and accelerates the formation of thrombi.

To overcome the wrinkling problem it has been
20 proposed to manufacture the balloons from a polyurethane or a silicone.

Alternatively, it was proposed to prevent excessive expansion and the expansion of the balloon was made uniform by incorporating a knitted fabric into a balloon.

25 In addition, it was proposed to use rubber as the balloon raw material and to form the balloon in a guitar-shape by pinching the balloon with a knitted fabric with several stripes of bands. When trialled in a constricted valve of the heart the balloon did not slip out.

30 However, even when such balloons were used, the outer diameters became too large from their structural points and exposure and incision of a blood vessel was required when the balloon was inserted into them. Consequently it took longer for the patient to recover
35 from the procedure and was generally inconvenient.

It was also previously proposed to incorporate a knitted fabric in a balloon. This was found to be very



complicated work and comprised the steps of knitting a
parallelly-twisted yarn of a polyamide yarn and a
urethane yarn and using such material to cover the outer
surface of the balloon expanded to a required expanded
5 diameter. It was found that the diameter of the balloon
could not be made smaller as the balloon could not be
stretched in the longitudinal direction.

The object of the present invention is to alleviate
all or some of the above mentioned problems.

10 According to a first aspect of the present invention
there is provided a catheter with a balloon, said balloon
being reinforced with a composite yarn adhered to or
embedded in the balloon, said composite yarn comprising
an elastic yarn and a non-elastic yarn with a larger free
15 length than that of said elastic yarn.

According to a second aspect of the present
invention there is provided a catheter with a balloon,
said balloon being reinforced with a knitted or woven
fabric of a composite yarn adhered to or embedded in the
20 balloon, said composite yarn comprising an elastic yarn
and a non-elastic yarn with a larger free length than
that of said elastic yarn.

As the catheter with a balloon of an embodiment of
the present invention exhibits larger balloon
25 stretchability, it is possible to make the outer diameter
smaller for a balloon with large expansion diameter used
in a tearing and opening technique for a valve of the
heart etc., and the catheter with the balloon can be
inserted in a blood vessel by means of a simple
30 Celludinga method. Therefore, there exist such
advantages that a cut wound for surginal operation could
be smaller; treatment and management after the surginal
operation are simple and the time required for recovery
is extremely short; smaller daily burdens of a patient
35 and an attendant are needed.

In addition, when the catheter with a balloon of the
present embodiment is pushed into a blood vessel and



inner heart to a diseased part, an inserting operation putting a guide wire and an expanded balloon in the blood flow can be easily preformed and as the apex of the balloon is soft, there is few possibility of damaging the
5 inner wall of the blood vessel and the workability thereof is also good. Moreover, preparation of a cylindrical body made of a composite yarn and a cylindrical elastic film and assembling them for manufacturing the balloon are easy and the yield is good.
10 The cost can be lowered, too.

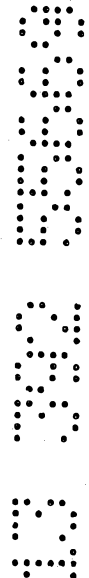
An embodiment of the invention will now be described by way of example only with reference to the accompanying drawings in which:-

Figure 1 is a rough figure of a catheter with a
15 balloon of an embodiment of the present invention and Figure 2 is a figure illustrating an expanded state of the balloon part of the catheter with the balloon in Figure 1.

- 20 1: A tube part
2: A balloon part

BEST EMBODIMENT FOR PRACTICING THE PRESENT INVENTION

As illustrated in Figure 1, in the catheter with a balloon 2 constituting a three-layered structure with a cylindrical body consisting of a cylindrical elastic film



and a composite yarn is attached on the apex of a tube 1 with a smooth surface and a round cross-section and on the rear end of the tube 1, if necessary, a fluid inlet and an take-up opening for an air removal tube placed in a fluid transferring path are provided. In the catheter with a balloon of the present invention, the ratio of the free length of an elastic yarn to a non-elastic yarn is preferably 0.15-0.5 and more preferably 0.2-0.35. The term free length refers to the length of the fibre when in the form of a straight line under no tension. If this value is too large, the balloon tends to be insufficiently expanded, and if this value is too small, a reinforcing effect by means of the non-elastic yarn is easily fluctuated and breakage of the balloon when expanded easily occurs. However, if a careful consideration is paid when manufacturing the balloon to avoid the breakage, even when said ratio is larger than the above described value, the effect of the present invention can be utilized. The ratio of the free length can be known by measuring the bending point of the stress-strain curve of said composite yarn to the higher tension side. When the free length of the non-elastic yarn is fluctuated and the bending point is therefore unclear, it can be known by extrapolating the straight lines before and behind the bending point.

