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**Miyagawa et al.**

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(54) **CATHETER**

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(52) **U.S. Cl.**

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(56)

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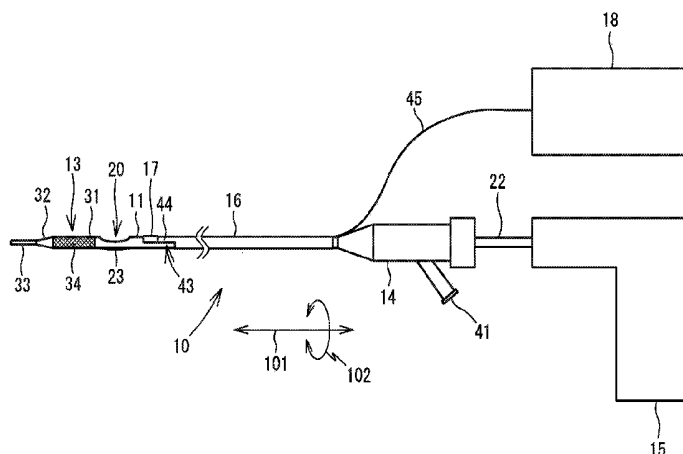
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**ABSTRACT**

[Object] To provide a catheter having a simple structure capable of excising an atheroma in a blood vessel and capable of obtaining ultrasonic images of a blood vessel.

[Solution] A catheter **10** has a shaft **11** having an opening **20** in a part of the side wall on the distal end side, a cutter **12** which is located in the vicinity of the opening **20** in the internal space of the shaft **11** and which can move in the axial direction **101** of the shaft **11**, a balloon **23** which is disposed on the side opposite to the opening **20** with respect to the axis of the shaft **11** and which outwardly expands from the side wall of the shaft **11**, and a phased array ultrasound probe **17** disposed along the circumferential direction **102** of the outer peripheral surface of the side wall in the vicinity of

(Continued)



the opening 20 at least on the same side as the side where the opening 20 is provided with respect to the axis of the shaft 11.

### 5 Claims, 11 Drawing Sheets

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*A61B 90/00* (2016.01)  
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- (52) **U.S. Cl.**  
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- (58) **Field of Classification Search**  
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FIG. 1

