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Description**FIELD OF INVENTION**

The present invention relates generally to medical devices directed to the prevention of luminal occlusion, and more particularly to stents and methods for making and utilizing these stents in the treatment of both benign and malignant conditions wherein the functionality of the stents is determined by geometrical variability of the scaffolding and concomitant interstices.

BACKGROUND OF THE INVENTION

Stents are devices that are inserted into a vessel or passage to keep the lumen open and prevent closure due to a stricture, external compression, or internal obstruction. In particular, stents are commonly used to keep blood vessels open in the coronary arteries and they are frequently inserted into the ureters to maintain drainage from the kidneys, the bile duct for pancreatic cancer or cholangiocarcinoma or the esophagus for strictures or cancer. Vascular as well as non-vascular stenting has evolved significantly; unfortunately there remain significant limitations with respect to the technology for producing stents suitable to various portions of a patient's anatomy.

Historically, in order to provide a stent with varying characteristics, the stent had to be manufactured from multiple materials, at least one for each characteristic desired. As a result, many of these stents are woven from two or more metals having differing shape-memories for example. Unfortunately, braided stents are vulnerable to premature obsolescence. Moreover, providing multiple material types in a single stent may lead to inconsistent characteristics along the surface area of the stent. This is particularly undesirable when the stent is to be placed in vascular or nonvascular lumens that have been occluded for one reason or another. The stent needs to be stiffer in some regions while more flexible in others.

Additionally, medical device companies have identified the need to cover stents at least partially to prevent the epithelialization of the scaffolding. Most covered stents however have an elastomeric cover that is subject to bunching particularly about stenotic tissue. This can lead to additional tissue granulation.

Additionally the ends of the stent tend to be exposed in order to encourage granulation tissue formation, which helps to anchor the stent in place. With metal stents, the direct metal to tissue contact accelerates tissue granulation and galvanic current generation is also an undesirable byproduct. Such direct current can have indirect effects on tissue granulation and direct effects on fluid flow dynamics.

Moreover, since many medical device companies have chosen to use poorly adapted cardiovascular stents for Pulmonary, GI and Peripheral Vascular indications, many of the anatomical differences in the lumens are not accounted for in stent design. For example, the pulsation of the cardiovascular lumen and the concomitant radial force requirements of a cardiovascular stent differ substantially from that of a tightly constricted lumen such as the trachea during repeated coughing. When a stent developed for the former is indicated for the latter, the stent tends to fail under the extreme conditions and lose its elasticity and therefore its ability to ensure airway patency. Non-vascular lumens also tend to have ciliated epithelia so as to facilitate clearance of fluids and particulates. As a general principal, covered stents were not specifically designed for coated lumens in that the external covering damages the cilia and prevents the body's natural clearing function. Moreover, the covering itself is usually made of a predominately hydrophilic polymer, which can lead to mucous formation and/or fluid stagnation. Stagnation of fluids or material passing through the lumen can lead to additional complications such as in stent restenosis or bacterial infections.

The document WO-A-02/30329 (Rex Medical, L.P.) describes a system for treating stenosis in a target blood vessel, such as the common carotid artery, comprising a graft portion having a main portion and a branch portion extending therefrom. The branch portion extends from the intermediate portion of the main portion at an angle thereto and is in fluid communication with the main portion. A first stent is associated with the main portion and is expandable from a first configuration to a second configuration to retain the main portion in position within the target vessel. A second stent is associated with the branch portion and is expandable from a first configuration to a second configuration to retain the branch portion in position within a branching vessel. In certain described embodiment, the stent has a graft material or layer on the inside or outside of the stent.

There remains an existing need for a therapeutic stent that can have varying characteristics along its surface area while being stamped, not braided, from a single base material. Moreover, there is a need for such a therapeutic stent where the relative hardness, softness, flexibility, stiffness and radial force can be modified as a function of geometric considerations rather than material considerations. In particular, there is a need for a stent that is divided into zones so as to allow the stent to have predetermined characteristics in one zone and could conceivably have drastically different characteristics in an adjacent zone so as to allow for stents that can be tailored to anatomical lumens in general and the particular lumen topography of a specific patient in particular. An additional need remains for a stent that is covered in a manner that limits the amount of covering surface area that is in direct contact with the target lumen. In particular, there is a need for a covered stent that is preferably covered internally such that the outer scaffolding surface of the stent is raised from the outer surface of the cover. To this end, cilia function is only partially limited and mucociliary clearance is not significantly affected. A need also remains for a cover that itself has anti-adherent properties or is complexed with an anti-adherent such that bacteria, fungi or other microbes cannot colonize the cover in particular and the stent generally.