As the structure of said composite yarn, a coresheath yarn wherein the core is made of an elastic yarn and the sheath is made of a non-elastic yarn is preferable



because both yarns are hardly separated. Namely, to make it easily stretch by an inner pressure or to preset the part order of stretching and the final stretched state, in some cases, forming a characteristic is performed by stretching and shrinking the yarns in advance and in these cases, there is the possibility that separation of both constituting yarns badly influences on the effect of the present invention.

As the method for manufacturing the core-sheath yarn, a method wherein twisting or entangling by means of a fluid is performed while a non-elastic yarn is overfed is preferable and a core-sheath yarn made by winding spirally a twisted yarn of a synthetic fiber on a urethane yarn is especially preferable as it is compact. As the non-elastic yarn, a textured yarn is preferable as its stretchability, adhesiveness with an elastic film etc., are excellent. Among textured yarns, false-twisted textured yarns with excellent stretchability are preferable. When a non-textured yarn is used, there exists a defect that separation thereof from an elastic yarn easily occurs but said defect can be avoided by taking such countermeasures as increasing sufficiently the number of twists.

It is preferable that the balloon consists of a three-layered structure of an elastic film/a cylindrical



body of a composite yarn/an elastic film. Namely, the inner elastic film provides an action utilizing a reinforcing effect caused by the composite yarn and the outer elastic film provides an action making passing through the blood vessel better. The balloon based on the present invention can be applied to most catheters with a balloon, but a furthermore excellent effect can be expected when it is applied to a catheter with a balloon wherein one end of the balloon is fixed on the inner tube (the inner tube may be an inner-filled rod) and another end thereof is fixed on the outer tube of a double tube wherein both inner and outer tubes can be concentrically slidable each other. Namely, in said catheter with a balloon, the outer diameter of the balloon can be furthermore decreased by sliding both tubes and thereby stretching the balloon in the axial direction when it is inserted in a blood vessel and inside of the heart or taken out thereof. If the rigidity of the inner tube is too small not to stretch the balloon, the balloon can be effectively stretched in the axial direction by providing a stopper on the apex of said inner tube and pushing it with a rod insertable in said inner tube. If the rigidity of the outer tube is insufficient, it is preferable that a high-modules fiber is fixed on the apex of the outer tube and it is guided to the opposite end.



The elastic film is often made of a thin membrane with large stretchability and smooth surface in such a way that the thickness of the front end side is thinner than that of the rear end. As the raw material, polyurethanes and rubbers are used and especially, rubbers having small initial moduli are excellent materials as raw materials for the elastic film.

The thickness of the elastic film is preferably in the range of 0.1-0.4 mm and it is not necessarily the same thickness between the inner and the outer layers.

Difference in thickness between the front end side and the rear end side is preferably about 5-20% and the difference can be obtained by making it thinner, for example, when a cylindrical elastic film is prepared or by expanding a corresponding part of the prepared cylindrical elastic film and fixing the residual elongation thereof as a simple method.

As the cylindrical body made of the composite yarn, a knitted fabric, a woven fabric, a braid or a wound body in a cylindrical shape with a large twill angle can be used and among them, a cylindrically-knitted structure with stretchability is preferable.

When a guitar-shape balloon is prepared, for example, an elastic band are adhered on the middle part of an elastic body with an adhesive such as a rubber paste.



As the elastic yarn being a core yarn for a textured yarn, any fiber with stretchability can be used without specific limitation. For example, a single yarn or a twisted yarn made of a rubber such as natural rubbers and synthetic rubbers or a polyurethane is suitable and as the yarn to be wound made of a synthetic fiber, not only high-tenacity nylon, polyester and "Teflon" yarns but high-tenacity polyimide and polyethylene yarns can be used as a single yarn, but a twisted yarn being flexible is preferable. The thicknesses of these yarns are not specifically limited as the required pressure-resistant characteristics are different depending upon the position to be worked and the purpose, but in many cases, the thickness of the elastic yarn being a core yarn is selected in the range of 10-50D and that of the tenacity or high tenacity yarn to be wound is selected in the range of 30-150D.