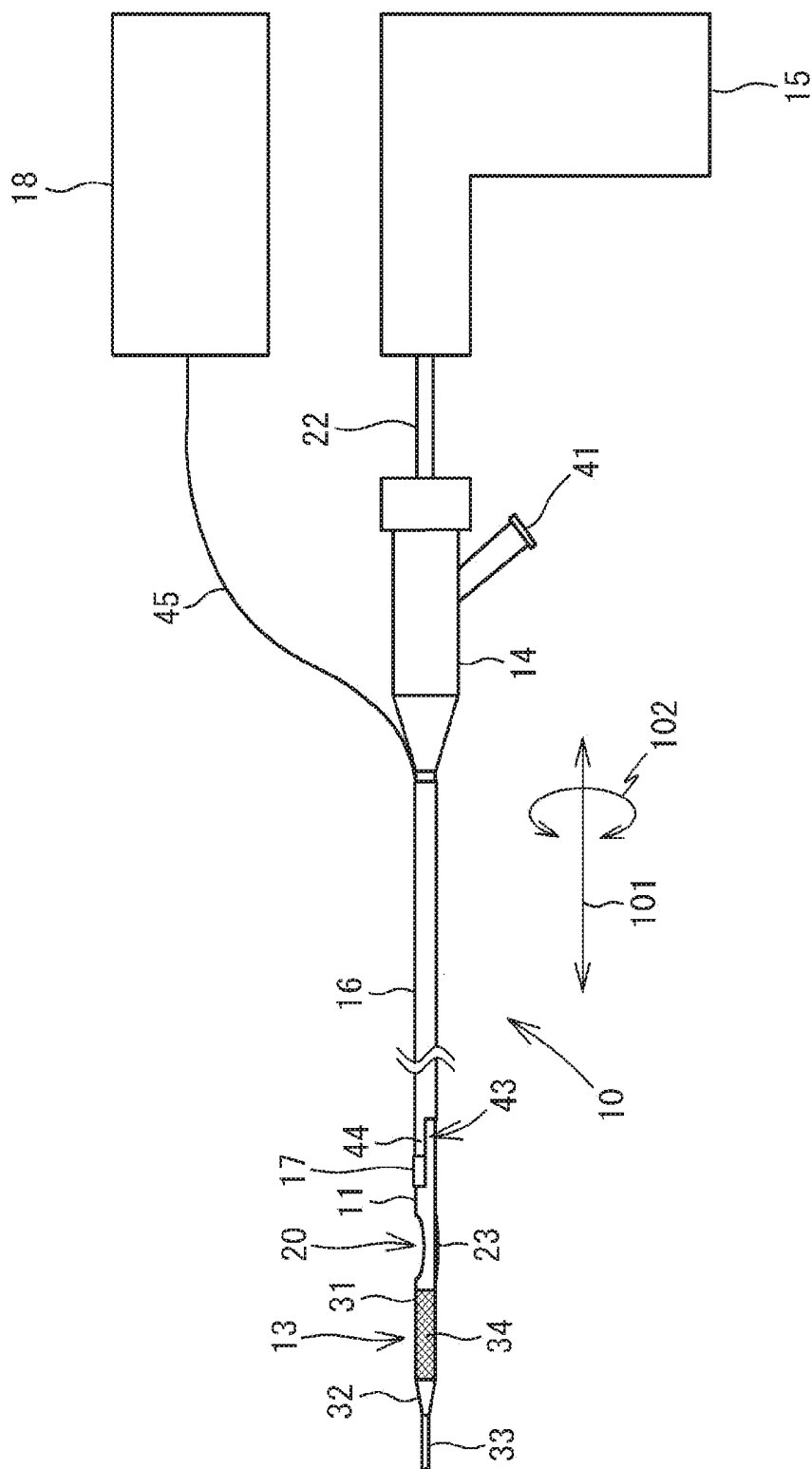


FIG. 2

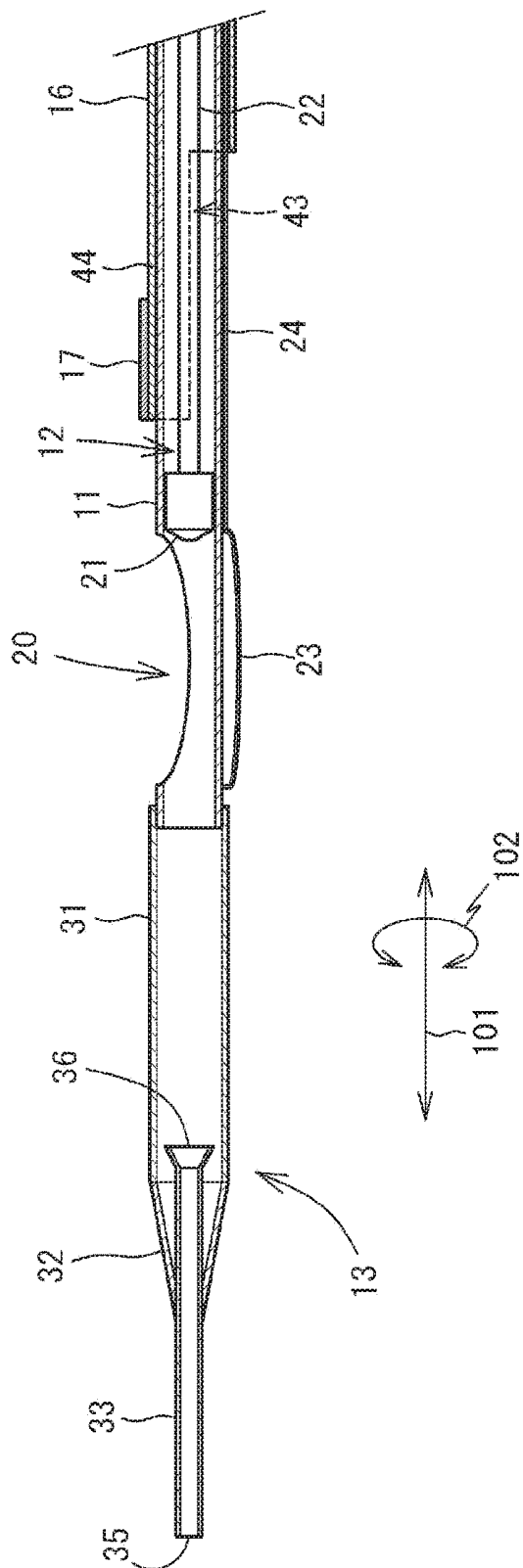


FIG. 3

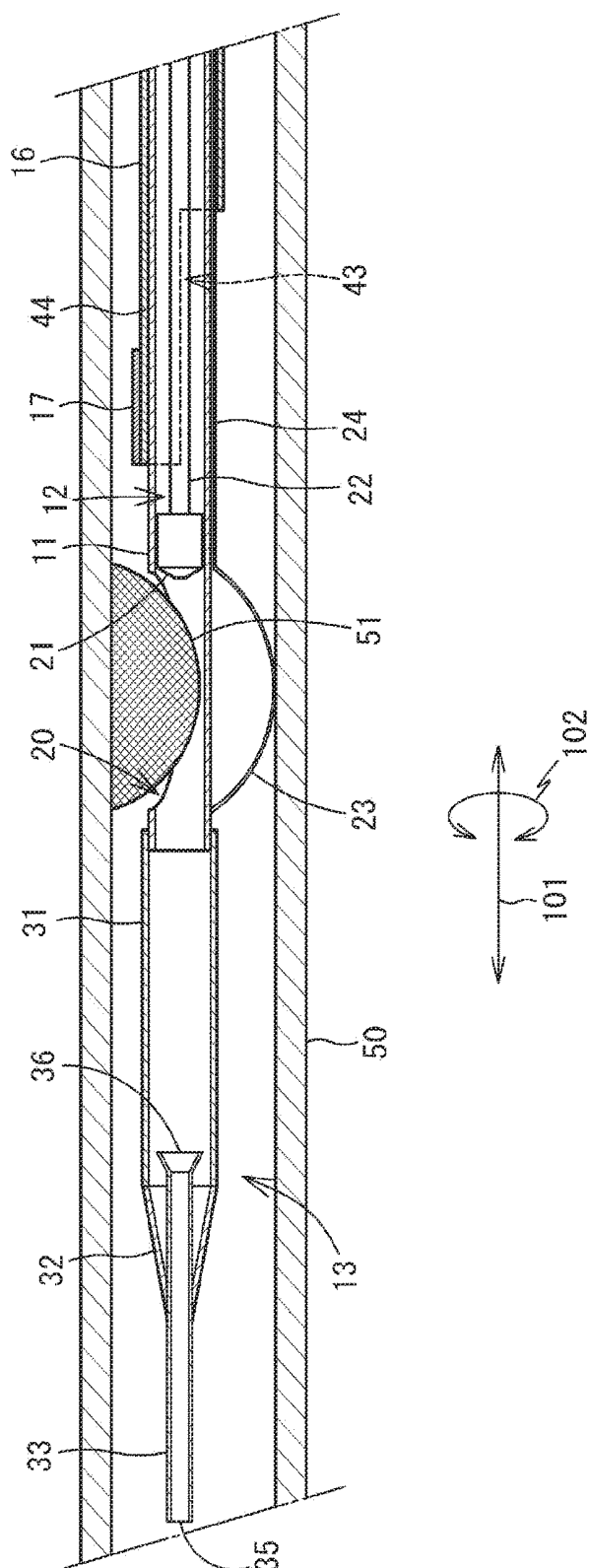


FIG. 4

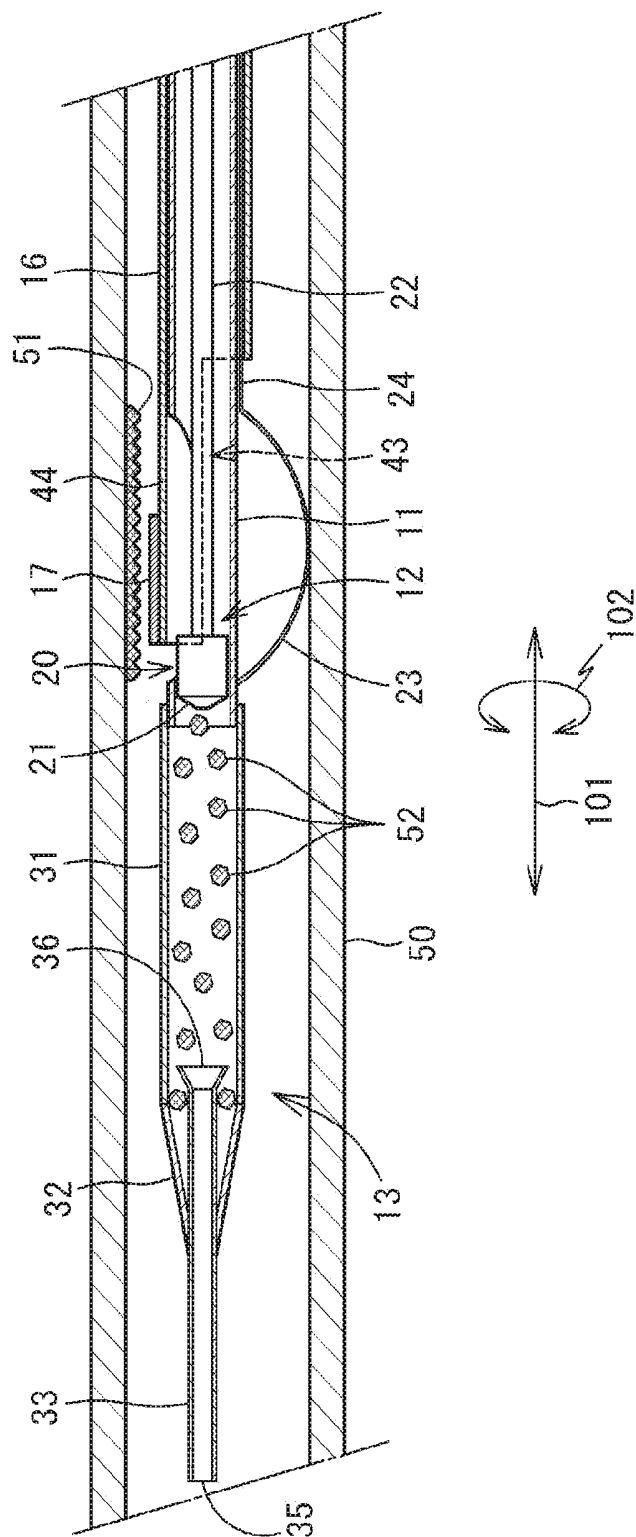


FIG. 5

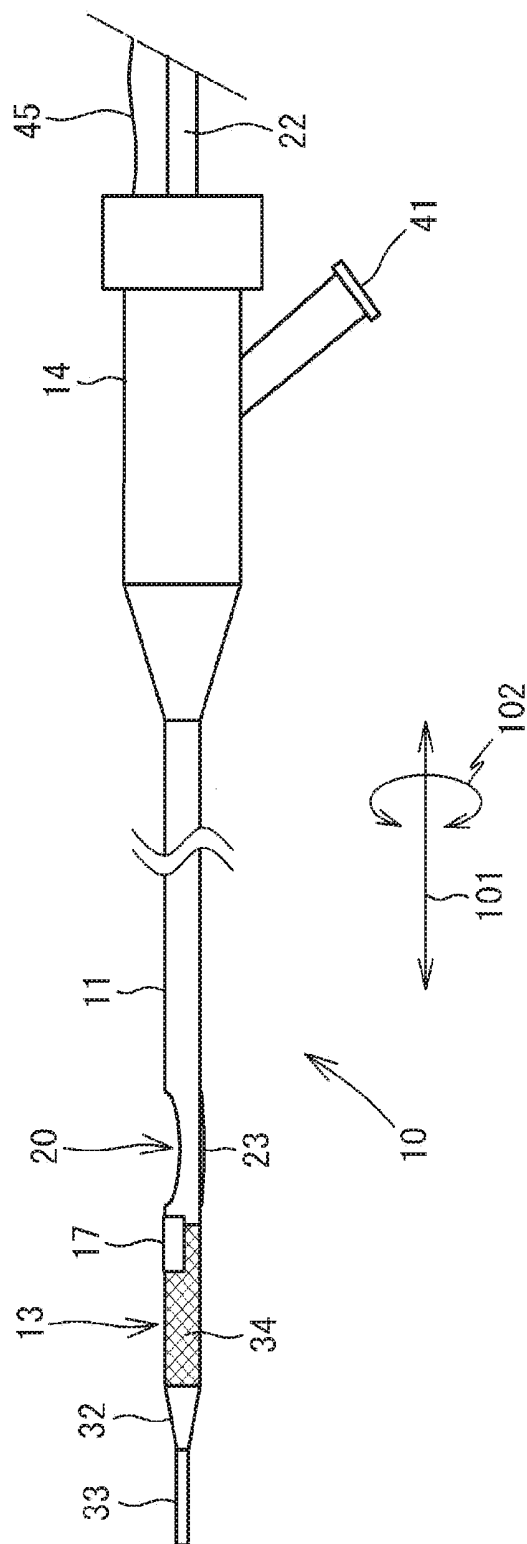


FIG. 6

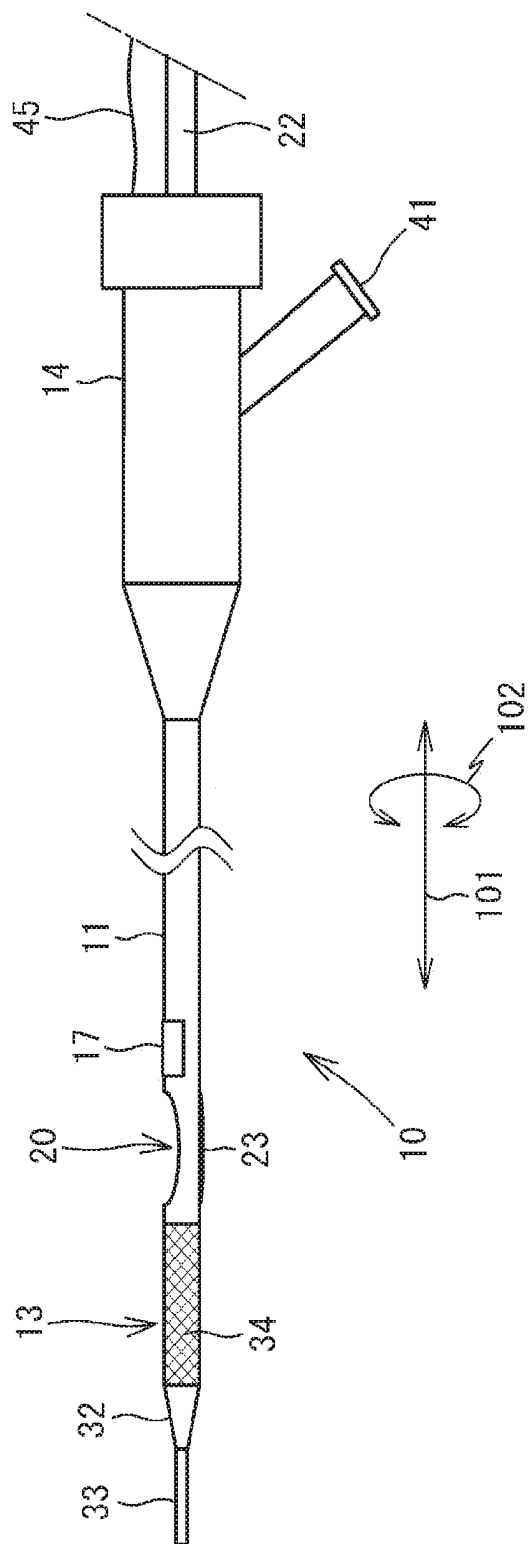




FIG. 7

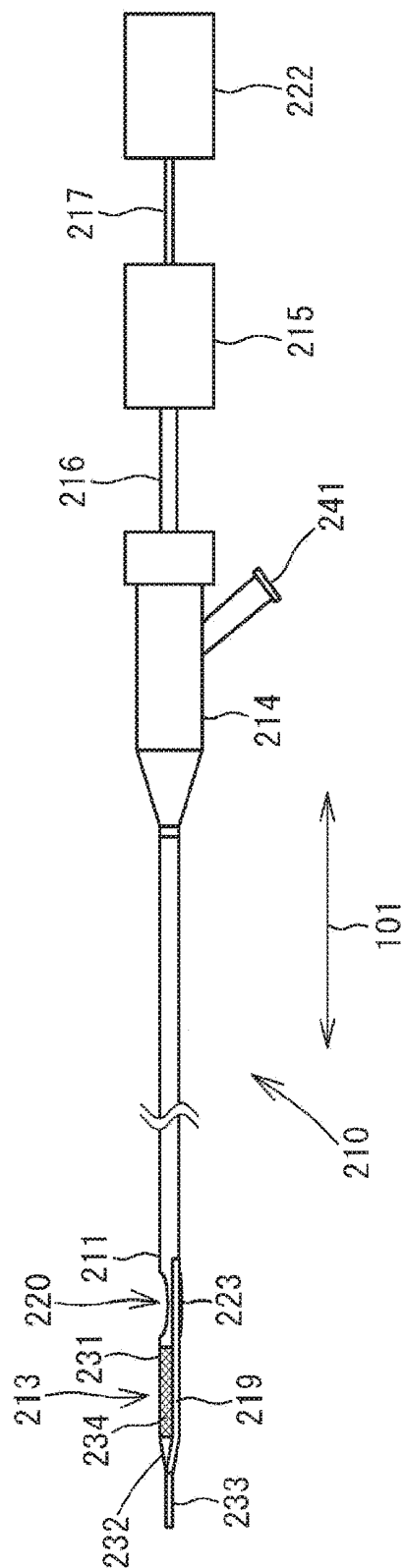


FIG. 8

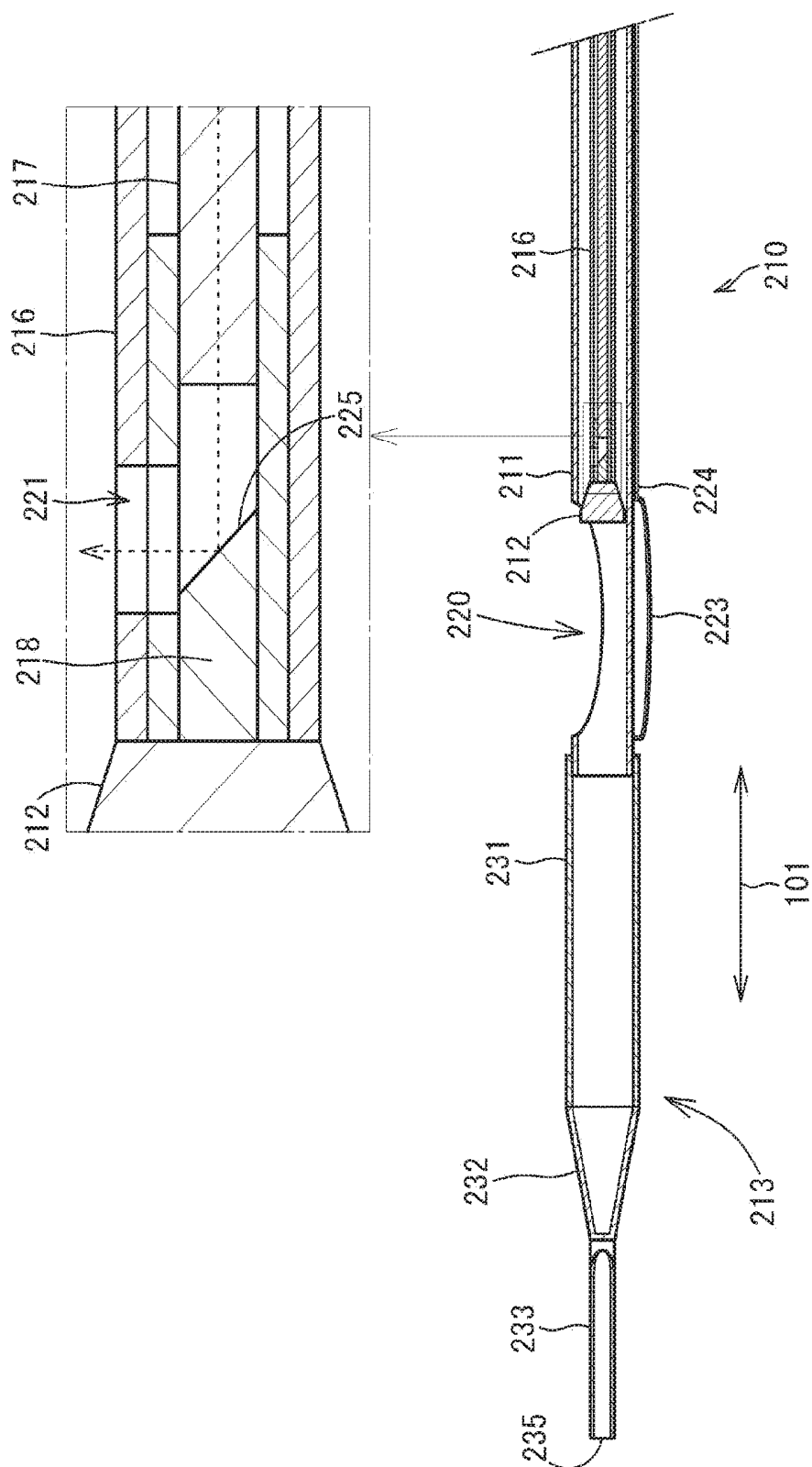


FIG. 9

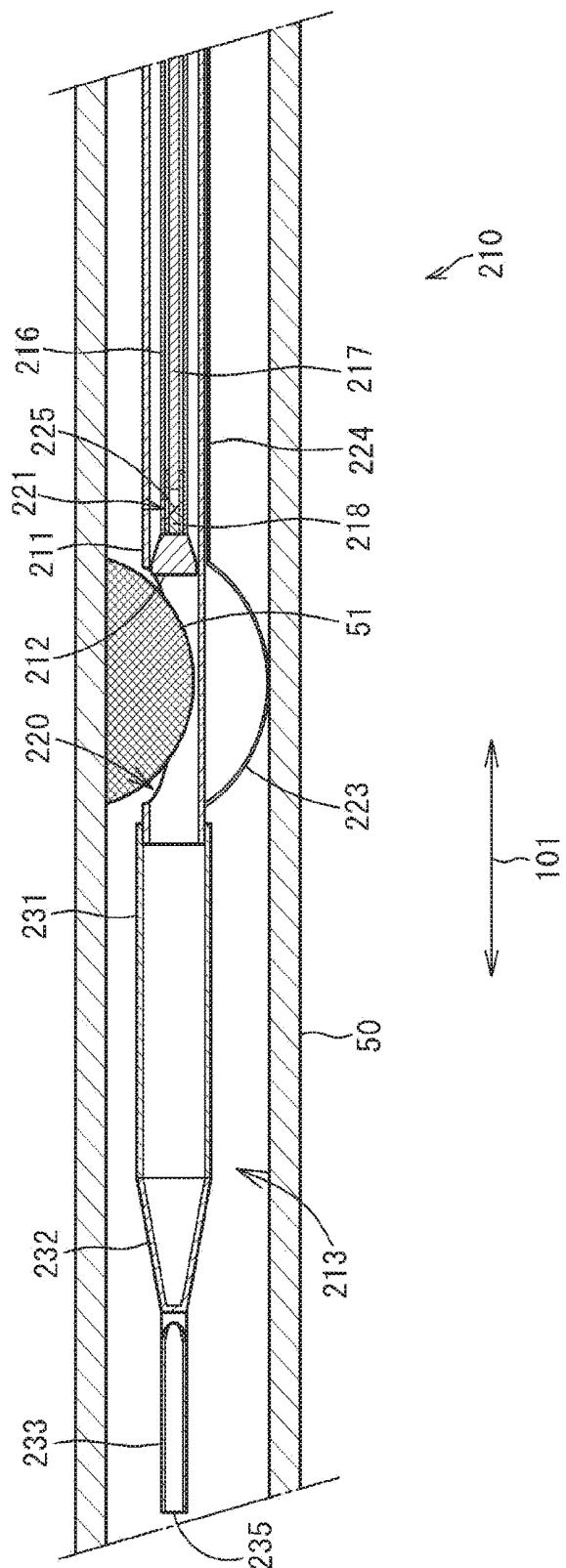
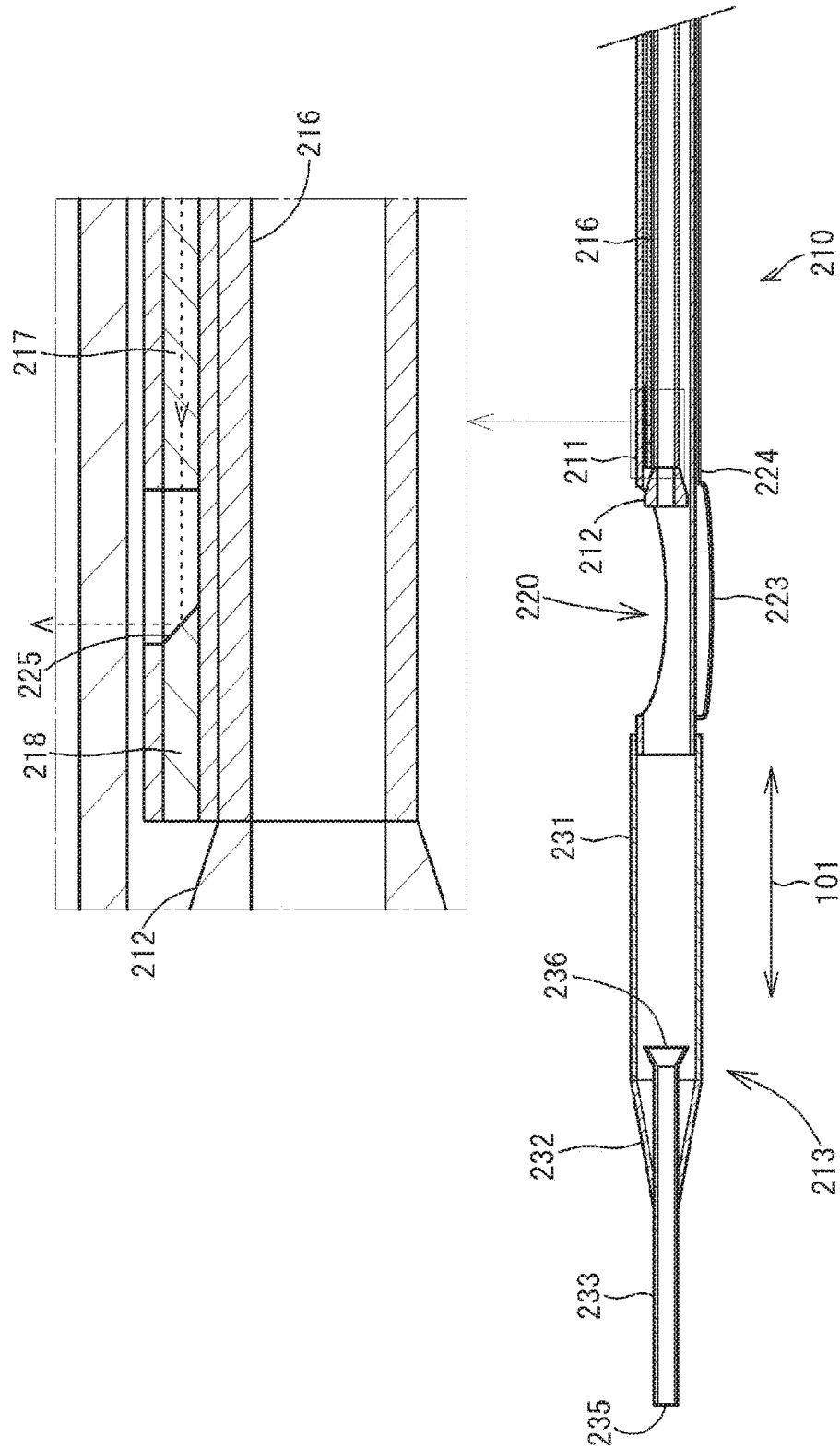




FIG. 11



# 1 CATHETER

## CROSS-REFERENCE TO RELATED APPLICATION

This application is a 35 U.S.C. 371 National Phase Entry Application from PCT/JP2015/053101, filed Feb. 4, 2015, which claims the benefit under 35 U.S.C. § 119(a) of the filing date of Japanese patent application No. 2014-020927, filed Feb. 6, 2014, and Japanese patent application No. 2014-104489, filed May 20, 2014, the respective disclosure(s) which is(are) incorporated herein by reference.

## TECHNICAL FIELD

The present invention relates to a catheter to be inserted into a blood vessel in order to excise an atheroma adhering to the inner wall of a blood vessel.

## BACKGROUND ART

Heretofore, medical treatment of inserting a catheter into a blood vessel to excise an atheroma generated in the blood vessel has been performed. An analysis of whether or not an atheroma is generated in a blood vessel has been performed by inserting a catheter having an ultrasound probe into the blood vessel to obtain an ultrasonic image. Moreover, for the analysis of whether or not an atheroma is present in a blood vessel, image analysis by an optical coherence tomography (hereinafter also referred to as "OCT"), for example, can be conducted (Patent Literature 4).

In order to excise an atheroma found in a blood vessel, an atheroma excision catheter has been used which excises an atheroma with a cutter through an opening in the side wall. The atheroma excision catheter is inserted into a blood vessel to the position where an opening faces the atheroma, and then a balloon provided in the vicinity of the opening is expanded, so that the opening is brought close to the atheroma, so that the atheroma enters a lumen of the atheroma excision catheter through the opening. In the lumen of the atheroma excision catheter, a cutter is movably provided in the axial direction. The cutter receives drive transmitted from a motor to be rotated in the lumen. The cutter is moved in the lumen while being rotated to thereby excise the atheroma entering the lumen. The excised atheroma is accommodated in the lumen of the atheroma excision catheter (Patent Literatures 1 and 2).

In the excision of the atheroma described above, ultrasonic images are obtained immediately before the excision in order to align the positions of the atheroma and the opening of the atheroma excision catheter. Thereafter, the atheroma is excised by the atheroma excision catheter. Furthermore, ultrasonic images are obtained again in order to check the state of the blood vessel after the excision of the atheroma. When it is temporarily judged that the atheroma is not sufficiently excised, the remaining atheroma is excised again by the atheroma excision catheter, and thereafter the state of the blood vessel is checked again. In such an operation, the catheter having the ultrasound probe and the atheroma excision catheter are alternately inserted into the blood vessel, which is likely to complicate the operation. In view of such circumstances, a catheter has been devised in which an atheroma excision catheter is provided with an ultrasound probe capable of rotating around the axis (Patent Literature 3).

# 2 CITATION LIST

## Patent Literature

- 5 [Patent Literature 1] Japanese Unexamined Patent Publication Application No. 6-30943
- [Patent Literature 2] Japanese Unexamined Patent Publication Application No. 5-56984
- [Patent Literature 3] Japanese Unexamined Patent Application Publication (Translation of PCT Application) No. 9-182754
- 10 [Patent Literature 4] Japanese Unexamined Patent Application Publication (Translation of PCT Application) No. 2008-523954

## SUMMARY OF INVENTION

### Technical Problems

- 20 The atheroma excision catheter provided with an ultrasound probe described above has problems that the attachment structure of the cutter and the ultrasound probe and a structure for rotating each of the cutter and the ultrasound probe are complicated, the catheter size increases, and instability of the operation due to failures and the like increases.

- In the excision of the atheroma described above, in order to align the positions of the atheroma and the opening of the atheroma excision catheter, image analysis by the OCT is conducted before the excision, and thereafter, the atheroma is excised by the atheroma excision catheter. Furthermore, image analysis by the OCT is conducted again in order to check the state of the blood vessel after the excision of the atheroma. When it is temporarily judged that the atheroma is not sufficiently excised, the remaining atheroma is excised again by the atheroma excision catheter, and thereafter the state of the blood vessel is checked again. In such an operation, the catheter for the OCT and the atheroma excision catheter are alternately inserted into the blood vessel, which is likely to complicate the operation.

- The present invention has been made in view of such circumstances. It is an object of the present invention to provide a catheter having a simple structure capable of excising an atheroma in a blood vessel and capable of obtaining ultrasonic images of the blood vessel.

- It is another object of the present invention to provide a catheter having a simple structure capable of excising an atheroma in a blood vessel and capable of obtaining images of the blood vessel by the OCT.

### Solution to Problems

- (1) A catheter according to the present invention has a tube body having an opening in a part of the side wall on the distal end side, a cutter which is located in the vicinity of the opening in the internal space of the tube body and which can move in the axial direction of the tube body, a balloon which is disposed on a side opposite to the opening with respect to the axis of the tube body and which outwardly expands from the side wall of the tube body, and a phased array ultrasound probe disposed along the circumferential direction of the outer peripheral surface of the side wall in the vicinity of the opening at least on the same side as the side where the opening is provided with respect to the axis of the tube body.
- 60 The catheter is inserted into a blood vessel from the distal end side. Due to the fact that the balloon is expanded in a state where the opening of the tube body is aligned with an

atheroma in the blood vessel, the tube body is fixed to the blood vessel, so that the atheroma enters the internal space of the tube body from the opening. By the cutter moved in the axial direction of the tube body, the atheroma entering the internal space of the tube body is excised. By the ultrasound probe disposed in the vicinity of the opening of the tube body, ultrasonic images of the excised atheroma are obtained.

(2) The catheter further has an outer tube body which is provided on the outside of the tube body and which can relatively move in the axial direction with respect to the tube body, in which the ultrasound probe may be provided on the outer tube body.

Since the ultrasound probe can move in the blood vessel in connection with the relative movement of the outer tube body, ultrasonic images of cross sections or partial cross sections of the blood vessel different in the position in the length direction can be obtained by moving the ultrasound probe with respect to the atheroma.

(3) The outer tube body has a notch portion in which a part corresponding to the balloon is cut out, and a support portion which is disposed on a side opposite to the notch portion with respect to the axis of the outer tube body and which can be moved to a first position where the support portion covers the opening and a second position where the support portion opens the opening by the relative movement, in which the ultrasound probe is provided on the support portion and the notch portion may not abut on the balloon in the state of being outwardly expanded at the first position.

Even when the outer tube body is relatively moved to the first position where the outer tube body is overlapped with the opening in the state where the balloon is expanded, the outer tube body and the balloon do not interfere with each other by the notch portion. In the support portion at the first position, the atheroma located at a position corresponding to the opening and the ultrasound probe face each other, so that ultrasonic images of the atheroma can be obtained.

(4) The ultrasound probe may be fixed on the distal end side relative to the opening in the tube body.

(5) The ultrasound probe may be fixed on the proximal end side relative to the opening in the tube body.

(6) A catheter according to the present invention has a tube body having an opening in a part of the side wall on the distal end side, a cutter which is located in the vicinity of the opening in the internal space of the tube body and which can move in the axial direction of the tube body, a torque shaft which is inserted into and passed through the internal space of the tube body and is connected to the cutter in such a manner as to be able to transmit a rotation torque to the cutter, a light guide material provided along the torque shaft, a reflective material which reflects light emitted from the light guide material in a second direction crossing a first direction in which the light guide material is extended, and a balloon which is disposed on a side opposite to the opening with respect to the axis of the tube body and outwardly expands from the side wall of the tube body.

The catheter is inserted into a blood vessel from the distal end side. Due to the fact that the balloon is expanded in a state where the opening of the tube body is aligned with an atheroma in the blood vessel, the tube body is fixed to the blood vessel, so that the atheroma enters the internal space of the tube body from the opening. By the cutter moved in the axial direction of the tube body, the atheroma entering the internal space of the tube body is excised. Light emitted from the light guide material provided along the torque shaft is reflected by the reflective material to be emitted to the blood vessel. Due to the fact that the reflected light from the

blood vessel is treated as an interference signal by an OCT system through the reflective material and the light guide material, images of the blood vessel in the vicinity of the atheroma are obtained.

(7) Preferably, the light guide material and the reflective material are disposed in the internal space of the torque shaft and the reflective material emits light reflected in the second direction to the outside of the torque shaft through the opening formed in the side wall of the torque shaft.

Thus, light can be guided to be reflected in the torque shaft.

(8) Preferably, the light guide material and the reflective material can be rotated integrally with the torque shaft.

Thus, the excision of an atheroma and the OCT can be realized by controlling the number of rotations of a rotating device, such as a motor, connected to the torque shaft.

(9) Preferably, the light guide material and the reflective material can move integrally with the torque shaft along the first direction.

Thus, images of a blood vessel can be obtained along the first direction.

(10) Preferably, the light guide material and the reflective material are disposed in the internal space of the torque shaft.

Thus, the light guide material and the reflective material are covered with the torque shaft to be protected, and therefore the light guide material and the reflective material are hard to be damaged.

(11) Preferably, the light guide material and the reflective material are disposed on the outer peripheral surface side of the torque shaft.

Thus, even when a guide wire is inserted into and passed through the internal space of the torque shaft, light reflected from the reflective material is not blocked by the guide wire.

(12) Preferably, a guide wire lumen is provided along the tube body.

Thus, the catheter can be inserted into a blood vessel along the guide wire.

#### Advantageous Effects of Invention

According to the catheter of the present invention, an atheroma in a blood vessel can be excised and ultrasonic images of the blood vessel can be obtained by a simple structure.

Moreover, according to the catheter of the present invention, an atheroma in a blood vessel can be excised and images of the blood vessel by the OCT can be obtained by a simple structure.

#### BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a view illustrating the external configuration of a catheter 10 in a state where a balloon 23 is contracted.

FIG. 2 is an enlarged cross sectional view illustrating the internal configuration in the vicinity of a distal end portion 13 of the catheter 10.

FIG. 3 is a schematic view illustrating a state where the balloon 23 is expanded in a blood vessel 50.

FIG. 4 is a schematic view illustrating a state where a support portion 44 is located at a first position after an atheroma 51 is excised in the blood vessel 50.

FIG. 5 is a view illustrating a modification of the catheter 10.

FIG. 6 is a view illustrating a modification of the catheter 10.

FIG. 7 is a view illustrating the external configuration of a catheter **210** in a state where a balloon **223** is contracted.

FIG. 8 is an enlarged cross sectional view illustrating the internal configuration in the vicinity of a distal end portion **213** of the catheter **210**.

FIG. 9 is a schematic view illustrating a state where the balloon **223** is expanded in the blood vessel **50**.

FIG. 10 is a schematic view illustrating the state where the atheroma **51** is excised in the blood vessel **50**.

FIG. 11 is an enlarged cross sectional view illustrating a modification of the catheter **210**.