Then, if the outer elastic film is adhered on a cylindrical body of a composite yarn and the inside of the cylindrical body of the composite yarn between an elastic film is coated with a silicone oil etc., a gap caused by expansion or shrinkage is not produced and a uniform expansion can be obtained. The length of the balloon can be freely set and in many cases, it is preferably in the range of 15-70 mm.



It is preferable that both ends and apexes of a balloon and the connected parts with tubes can be finished with an adhesive to make their surfaces smooth.

As the raw material used for a soft inner tube in coaxially (concentrically)-assembled tubes, for example, polyvinyl chloride and polyurethane are preferably used. The length of this tube is preferably 5-25 cm and it is preferable that the diameter is 14 Fr or smaller as it is inserted by means of Celludinga method.

In addition, as the relatively hard outer tube with torque characteristics, for example, a tube made of a synthetic resin such as polyethylene, "Teflon", polypropylene and nylon is used. Besides these tubes, for example, hard tubes such as a tube made by extracting a plasticizer for polyvinyl chloride with a solvent or a tube cured by crosslinking by means of irradiation. It is preferable that the diameter of this outer tube is 14 Fr or smaller because of the above described reason. In addition, the length is not specifically defined as it is different depending on age and size of the patient and the distance from the inserted part to the position to be cured, but a product with a total length in the range of 50-200 cm is preferably used.

In addition, depending on the position to be cured, a relatively hard tube similar to the outer tube can be



sometimes preferably substituted for the soft inner tube as the relatively hard tube provides an action making insertion of a rigid body easy.

Furthermore, if a contrast-medium such as barium sulfate, bismuth compd. or a metal powder such as tungsten is mixed in these tubes, it is preferable that the position in the blood vessel can be helpfully thereby detected by X-ray irradiation.

In addition, as the raw material of the yarn or the knitted fabric buried in or set along with the soft inner tube, a thin and strong one is preferable and for example, said yarns can be all used but polyamide yarns are the best among them.

In addition, it is necessary that an air removal tube set between an inner tube and an outer tube does not interference the flexibility of the soft tube and a tube with a thin thickness and a small diameter, for example, a tube made of polyimide or "Teflon" or a polyethylene tube is used.

As a rigid body inserted in an inner tube and stretching a balloon, for example, a stainless pipe or thin rod can be used.

In addition, as a fluid expanding a balloon, for example, physiological saline, carbon dioxide or a contrast-medium is used. These fluids are used by a



method wherein a specified amt. thereof is injected thereinto by means of an injection cylinder etc.

The catheter with a balloon of the present invention has such features that the stretchability is large and it exhibits pressure resistance and a uniform expanding property by using a cylindrical body made of a composite yarn with enlarged stretchability.

By the present ^{embodiment} ~~invention~~, even a balloon with a large diameter can be inserted in a blood vessel in such a way that it is made long and slender and small and in addition, there is no possibility of damaging the inner wall of the blood vessel when the apex part of a tube rubs it and the balloon can be transferred to the position to be cured through the blood vessel and the heart by utilizing a guide wire and the floating property of the balloon.

Therefore, the time required for curing is short and management after surgical operation is simple and recovery can be rapidly performed. Furthermore, this method can be provided as a method wherein expansion of the occluded part of a blood vessel or an opening technique for a valve of the heart by means of a balloon can be non- operationally, simply and easily performed. In addition, the balloons of the present invention can be furthermore simply manufactured in such a way that products with



various sizes are separately manufactured to provide catheter with balloons with good yield and with low cost.

(Examples)

The present invention will be explained in more detail by using examples as follows.

Example 1.

A false-twisted textured yarn consisting of a polyester yarn (a non-elastic yarn) of 70D-24f was inter-twisted to a polyurethane yarn (an elastic yarn: "Opelon") of 20D at a speed of 400 T/m while the former was overfed 300% to said urethane yarn to manufacture a composite yarn.

The ratio of free length of said yarn was 0.26 read from the S-S curve.

A balloon was prepared by using this composite yarn.