## DESCRIPTION OF EMBODIMENTS

Hereinafter, preferable embodiments of the present invention are described. It is a matter of course that the embodiments merely describe one embodiment of the present invention and the embodiments can be altered insofar as the scope of the present invention is not altered.

### First Embodiment

As illustrated in FIGS. 1 and 2, a catheter **10** has a shaft **11**, a cutter **12** provided in the shaft **11**, a distal end portion **13** configuring the distal end of the shaft **11**, a proximal end portion **14** connected to the proximal end of the shaft **11**, an actuator **15** which gives rotary drive to the cutter **12**, an IVUS shaft **16**, and an ultrasound probe **17** provided on the IVUS shaft **16**. The catheter **10** is used as a medical instrument which is inserted into a blood vessel to excise an atheroma or photographs ultrasonic images of a blood vessel.

The shaft **11** is a tube capable of containing the cutter **12** therein. The shaft **11** is configured from a surgical stainless steel circular tube or a synthetic resin circular tube, for example, and has flexibility which allows the shaft **11** to be elastically curved corresponding to the curved shape of a blood vessel. The distal end and the proximal end of the shaft **11** each are opened. The outer diameter of the shaft **11** is set according to the inner diameter of a blood vessel into which the shaft **11** is to be inserted, e.g., coronary arteries. The inner diameter of the shaft **11** is set according to the outer diameter of the cutter **12**. The outer diameter and the inner diameter of the shaft **11** are almost uniform over the axial direction **101** of the shaft **11**. The length in the axial direction **101** of the shaft **11** is set in consideration of the length from catheter insertion portions, such as the human limbs, to the affected portion.

As illustrated in FIG. 2, an opening **20** is formed in the vicinity of the distal end portion **13** in the shaft **11**. The opening **20** is formed by cutting out a part of the side wall of the shaft **11**. The shape and the size of the opening **20** are set in consideration of the shape and the size of an atheroma which is probably formed in the affected portion. The shaft **11** is equivalent to the tube body.

As illustrated in FIG. 2, the cutter **12** is provided in the vicinity of the opening **20** in the internal space of the shaft **11**. The cutter **12** has a cutting portion **21** and a shaft **22**. The cutting portion **21** has an approximately cylindrical shape and the outer diameter is a little smaller than the inner diameter of the shaft **11**. On the distal end side of the cutting portion **21**, a plurality of blades are formed in such a manner as to radially extend from the center. Although not illustrated in FIG. 2, a through-hole is formed along the axial direction **101** in the center of the cutting portion **21**. The shaft **22** is extended from the proximal end of the cutting portion **21** to the outside of the proximal end portion **14**. The shaft **22** is

a long and narrow tube and the internal space communicates with the through-hole of the cutting portion **21**. The internal space of the shaft **22** and the through-hole of the cutting portion **21** are configured so that a guide wire is inserted into and passed through the internal space and the through-hole.

As illustrated in FIG. 1, the shaft **22** is connected to the actuator **15**. Due to the fact that the shaft **22** receives drive transmitted from the actuator **15** to rotate, the cutting portion **21** rotates. Due to the fact that the shaft **22** is moved in the axial direction **101**, the cutting portion **21** moves in the axial direction **101** in the internal space of the shaft **11**.

As illustrated in FIGS. 1 and 2, a balloon **23** is provided at a position opposite to the opening **20** with respect to the axis of the shaft **11**. The balloon **23** can outwardly expand from the side wall of the shaft **11** and is folded and stuck to the side wall of the shaft **11** until the catheter **10** is inserted into a blood vessel. As raw materials of the balloon **23**, materials having biocompatibility are preferable. Specific examples of the materials include polyurethane, polyethylene, polyester, polypropylene, polyamide, polyamide elastomer, polytetrafluoroethylene, polyvinylidene fluoride, and the like.

As illustrated in FIG. 2, the proximal end side of the balloon **23** is connected to a tube body for balloon **24** provided along the side wall of the shaft **11**. The internal space of the tube body for balloon **24** is caused to communicate with the internal space of the balloon **23**. The tube body for balloon **24** is extended to the proximal end portion **14** and the internal space of the tube body for balloon **24** is connected to a port **41** of the proximal end portion **14**. Due to the fact that liquid, such as physiological saline, injected from the port **41** of the proximal end portion **14** flows into the balloon **23**, the balloon **23** is expanded in a blood vessel. The tube body for balloon **24** is a molded body of flexible plastic which can be elastically deformed, such as polyamide, polyamide elastomer, and polyetheramide.

As illustrated in FIGS. 1 and 2, the distal end portion **13** is connected to the distal end of the shaft **11**. As illustrated in FIG. 2, the distal end portion **13** has a blade tube **31**, a reduced diameter portion **32**, and a distal end tip **33**.

As illustrated in FIGS. 1 and 2, the blade tube **31** is a circular tube in which both sides are opened. The blade tube **31** is connected to the distal end of the shaft **11** and the internal space thereof is caused to communicate with the internal space of the shaft **11**. The blade tube **31** is one in which flexible plastic which can be elastically deformed, such as polyamide, polyamide elastomer, and polyetheramide, is reinforced with a core material **34**. The core material **34** is embedded in the side wall of the blade tube **31**. The core material **34** is formed into a cylindrical shape by braiding wire rods, such as surgical stainless steel, into a mesh. The inner diameter of the blade tube **31** is almost equal to the outer diameter of the shaft **11** and the blade tube **31** is fitted to the distal end of the shaft **11** from the outside. The outer diameter and the inner diameter of the blade tube **31** are almost uniform over the axial direction **101**. In each figure other than FIG. 1, the core material **34** is not illustrated.

As illustrated in FIGS. 1 and 2, the reduced diameter portion **32** is a circular tube in which both sides are opened and the outer diameter decreases in a tapered shape. The reduced diameter portion **32** is connected to the distal end of the blade tube **31** and the internal space thereof is caused to communicate with the internal space of the blade tube **31**. The reduced diameter portion **32** contains flexible plastic which can be elastically deformed, such as polyamide and polyetheramide. The inner diameter on the proximal end



side of the reduced diameter portion 32 is almost equal to the outer diameter of the distal end of the blade tube 31. The reduced diameter portion 32 is fitted to the distal end of the blade tube 31 from the outside, and then thermally fused thereto. The inner diameter on the distal end side of the reduced diameter portion 32 is almost equal to the outer diameter of the central portion of the distal end tip 33. On the distal end side of the reduced diameter portion 32, the thickness decreases toward the distal end side.

As illustrated in FIGS. 1 and 2, the distal end tip 33 is a circular tube in which both sides are opened and the outer diameter on a proximal end 36 side increases in a tapered shape. The distal end tip 33 is connected to the distal end of the reduced diameter portion 32 and the internal space thereof is caused to communicate with the internal space of the reduced diameter portion 32. The distal end 35 of the distal end tip 33 is projected from the distal end of the reduced diameter portion 32 to the outside in the axial direction 101. The proximal end 36 side of the distal end tip 33 is extended in the axial direction 101 in the internal space of the reduced diameter portion 32 and the proximal end 36 reaches the internal space of the blade tube 31. More specifically, a portion including a distal end side portion of the blade tube 31 and the reduced diameter portion 32 has a double tube structure in which the blade tube 31 and the reduced diameter portion 32 serve as the outer side and the distal end tip 33 serves as the inner side.

Although the diameter of the distal end tip 33 is increased on the proximal end 36 side, the outer diameter and the inner diameter of the other portion are almost uniform. The outer diameter of the uniform portion is smaller than the inner diameter of the blade tube 31 and is almost equal to the inner diameter of the distal end of the reduced diameter portion 32. Although the diameter of the proximal end 36 side is increased, the maximum diameter is smaller than the inner diameter of the blade tube 31.

The distal end tip 33 contains flexible plastic which can be elastically deformed, such as polyamide and polyetheramide. The distal end tip 33 is inserted into the distal end of the reduced diameter portion 32, and then thermally fused thereto. On the distal end of the distal end tip 33, a marker which can be confirmed by X-rays or the like may be provided.

The proximal end portion 14 is provided on the proximal end of the shaft 11. The proximal end portion 14 is a cylindrical member having an internal space continuing to the internal space of the shaft 11. The proximal end portion 14 is a molded body of resin, such as polypropylene or ABS. The proximal end portion 14 may serve as a handle in an operation of inserting and removing the shaft 11 into/from a blood vessel.

The proximal end portion 14 is provided with the port 41 extended in a direction crossing the axial direction 101. Another device, such as a syringe, is connected to the port 41, and then fluid, such as physiological saline, which is flown in and out from the device, flows in and out of the tube body for balloon 24 from the proximal end portion 14. The proximal end portion 14 may be provided with another port continuing to the internal space of the shaft 11. Such a port is used for the purpose of, for example, collecting an excised atheroma entering the inside of the shaft 11.

From an opening on the proximal end side of the proximal end portion 14, the shaft 22 of the cutter 12 is extended. The actuator 15 is connected to the shaft 22. In the actuator 15, a motor, a battery, and the like are built. To the shaft 22, rotation of the motor of the actuator 15 is transmitted.

As illustrated in FIGS. 1 and 2, the IVUS shaft 16 is provided on the outside of the shaft 11. The IVUS shaft 16 is a tube which allows the insertion of the shaft 11 into the IVUS shaft 16 in such a manner that the shaft 11 can relatively move in the axial direction 101. The IVUS shaft 16 is configured from a surgical stainless steel circular tube or a synthetic resin circular tube, for example, and has flexibility which allows the IVUS shaft 16 to be elastically curved corresponding to the curved shape of a blood vessel. The distal end and the proximal end of the IVUS shaft 16 each are opened. The outer diameter of the IVUS shaft 16 is set according to the inner diameter of a blood vessel into which the IVUS shaft 16 is to be inserted, e.g., coronary arteries. The inner diameter of the IVUS shaft 16 is set according to the outer diameter of the shaft 11. The outer diameter and the inner diameter of the IVUS shaft 16 are almost uniform over the axial direction 101. The length in the axial direction 101 of the IVUS shaft 16 is a little shorter than the length in the axial direction 101 of the shaft 11. Therefore, in the state where the IVUS shaft 16 is moved to the proximal end side with respect to the shaft 11 until the IVUS shaft 16 abuts on the proximal end portion 14, a distal end side relative to the opening 20 of the shaft 11 is projected from the distal end of the IVUS shaft 16 to be exposed to the outside as illustrated in FIG. 1. The IVUS shaft 16 is equivalent to the outer tube body.

As illustrated in FIG. 2, a notch portion 43 extending to the proximal end side along the axial direction 101 is formed at a position on a side corresponding to the side of the balloon 23 of the shaft 11 in the distal end portion of the IVUS shaft 16. The length in the circumferential direction 102 of the notch portion 43 is longer than the length in the circumferential direction 102 of the balloon 23 and about the half in the circumferential direction 102 of the IVUS shaft 16 is cut out. The length in the axial direction 101 of the notch portion 43 is a length in which the notch portion 43 does not abut on the balloon 23 in the state (FIG. 4) where the distal end portion of the IVUS shaft 16 is relatively moved to the distal end side with respect to the shaft 11 to a position where the distal end portion of the IVUS shaft 16 covers the opening 20 and is almost equal to the length in the axial direction 101 of the opening 20.

A portion which is formed into an approximately half tube shape by the notch portion 43 of the distal end portion of the IVUS shaft 16 is a support portion 44 for the ultrasound probe 17. The support portion 44 is located on the same side as the side on which the opening 20 is provided of the shaft 11 in the circumferential direction 102. When the IVUS shaft 16 is moved to the distal end side of the shaft 11, the support portion 44 is located at a first position (FIG. 4) where the support portion 44 covers the opening 20. When the IVUS shaft 16 is moved to the proximal end side of the shaft 11, the support portion 44 is located at a second position (FIG. 3) where the support portion 44 opens the opening 20.

The ultrasound probe 17 is provided on the outside of the support portion 44. The ultrasound probe 17 is a so-called phased array type, in which a plurality of elements are arranged along the circumferential direction 102 on the outer peripheral surface of the support portion 44. The number of the elements is not particularly limited. However, when the elements are arranged corresponding to the number of the elements provided on the half of the circumference of the IVUS shaft 16, for example, 32 elements are arranged in the circumferential direction. When the elements are electrically ignited in order, tomographic images of a blood vessel around the ultrasound probe 17 are collected. A cable 45 is

connected to the elements of the ultrasound probe 17. The cable 45 is inserted into and passed through the internal space of the IVUS shaft 16, extended to the outside from the proximal end, and then connected to the control device 18. The control device 18 is one which supplies electric power to the ultrasound probe 17 and forms tomographic images based on electric signals obtained from the ultrasound probe 17 and is a known control device for use in intravascular ultrasound (IVUS).

Although not illustrated in each figure, the IVUS shaft 16 may be provided with a fitting structure which regulates the movement in the circumferential direction in such a manner that the IVUS shaft 16 does not relatively rotate in the circumferential direction (around the axial direction 101) with respect to the shaft 11 and can relatively move only with respect to the axial direction 101. Due to the fact that the fitting structure or the like is provided, the position of the notch portion 43 of the IVUS shaft 16 and the position of the ultrasound probe 17 are not displaced with respect to the circumferential direction 102 of the shaft 11.

[Usage Directions for Catheter 10]

Hereinafter, the usage directions for the catheter 10 are described with reference to FIGS. 3 and 4.

The catheter 10 is used when excising an atheroma 51 formed in the inner wall of a blood vessel 50. The position of the atheroma 51 is confirmed by the IVUS or the like beforehand. The catheter 10 is inserted into the blood vessel 50 from the distal end portion 13 in the state (FIG. 1) where the balloon 23 is contracted. Although not illustrated in each figure, a guide wire is inserted into the blood vessel 50 beforehand in the insertion of the catheter 10 into the blood vessel 50. The insertion of the guide wire into the blood vessel 50 is performed by a known technique. The catheter 10 is inserted into the blood vessel 50 from the distal end portion 13 while inserting the guide wire, which is inserted into the blood vessel 50, into the internal space of the distal end tip 33 of the distal end portion 13, the internal space of the shaft 11, the through-hole of the cutting portion 21 of the cutter 12, and then the internal space of the shaft 22 in order.

The distal end portion 13 is advanced to the atheroma 51 in the blood vessel 50 while being elastically curved along the guide wire at a portion where the blood vessel 50 is curved, such as coronary arteries. When the distal end portion 13 reaches the atheroma 51, so that the opening 20 of the shaft 11 faces the atheroma 51, the insertion of the shaft 11 into the blood vessel 50 is ended. By moving the support portion 44 of the IVUS shaft 16 to the first position where the support portion 44 covers the opening 20, and then electrically igniting the ultrasound probe 17, tomographic images of the blood vessel 50 including the opening 20 are collected. By confirming the formed tomographic images of the blood vessel 50 by the control device 18, it can be confirmed that the opening 20 has reached a position corresponding to the position of the atheroma 51 and the state of the atheroma 51 can be confirmed. Thereafter, the guide wire is drawn out of the proximal end portion 14 side of the catheter 10. The actuator 15 is connected to the shaft 22 of the cutter 12.

As illustrated in FIG. 3, in the state where the opening 20 of the shaft 11 faces the atheroma 51 and where the support portion 44 is located at the second position where the support portion 44 of the IVUS shaft 16 opens the opening 20, the balloon 23 in the contracted state is expanded by fluid which is caused to flow into the tube body for balloon 24 from the port 41. When the expanded balloon 23 abuts on the inner wall of the blood vessel 50 on a side opposite to the atheroma 51, the opening 20 is stuck to the atheroma 51, so

that the atheroma 51 partially enters the internal space of the shaft 11 from the opening 20. In the state above, the catheter 10 is fixed to the blood vessel 50.

Subsequently, a motor of the actuator 15 is driven, so that the cutting portion 21 is rotated through the shaft 22 of the cutter 12. Due to the fact that the shaft 22 is advanced to the distal end side in the axial direction 10 with respect to the shaft 11 on the proximal end portion 14 side, the rotating cutting portion 21 abuts on the atheroma 51, so that the atheroma 51 is excised by the cutting portion 21. Fragments 52 of the excised atheroma 51 enter the internal space of the blade tube 31 through the internal space of the shaft 11.

After the atheroma 51 is excised by the cutter 12, the support portion 44 of the IVUS shaft 16 is located at the first position where the support portion 44 covers the opening 20. Then, the ultrasound probe 17 is electrically ignited, whereby tomographic images of the blood vessel 50 in the state where the atheroma 51 is excised are collected. Thus, the state where the atheroma 51 is excised can be immediately confirmed, i.e., without drawing out the catheter 10 from the blood vessel 50. Therefore, when the excision of the atheroma 51 is insufficient, for example, the support portion 44 of the IVUS shaft 16 is located at the second position again, and then the remaining atheroma 51 can be excised by the cutter 12. The collection of the tomographic images of the blood vessel 50 may be performed by electrically igniting the ultrasound probe 17 while moving the IVUS shaft 16 in the axial direction 101 with respect to the shaft 11. Thus, the tomographic images are continuously collected in the length direction (which is almost in agreement with the axial direction 101) of the blood vessel 50. Then, when the excision of the atheroma 51 is completed, the balloon 23 is contracted, and then the catheter 10 is drawn out of the blood vessel 50 to be collected.

#### Operational Effects of First Embodiment

According to the catheter 10 of the first embodiment, the atheroma 51 in the blood vessel 50 can be excised and the ultrasonic images of the blood vessel 50 can be obtained by the simple structure.

Moreover, since the IVUS shaft 16 can be relatively moved in the axial direction 101 with respect to the shaft 11 and the ultrasound probe 17 can be moved in the length direction (axial direction 101) in the blood vessel 50 in connection with the relative movement, the ultrasound probe 17 can be moved with respect to the atheroma 51, so that tomographic images different in the position in the length direction of the blood vessel 50 can be obtained.

Moreover, the notch portion 43 is provided in the IVUS shaft 16, and therefore even when the support portion 44 provided on the side opposite to the notch portion 43 is located at the first position where the support portion 44 covers the opening 20, the balloon 23 which is outwardly expanded and the IVUS shaft 16 do not abut on each other by the notch portion 43. Therefore, in the support portion 44 at the first position, tomographic images including the atheroma 51 at a position facing the opening 20 can be obtained.

#### Modification of First Embodiment

In the first embodiment described above, although the ultrasound probe 17 is provided in the IVUS shaft 16, the IVUS shaft 16 may not be provided and the ultrasound probe 17 may be provided on the outer peripheral surface of the shaft 11.

As illustrated in FIG. 5, the ultrasound probe 17 is provided in the vicinity of the opening 20 in the shaft 11 and is provided on the distal end side relative to the opening 20. In this modification, the ultrasound probe 17 and the balloon 23 are arranged in such a manner as not to interfere with each other, the ultrasound probe 17 may have 32 elements corresponding to the number of the elements provided on the half of the circumference of the shaft 11 as in the first embodiment described above or may have 64 elements corresponding to the number of the elements provided on the entire circumference of the shaft 11. Although not illustrated in FIG. 5, the cable 45 extended from the ultrasound probe 17 is inserted into and passed through the internal space of the shaft 11 to be connected to the control device 18. As illustrated in FIG. 6, the ultrasound probe 17 may be provided in the vicinity of the opening 20 in the shaft 11 and on the proximal end side relative to the opening 20.

Since the IVUS shaft 16 is not provided according to such a modification, a reduction in size, particularly a reduction in diameter, of the catheter 10 is achieved, and the structure becomes simpler. By deflating the balloon 23 and moving the ultrasound probe 17 in the length direction of the blood vessel 50 together with the catheter 10, tomographic images of the blood vessel 50 containing the atheroma 51 and tomographic images continuing in the length direction of the blood vessel 50 can be collected. Moreover, due to the fact that the ultrasound probe 17 is provided over the entire circumference of the shaft 11, tomographic images of the entire circumference of the blood vessel 50 can be collected.

#### Second Embodiment

As illustrated in FIGS. 7 and 8, a catheter 210 has a shaft 211 (which is equivalent to the tube body), a torque shaft 216 and a cutter 212 provided in the shaft 211, a distal end portion 213 configuring the distal end of the shaft 211, a proximal end portion 214 connected to the proximal end of the shaft 211, an actuator 215 which gives rotation to the cutter 212, an OCT image wire 217 (which is equivalent to the light guide material) and a reflective material 218 provided in the torque shaft 216, and a guide wire tube 219. The catheter 210 is used as a medical instrument which is inserted into a blood vessel to excise an atheroma or photographs ultrasonic images of a blood vessel.

The shaft 211 is a tube capable of containing the cutter 212 therein. The shaft 211 is configured from a surgical stainless steel circular tube or a synthetic resin circular tube, for example, and has flexibility which allows the shaft 211 to be elastically curved corresponding to the curved shape of a blood vessel. The distal end and the proximal end of the shaft 211 each are opened. The outer diameter of the shaft 211 is set according to the inner diameter of a blood vessel into which the shaft 211 is to be inserted, e.g., coronary arteries. The inner diameter of the shaft 211 is set according to the outer diameter of the cutter 212. The outer diameter and the inner diameter of the shaft 211 are almost uniform over the axial direction 101 of the shaft 211. The length in the axial direction 101 of the shaft 211 is set in consideration of the length from catheter insertion portions, such as the human limbs, to the affected portion.

As illustrated in FIG. 8, an opening 220 is formed in the vicinity of the distal end portion 213 in the shaft 211. The opening 220 is formed by cutting out a part of the side wall of the shaft 211. The shape and the size of the opening 220 are set in consideration of the shape and the size of an atheroma which is probably formed in the affected portion.