Namely, as an outer tube, a polyethylene tube containing a contrast-medium with 13.5 Fr and a length of 560 mm and a polyvinyl chloride tube containing 45 parts of DOP with 13.5 Fr and a length of 130 mm were connected through a stainless pipe with a diameter of 3.5 mm and a stainless pipe with a diameter of 3 mm and a length of 6 mm was attached on the apex of the polyvinyl chloride tube. In this time, a polyimide twisted yarn of 400D was placed on the hollow side of the polyvinyl chloride and fixed on stainless pipes on both sides. As an inner tube,



on the apex of a polyvinyl chloride tube containing 45 parts of DOP with 6.6 Fr, a stainless pipe with a diameter of 1.6 mm and a length of 7 mm having a projected part inside made by means of a corset was attached and a stainless pipe with 14G and a length of 70 mm and a needle base was inserted in the rear end. Connection of the inner and outer pipes and the inserted stainless pipe were bound and fixed with a nylon yarn with a thickness of 70 μ m. This inner tube was passed through a W-connector and furthermore inserted into the outer tube and then said outer tube was fixed on the W-connector to prepare a tube part.

Then, as an inner elastic film of the balloon, a rubber with a thickness of 0.25 mm and a length of 25 mm was bound and fixed on each apex of the inner tube and the outer tube with a nylon yarn with a thickness of 70 μ m and the surface thereof was coated with a small amount of a silicone oil. Separately, a cylindrically-knitted fabric was prepared by knitting said composite yarn by means of a knitting machine wherein 50 knitting needles were arranged on a circumference with a diameter of 20 mm. On the middle part of said knitted fabric with a length of 30 mm, a rubber with a thickness of 0.25 mm and a length of 7 mm was adhered with a rubber paste to prepare a band. Furthermore on the outside thereof, a part prepared by



adhering a rubber tube with a thickness on the front side of 0.25 mm and a thickness on the rear side of 0.3 mm was covered. They were all fitted together in such a way that the total length was 25 mm and both ends thereof were bound and fixed with a nylon yarn with a thickness of 70 μ m. Then, excess knitted fabric and rubbers were cut off to complete a balloon part.

Then, after a polyimide tube with a diameter of 0.9 mm was inserted into a fluid transferring path between the inner and outer tubes from one of remaining openings of the W-connector up to 10 mm to the balloon, the rear part up to the inside of the W-connector was covered with a reinforcing polyethylene tube. After a two directional stopcock was fixed on the rear end thereof and the gaps of the openings were filled with an adhesive material, finally, the surfaces of the both ends of the balloon, the apex and the connected parts of tubes were finished smoothly with an epoxy resin to complete a catheter with a balloon of the present invention.

This catheter with a balloon was attached on an injection cylinder and after water was substituted for air while air was evacuated from an air vent, about 7 ml of water was injected therein. Only apex part of this balloon was expanded to about 23 mm. Then, when the injected amount was made to 23 ml, the balloon turned into



a cylindrical shape with a diameter of about 28 mm.

Then, water was pulled out and the surface was smoothly shrunk without producing wrinkles. Then, a stainless steel tube with a diameter of 1.19 mm was inserted into the inner tube to push the apex part of the inner tube and to stretch thereby the balloon from 25 mm to 55 mm. The diameter of the balloon was changed thereby from about 7 mm to 4.5 mm. When the stainless tube was pulled backward, the balloon was returned to its original diameter and length.

Example 2.

A catheter with a balloon similar to the one of Example 1 was used for a clinical treatment for an inter-related tearing and opening technique for the mitral valve. Namely, using a method perforating the atrium partition, a specially-made guide wire was inserted into the left atrium through the skin, the crotch vein, the right atrium and the atrium partition. Then, a dilator of 14 Fr was forwarded along with this guide and the perforated parts on the crotch vein and the atrium partition were expanded. A catheter with a balloon of the present invention was successively inserted into the left atrium from the right atrium through the atrium partition and only the apex of the balloon was expanded to a diameter of 10 mm by using carbon dioxide and it was run



into a valve opening with the blood flow as a Swanganz catheter was. Then, it was inserted into the left ventricle and the apex side of the balloon was furthermore expanded by using a contrast-medium and it was lightly drawn nearby and brought into contact with the valve opening to be teared and opened. At this position, the balloon was furthermore expanded and the valve opening was pinched with the constricted balloon. The balloon was expanded as they were under this condition and the valve opening can be enlarged without slipping out of the balloon.

The tearing and opening technique needed 1 hour containing 5 sec for expansion and shrinkage of the balloon without any side effect. 3 days were needed from entering a hospital to leaving the hospital. The time needed was only less than a half in comparison with a tearing and opening technique accompanied with exposure and cutting-out of the blood vessel.

Example 3.

A catheter with a balloon was prepared in the same way as in Example 1, but the denier of "Opelon" yarn as the elastic yarn was 30D and the ratio of overfeeding a polyester yarn and the number of twist thereof were changed to 270% and 400 T/m respectively.