As illustrated in FIG. 8, the cutter 212 is provided in the vicinity of the opening 220 in the internal space of the shaft 211. The cutter 212 has an approximately cylindrical shape and the outer diameter is a little smaller than the inner diameter of the shaft 211. Therefore, the cutter 212 can move along the axial direction 101 in the internal space of the shaft 211. On the distal end side of the cutter 212, a plurality of blades are formed in such a manner as to radially extend from the center. Although not illustrated in FIG. 8, a through-hole is formed along the axial direction 101 in the center of the cutter 212.

The torque shaft 216 is inserted into and passed through the internal space of the shaft 211. The distal end side thereof is connected to the cutter 212, and the proximal end side thereof is extended to the outside of the proximal end portion 214 of the shaft 211. The torque shaft 216 has flexibility which allows the torque shaft 216 to be elastically curved corresponding to the curved shape of a blood vessel together with the shaft 211 and has torsional rigidity which transmits rotation around the axial direction 101. The torque shaft 216 is configured to form a tube shape as a whole by continuing surgical stainless steel in a spiral shape, for example.

As illustrated in FIG. 7, the torque shaft 216 is connected to the actuator 215. Due to the fact that the torque shaft 216 receives drive transmitted from the actuator 215 to rotate, the cutter 212 rotates. Due to the fact that the torque shaft 216 is moved in the axial direction 101, the cutter 212 moves in the axial direction 101 in the internal space of the shaft 211.

As illustrated in FIG. 8, an opening 221 is formed in the vicinity of the distal end connected to the cutter 212 in the torque shaft 216. The opening 221 is formed by cutting out a part of the side wall of the torque shaft 216. The shape and the size of the opening 221 are set in consideration of the shape and the size of near-infrared rays which are reflected by the reflective material 218 to be emitted in a direction orthogonal to the axial direction 101.

As illustrated in FIG. 8, an OCT image wire 217 is inserted into and passed through the internal space of the torque shaft 216 from the proximal end portion 214 to be extended to the opening 221. The inner diameter of the internal space of the torque shaft 216 is equivalent to the outer diameter of the OCT image wire 217. Therefore, the axis of the OCT image wire 217 and the axis of the torque shaft 216 are almost in agreement with each other. Although not illustrated in detail in each figure, the OCT image wire 217 is one in which an optical fiber is built in a transparent outer casing and which is provided with a lens emitting near-infrared rays on a distal end portion. The near-infrared rays emitted from the lens are emitted along the axial direction of the OCT image wire 217. The OCT image wire 217 propagates near-infrared rays to be supplied from a light source built in an OCT body display portion 222 to the distal end side.

In the internal space of the torque shaft 216, the reflective material 218 is disposed facing the distal end of the OCT image wire 217 in the axial direction 101. In the reflective material 218, a reflective surface 225 facing the distal end of the OCT image wire 217 is a surface inclined to form an angle of 45° with respect to the axis of the OCT image wire 217. The reflective surface 225 is exposed to the outside of the torque shaft 216 through the opening 221 of the torque shaft 216. The reflective material 218 is a columnar body containing an optical fiber, resin, or the like. The outer diameter thereof is equivalent to the inner diameter of the internal space of the torque shaft 216. Therefore, the axis of the reflective material 218 and the axis of the torque shaft

**216** are almost in agreement with each other. On a surface including the reflective surface **225** of the reflective material **218**, metal layers are laminated. The metal layer is formed by, for example, plating or sputtering nickel, gold, aluminum, chromium, and the like alone or a mixture thereof onto the surface of the reflective material **218**. By the reflective surface **225**, near-infrared rays emitted along the axial direction **101** from the OCT image wire **217** are reflected in a direction (second direction) orthogonal to the axial direction **101** to be emitted to the outside of the torque shaft **216** through the opening **221**. The angle of the reflective surface **225** with respect to the axis of the OCT image wire **217** is an example and may not necessarily be 45°. More specifically, the near-infrared rays to be emitted along the axial direction **101** from the OCT image wire **217** may be reflected in a direction crossing the axial direction **101**.

The OCT image wire **217** and the reflective material **218** can rotate around the axis (axial direction **101**) integrally with the torque shaft **216** and can slide in the axial direction **101** in a state of holding the mutual positional relationship, i.e., the clearance and the angle of the reflective surface **225**. The rotation and the slide of the OCT image wire **217** and the reflective material **218** are controlled by directly or indirectly operating the proximal end side of the torque shaft **216** extended from the proximal end portion **214**. Specifically, driving force from the actuator **215** is given to the proximal end side of the torque shaft **216**, whereby the torque shaft **216** is rotated and slid.

As illustrated in FIGS. 7 and 8, a balloon **223** is provided at a position opposite to the opening **220** with respect to the axis of the shaft **211**. The balloon **223** can outwardly expand from the side wall of the shaft **211** and is folded and stuck to the side wall of the shaft **211** until the catheter **210** is inserted into a blood vessel. As raw materials of the balloon **223**, materials having biocompatibility are preferable. Specific examples of the materials include polyurethane, polyethylene, polyester, polypropylene, polyamide, polyamide elastomer, polytetrafluoroethylene, polyvinylidene fluoride, and the like.

As illustrated in FIG. 8, the proximal end side of the balloon **223** is connected to a tube for balloon **224** provided along the side wall of the shaft **211**. The internal space of the tube for balloon **224** is caused to communicate with the internal space of the balloon **223**. The tube for balloon **224** is extended to the proximal end portion **214** and the internal space of the tube for balloon **224** is connected to a port **241** of the proximal end portion **214**. Due to the fact that liquid, such as physiological saline, injected from the port **241** of the proximal end portion **214** flows into the balloon **223**, the balloon **223** is expanded in a blood vessel. The tube for balloon **224** is a molded body of flexible plastic which can be elastically deformed, such as polyamide, polyamide elastomer, and polyetheramide.

As illustrated in FIGS. 7 and 8, the distal end portion **213** is connected to the distal end of the shaft **211**. As illustrated in FIG. 8, the distal end portion **213** has a blade tube **231**, a reduced diameter portion **232**, and a distal end tip **233**.

As illustrated in FIGS. 7 and 8, the blade tube **231** is a circular tube in which both sides are opened. The blade tube **231** is connected to the distal end of the shaft **211** and the internal space thereof is caused to communicate with the internal space of the shaft **211**. The blade tube **231** is one in which flexible plastic which can be elastically deformed, such as polyamide, polyamide elastomer, and polyetheramide, is reinforced with a core material **234**. The core material **234** is embedded in the side wall of the blade tube **231**. The core material **234** is formed into a cylindrical shape

by braiding wire rods, such as surgical stainless steel, into a mesh. The inner diameter of the blade tube **231** is almost equal to the outer diameter of the shaft **211** and the blade tube **231** is fitted to the distal end of the shaft **211** from the outside. The outer diameter and the inner diameter of the blade tube **231** are almost uniform over the axial direction **101**. In each figure other than FIG. 7, the core material **234** is not illustrated.

As illustrated in FIGS. 7 and 8, the reduced diameter portion **232** is a circular tube in which the blade tube **231** side is opened and the outer diameter decreases toward the distal end in a tapered shape. The reduced diameter portion **232** is connected to the distal end of the blade tube **231** and the internal space thereof is caused to communicate with the internal space of the blade tube **231**. The reduced diameter portion **232** contains flexible plastic which can be elastically deformed, such as polyamide and polyetheramide. The inner diameter on the proximal end side of the reduced diameter portion **232** is almost equal to the outer diameter of the distal end of the blade tube **231**. The reduced diameter portion **232** is fitted to the distal end of the blade tube **231** from the outside, and then thermally fused thereto. The distal end side of the reduced diameter portion **232** is sealed. The thickness decreases toward the distal end side on the distal end side of the reduced diameter portion **232**.

As illustrated in FIGS. 7 and 8, the distal end tip **233** is connected to the distal end of the reduced diameter portion **232**. A distal end **235** of the distal end tip **233** is projected from the distal end of the reduced diameter portion **232** to the outside in the axial direction **101**. The distal end tip **233** contains flexible plastic which can be elastically deformed, such as polyamide and polyetheramide. The distal end tip **233** is thermally fused on the distal end side of the reduced diameter portion **232**. On the distal end of the distal end tip **233**, a marker which can be confirmed by X-rays or the like may be provided.

As illustrated in FIG. 7, a guide wire tube **219** is provided over the distal end tip **233**, the blade tube **231**, and a part of the distal end side of the shaft **211** along the outer peripheral surface of each of the distal end tip **233**, the blade tube **231**, and the distal end of the shaft **211**. The guide wire tube **219** is a molded body of flexible plastic which can be elastically deformed, such as polyamide, polyamide elastomer, and polyetheramide. The guide wire tube **219** is disposed at a position where the guide wire tube **219** is not overlapped with the opening **220** of the shaft **211** and the balloon **223**, e.g., in the vicinity of the boundary between the shaft **211** and the balloon **223**. The internal space of the guide wire tube **219** is a guide wire lumen, and a guide wire which is not illustrated is inserted into and passed through the guide wire lumen.

The proximal end portion **214** is provided on the proximal end of the shaft **211**. The proximal end portion **214** is a cylindrical member having an internal space continuing to the internal space of the shaft **211**. The proximal end portion **214** is a molded body of resin, such as polypropylene or ABS. The proximal end portion **214** may serve as a handle in an operation of inserting and removing the shaft **211** into/from a blood vessel.

The proximal end portion **214** is provided with the port **241** extended in a direction crossing the axial direction **101**. Another device, such as a syringe, is connected to the port **241**, and then fluid, such as physiological saline, which is flown in and out from the device, flows in and out of the tube for balloon **224** from the proximal end portion **214**. The proximal end portion **214** may be provided with another port continuing to the internal space of the shaft **211**. Such a port

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is used for the purpose of, for example, collecting an excised atheroma entering the inside of the shaft 211.

From an opening on the proximal end side of the proximal end portion 214, the torque shaft 216 is extended. The actuator 215 is connected to the torque shaft 216. In the actuator 215, a motor, a battery, and the like are built. To the torque shaft 216, rotation of the motor of the actuator 215 is transmitted.

The OCT image wire 217 disposed in the internal space of the torque shaft 216 is connected to the OCT body display portion 222 through the actuator 215. The OCT body display portion 222 has a light source which supplies near-infrared rays, an interferometer, a portable reference mirror, a monitor, an arithmetic unit, and the like. Near-infrared rays supplied from the light source are split by the interferometer to be supplied to each of the OCT image wire 217 and the portable reference mirror. Then, near-infrared rays reflected in a blood vessel and near-infrared rays reflected on the portable reference mirror are combined in the interferometer. Interference signals of the near-infrared rays are treated by the arithmetic unit to be displayed as tomographic images of the blood vessel on a monitor.

[Usage Directions for Catheter 210]

Hereinafter, the usage directions for the catheter 210 are described with reference to FIGS. 9 and 10.

The catheter 210 is used when excising an atheroma 51 formed in the inner wall of a blood vessel 50. The position of the atheroma 51 is confirmed by the blood vessel tomographic images by the OCT. The catheter 210 is inserted into the blood vessel 50 from the distal end portion 213 in the state (FIG. 7) where the balloon 223 is contracted. Although not illustrated in each figure, a guide wire is inserted into the blood vessel 50 beforehand in the insertion of the catheter 210 into the blood vessel 50. The insertion of the guide wire into the blood vessel 50 is performed by a known technique. The catheter 210 is inserted into the blood vessel 50 from the distal end portion 213 while inserting the guide wire inserted into the blood vessel 50 into a guide wire tube 219.

The distal end portion 213 is advanced to the atheroma 51 in the blood vessel 50 while being elastically curved along the guide wire at a portion where the blood vessel 50 is curved, such as coronary arteries. When the distal end portion 213 reaches the atheroma 51, and the opening 220 of the shaft 211 faces the atheroma 51, the insertion of the shaft 211 into the blood vessel 50 is ended. By rotating the torque shaft 216 by the actuator 215 and supplying near-infrared rays to the OCT image wire 217 from the OCT body display portion 222, tomographic images of the blood vessel 50 are displayed in the OCT body display portion 222. By confirming the tomographic images of the blood vessel 50, it can be confirmed that the opening 220 has reached the position corresponding to the atheroma 51 and the state of the atheroma 51 can be confirmed. Thereafter, the guide wire is drawn out of the proximal end portion 214 side of the catheter 210. The actuator 215 is connected to the torque shaft 216 of the cutter 212. In FIGS. 9 and 10, the guide wire is omitted.

In the state where the opening 220 of the shaft 211 faces the atheroma 51 as illustrated in FIG. 9, the balloon 223 in the contracted state is expanded by fluid which is caused to flow into the tube for balloon 224 from the port 241. When the expanded balloon 223 abuts on the inner wall of the blood vessel 50 on a side opposite to the atheroma 51, the opening 220 is stuck to the atheroma 51, so that the atheroma 51 partially enters the internal space of the shaft 211 from the opening 220. In the state above, the catheter 210 is fixed to the blood vessel 50.

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Subsequently, the rotation of the actuator 215 is transmitted to the cutter 212 through the torque shaft 216 of the cutter 212, so that the cutter 212 is rotated. Due to the fact that the torque shaft 216 is advanced to the distal end side in the axial direction 10 with respect to the shaft 211 on the proximal end portion 214 side, the rotating cutter 212 abuts on the atheroma 51, so that the atheroma 51 is excised by the cutter 212. Fragments 52 of the excised atheroma 51 enter the internal space of the blade tube 231 through the internal space of the shaft 211.

After the atheroma 51 is excised by the cutter 212, tomographic images of the blood vessel 50 in the state where the atheroma 51 is excised can be obtained by the same OCT as above. Thus, the state where the atheroma 51 is excised can be immediately confirmed, i.e., without drawing out the catheter 210 from the blood vessel 50. Therefore, when the excision of the atheroma 51 is insufficient, for example, the remaining atheroma 51 can be excised by rotating the cutter 212. The collection of the tomographic images of the blood vessel 50 may be performed while moving the reflective material 218 together with the torque shaft 216 in the axial direction 101 with respect to the shaft 211. Thus, the tomographic images continuing in the length direction (which is almost in agreement with the axial direction 101) of the blood vessel 50 can be obtained. Then, when the excision of the atheroma 51 is completed, the balloon 223 is contracted, and then the catheter 210 is drawn out of the blood vessel 50 to be collected.

#### Operational Effects of Second Embodiment

According to the catheter 210 of the second embodiment, the atheroma 51 in the blood vessel 50 can be excised and the ultrasonic images of the blood vessel 50 can be obtained by the simple structure.

Moreover, since the OCT image wire 217 and the reflective material 218 are disposed in the internal space of the torque shaft 216, near-infrared rays can be guided to the vicinity of the atheroma 51 in the blood vessel 50 to be reflected in the torque shaft 216.

Moreover, since the OCT image wire 217 and the reflective material 218 can be rotated integrally with the torque shaft 216, the excision of the atheroma 51 and the OCT can be realized by controlling the number of rotations of the motor in the actuator 215.

Moreover, since the OCT image wire 217 and the reflective material 218 can move along the axial direction 101 integrally with the torque shaft 216, tomographic images of the blood vessel 50 along the axial direction 101 can be obtained.

Moreover, since the OCT image wire 217 and the reflective material 218 are disposed in the internal space of the torque shaft 216, the OCT image wire 217 and the reflective material 218 are covered with the torque shaft 216 to be protected.

Moreover, since the guide wire tube 219 is provided along the shaft 211, the catheter 210 can be inserted into the blood vessel 50 along a guide wire.

#### Modification

In the embodiments described above, although the guide wire tube 219 has a so-called rapid exchange type structure in which the guide wire tube 219 is disposed on the outside of the shaft 211. However, as illustrated in FIG. 11, a

so-called over-the-wire type structure in which the internal space of the torque shaft **216** is used as a guide wire lumen may be adopted.

When the over-the-wire type structure is adopted, the OCT image wire **217** and the reflective material **218** are disposed in such a manner as to face each other with respect to the axial direction **101** along the outer peripheral surface of the torque shaft **216** as illustrated in FIG. **11**. Near-infrared rays to be emitted along the axial direction **101** from the OCT image wire **217** are reflected by the reflective material **218** in a direction (direction orthogonal to) crossing the axial direction **101** toward the outside of the torque shaft **216**. Thus, the near-infrared rays are not emitted to the guide wire inserted into and passed through the internal space of the torque shaft **216**. In the over-the-wire type structure, the opening **221** does not need to be formed in the torque shaft **216**.

When the over-the-wire type structure is adopted, a circular tube, in which both sides are opened and the outer diameter of a proximal end **236** side increases in a tapered shape as illustrated in FIG. **11** is adopted as the distal end tip **233**. The distal end tip **233** is connected to the distal end of the reduced diameter portion **232** and the internal space thereof is caused to communicate with the internal space of the reduced diameter portion **232**. A distal end **235** of the distal end tip **233** is projected from the distal end of the reduced diameter portion **232** to the outside in the axial direction **101**. The proximal end **236** side of the distal end tip **233** is extended in the axial direction **101** in the internal space of the reduced diameter portion **232** and the proximal end **236** reaches the internal space of the blade tube **231**. More specifically, a portion including a distal end side portion of the blade tube **231** and the reduced diameter portion **232** has a double tube structure in which the blade tube **231** and the reduced diameter portion **232** serve as the outer side and the distal end tip **233** serves as the inner side.

Although the diameter of the distal end tip **233** is increased on the proximal end **236** side, the other outer diameter and the inner diameter of the other portion are almost uniform. The outer diameter of the uniform portion is smaller than the inner diameter of the blade tube **231** and is almost equal to the inner diameter of the distal end of the reduced diameter portion **232**. Although the diameter of the proximal end **236** side is increased, the maximum diameter is smaller than the inner diameter of the blade tube **231**.

A through-hole along the axial direction **101** is formed in the cutter **212**. A guide wire lumen is formed along the axial direction **101** of the shaft **211** by the internal space of the torque shaft **216**, the through-hole of the cutter **212**, the internal space of the blade tube **231**, and the internal space of the distal end tip **233**.

The distal end surface of the OCT image wire **217** may be a surface inclined to form an angle of 45° with respect to the axial direction **101** and the reflective material **218** may be provided on the distal end surface. In that case, the reflective material **218** disposed at a position apart from the OCT image wire **217** is not present.

#### REFERENCE SIGNS LIST

**10** Catheter  
**11** Shaft (Tube body)

**12** Cutter  
**16** IVUS shaft (Outer tube body)  
**17** Ultrasound probe  
**20** Opening  
**23** Balloon  
**43** Notch portion  
**44** Support portion  
**210** Catheter  
**211** Shaft (Tube body)  
**212** Cutter  
**216** Torque shaft  
**217** OCT image wire (Light guide material)  
**218** Reflective material  
**219** Guide wire tube (Guide wire lumen)  
**220,221** Opening  
**223** Balloon

The invention claimed is:

**1.** A catheter comprising:

- a tube body having an opening in a part of a side wall on a distal end side;
- a cutter which is located in a vicinity of the opening in an internal space of the tube body and which can move in an axial direction of the tube body;
- a balloon which is disposed on a side opposite to the opening with respect to an axis of the tube body and which outwardly expands from the side wall of the tube body; and
- a phased array ultrasound probe disposed along a circumferential direction of an outer peripheral surface of the side wall in the vicinity of the opening at least on a same side as a side where the opening is provided with respect to the axis of the tube body.

**2.** The catheter according to claim **1**, further comprising: an outer tube body which is provided on an outside of the tube body and which can relatively move in the axial direction with respect to the tube body, wherein the ultrasound probe is provided in the outer tube body.

**3.** The catheter according to claim **2**, wherein the outer tube body has:

- a notch portion in which a part corresponding to the balloon is cut out; and
- a support portion which is disposed on a side opposite to the notch portion with respect to the axis of the outer tube body and which can be moved to a first position where the support portion covers the opening and a second position where the support portion opens the opening by the relative movement, wherein the ultrasound probe is provided on the support portion, and

the notch portion does not abut on the balloon in a state of being outwardly expanded at the first position.

**4.** The catheter according to claim **1**, wherein the ultrasound probe is fixed on a distal end side relative to the opening in the tube body.

**5.** The catheter according to claim **1**, wherein the ultrasound probe is fixed on a proximal end side relative to the opening in the tube body.

\* \* \* \* \*

专利名称(译)	导管		
公开(公告)号	<a href="#">US10405881</a>	公开(公告)日	2019-09-10
申请号	US15/116562	申请日	2015-02-04
[标]申请(专利权)人(译)	尼普洛株式会社		
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IPC分类号	A61B8/00 A61B17/3207 A61B17/22 A61B90/00 A61B8/12		
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优先权	2014020927 2014-02-06 JP 2014104489 2014-05-20 JP		
其他公开文献	US20160354111A1		
外部链接	<a href="#">Espacenet</a>		

# 摘要(译)

[目的]提供一种导管，其具有能够切除血管中的动脉粥样硬化并且能够获得血管的超声图像的简单结构。[解决方案]导管 10 具有轴 11，其在远端侧的侧壁的一部分中具有开口 20 12 位于轴 11 的内部空间中的开口 20 附近且可沿轴向移动 > 101 的轴 11，气球 23 设置在相对于开口 20 的相对侧的气囊上。轴 11 的轴线从轴的侧壁向外扩展 11 沿着圆周设置的相控阵超声探头 17 开口 20 附近的侧壁的外周表面的方向 102 20 的一侧相对于轴11的轴线设置。