In this catheter with a balloon, the ratio of free



length of the constituting yarns of the composite yarn was apparently 0.28 and the balloon exhibited the maximum expanded diameter of 30 mm and the minimum shrunk diameter of 4-5 mm and excellent stretching characteristics without generation of wrinkle.

Possibility of Industrial Application

The catheter with a balloon of the present invention can be widely used for a method for medical treatment wherein a shrunk balloon is got into a constricted or occluded blood vessel and a fluid is pressed into the balloon to expand the balloon and the expanded balloon enlarges the constricted or the occluded blood vessel to make the constricted or occluded part normal. Especially, it can be widely used for a non-surgical and through-skin type valve tearing and opening operation substituting for a valve incisive operation for the valve of the heart which is constricted by aging, calcination or a sequela of rheumatic heat and a valve substitutive operation for an artificial valve.



THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A catheter with a balloon, said balloon being reinforced with a composite yarn adhered to or embedded in the balloon, said composite yarn comprising an elastic yarn and a non-elastic yarn with a larger free length than that of said elastic yarn.

2. A catheter with a balloon as described in Claim 1 wherein the ratio of free length of an elastic yarn to a non-elastic yarn of 0.15-0.5.

3. A catheter with a balloon as described in Claim 1 wherein the ratio of free length of an elastic yarn to a non-elastic yarn of 0.2-0.35.

4. A catheter with a balloon as described in Claim 1 wherein said composite yarn is a core-sheath yarn and wherein an elastic yarn is the core and a non-elastic yarn is the sheath.

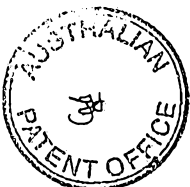
5. A catheter with a balloon as described in Claim 1 wherein said composite yarn is a core-sheath yarn and wherein a synthetic fiber-twisted yarn is spirally wound on a urethane yarn.

6. A catheter with a balloon as described in Claim 1 wherein the non-elastic yarn is a false-twisted textured yarn.

7. A catheter with a balloon, said balloon being reinforced with a knitted or woven fabric of a composite yarn adhered to or embedded in the balloon, said composite yarn comprising an elastic yarn and a non-elastic yarn with a larger free length than that of said elastic yarn.

8. A catheter with a balloon as described in Claim 7 wherein said knitted or woven fabric is a knitted fabric.

9. A catheter with a balloon as described in Claim 1 wherein the balloon is a three-layered structure consisting of an elastic film, a cylindrical fabric of the composite yarn, and an elastic film.



10. A catheter with a balloon as described in Claim 9 wherein the cylindrical fabric of the composite yarn is adhered with one or both of the elastic film(s).

11. A catheter with a balloon as described in Claim 1 wherein one end of the balloon is fixed on the inner tube of a double tube and wherein said tubes are concentrically slidable with respect to each other and another end thereof is fixed on the outer tube.

12. A catheter with a balloon as described in Claim 11 further comprising a shaft insertable in said inner tube and a stopper for said shaft on the apex of said inner tube.

13. A catheter with a balloon substantially as hereinbefore described with reference to the accompanying drawings and/or the examples.

DATED this 13th day of March 1992

Toray Industries, Inc.

By Its Patent Attorneys

DAVIES COLLISON CAVE



ABSTRACT

The present invention relates to a catheter with a balloon characterized by a feature that the balloon is reinforced with a composite yarn consisting of an elastic yarn and a non-elastic yarn with a larger free length than that of said elastic yarn.

As the catheter with a balloon has a balloon with large stretchability, when the catheter is inserted into a blood vessel or taken out on a surgical operation, it is possible not only to make surely the diameter small without producing any wrinkle but to stretch surely to the same size and the same shape at any time. Therefore, not only reliability of the surgical operation is improved but also a cut wound for the surgical operation could be smaller. There exist such advantages that treatment and management after the surgical operation are simple and the time required for recovery is extremely short and smaller daily burdens of a patient and an attendant are needed.



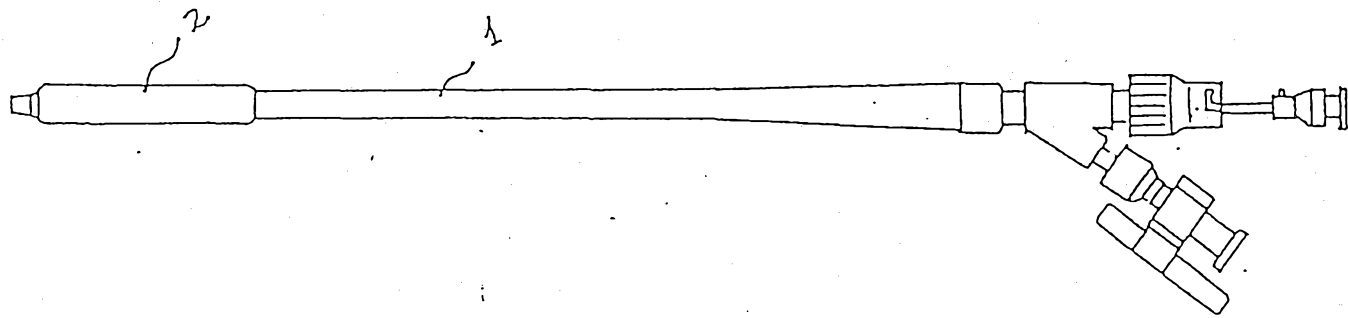


Figure 1

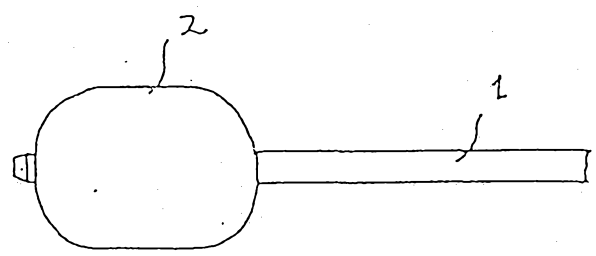


Figure 2

11

INTERNATIONAL SEARCH REPORT

International Application No PCT/JP90/00587

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC		
Int. Cl ⁵ A61M25/10		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
IPC	A61M25/00-25/10, A61L29/00	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
Jitsuyo Shinan Koho 1950 - 1990 Kokai Jitsuyo Shinan Koho 1971 - 1990		
III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	JP, A, 61-228877 (Kuraray Co., Ltd.), 13 October 1986 (13. 10. 86), Pages 1, 2 and 6; Fig. 1 (Family: none)	1 - 12
¹⁰ 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 "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 "&" document member of the same patent family		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
July 19, 1990 (19. 07. 90)	July 30, 1990 (30. 07. 90)	
International Searching Authority	Signature of Authorized Officer	
Japanese Patent Office		

国 際 調 査 報 告

国際出願番号PCT/JP90/00587

I. 発明の属する分野の分類		
国際特許分類 (IPC) Int. Cl. ⁸ A61M25/10		
II. 国際調査を行った分野		
調 査 を 行 っ た 最 小 限 資 料		
分 類 体 系	分 類 記 号	
I P C	A61M25/00-25/10, A61L29/00	
最小限資料以外の資料で調査を行ったもの		
日本国実用新案公報 1950-1990年 日本国公開実用新案公報 1971-1990年		
III. 関連する技術に関する文献		
引用文献の カテゴリー※	引用文献名 及び一部の箇所が関連するときは、その関連する箇所の表示	請求の範囲の番号
A	JP, A, 61-228877 (株式会社 クラレ), 13. 10月. 1986 (13. 10. 86), 第1-2頁, 第6頁第1図 (ファミリーなし)	1-12
※引用文献のカテゴリー 「A」特に関連のある文献ではなく、一般的技術水準を示すもの 「E」先行文献ではあるが、国際出願日以後に公表されたもの 「L」優先権主張に疑義を提起する文献又は他の文献の発行日若しくは他の特別な理由を確立するために引用する文献 (理由を付す) 「O」口頭による開示、使用、展示等に言及する文献 「P」国際出願日前で、かつ優先権の主張の基礎となる出願の日の後に公表された文献 「T」国際出願日又は優先日の後に公表された文献であって出願と矛盾するものではなく、発明の原理又は理論の理解のために引用するもの 「X」特に関連のある文献であって、当該文献のみで発明の新規性又は進歩性がないと考えられるもの 「Y」特に関連のある文献であって、当該文献と他の1以上の文献との、当業者にとって自明である組合せによって進歩性がないと考えられるもの 「&」同一パテントファミリーの文献		
IV. 認 証		
国際調査を完了した日 19.07.90	国際調査報告の発送日 30.07.90	
国際調査機関 日本国特許庁 (ISA/JP)	権限のある職員 特許庁審査官 安 田 達 夫®	406859