WO03/057075 discloses a stent which is dip-coated or coated with inner and outer liners such that the struts of the stents are surrounded by the cover.

SUMMARY OF EXEMPLARY EMBODIMENTS

The invention is defined in the appended claims.

It is a preferred purpose of the present invention to provide a stent, in accordance with an exemplary embodiment of the present invention, which combines many of the excellent characteristics of both silicone and metal stents while eliminating the undesirable ones. In particular, it is an objective of a preferred embodiment in accordance with the present invention to provide a stent that is easily installed, yet in alternative embodiments, removable. Moreover the stent in accordance with this embodiment of the present invention would not cause material infections and may be capable of reducing infection. Therefore, a principal objective of a preferred embodiment in accordance with the present invention is to provide a prosthesis that is suitable for both permanent and temporary use while being easy to insert, reposition

and remove.

A principal objective of an example is to provide a stent that may be stamped from preferably a single material that is capable of maintaining its axial working length when radially compressed. To this end, the stent does not have a seam that could aggravate luminal tissue. In particular, a stent in accordance with the present invention is formed using a tool that molds the stents outer contour as well as its interstices.

In another example a stent is provided that can be indicated for the treatment of benign and malignant disease and improve the way clinicians treat malignant obstructions.

Still another example is to provide a stent and method for installing the stent that is economical and suitable for routine purposes. Moreover, the stent will have minimal migration, cause minimal tissue granulation, will not foreshorten after deployment and mucociliary clearance will not be problematic.

Yet another example is to provide a prosthesis that will have superior internal to external diameter ratio, superior radial force with dynamic expansion, while being suitable for use in pediatric and adult patients with malignant and benign disease.

In another example a family of stents is provided where the relative hardness/softness of regions of the stent can differ from other regions of the stent to provide additional patient comfort and resistance to radial forces.

In another example a family of stents is provided with novel interstice configurations that facilitate flexibility, durability and/or proper installation.

In another example a self-expanding stent is provided having the above benefits that also defines a plurality of apertures about the termini of the stent for, inter alia, removal of the stent.

In a preferred embodiment of the present invention a prosthesis is provided that minimizes cilia destruction at the site of implantation. In the furtherance of this and other objectives, the preferred prosthesis is covered internally with a polyurethane such that the surfaces of the struts that come into contact with the lumen of the patient are elevated above the surface of the covering such that the cilia can move to allow for free fluid action of ciliated epithelium.

Further objectives, features and advantages of the invention will be apparent from the following detailed description taken in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE FIGURES

FIG. 1 shows a polarization microscopic image of a plurality of zones of a stent with a cover in accordance with a preferred embodiment of the present invention.

FIG. 2 shows an alternative perspective view of the polarization microscopic image of FIG. 1.

FIG. 3 shows an enlarged perspective view of the interstices of an exemplary zone as shown in FIG. 1.

DETAILED DESCRIPTION OF AN EMBODIMENT

An exemplary embodiment of the stent, in accordance with the present invention, provides a stent that prevents epithelialization of the stent and is not subject to premature elongation and foreshortening but is capable of engaging the desired implantation location. The stent also retains its axial length while undergoing radial compression.

The stent is preferably formed from a composite material selected from the group consisting essentially of Ni, C, Co, Cu, Cr, H, Fe, Nb, O, Ti and combinations thereof. The composite material is generally formed into a compressed tube from which the stent is etched and is formed on a suitable shaping device to give the stent the desired external geometry. Both the synthetic collar techniques and in vitro valuation techniques show the remarkable ability of stents in accordance with the present invention to convert acting force into deformation work absorbed by the angled structure, which prevents excessive scaffolding stress and premature material fatigue and accelerated obsolescence.

Though one skilled in the stent engineering art, once apprised of the present application, would be able to manufacture a stent consistent with the present invention by other methods, a preferred method of manufacturing such stents follows. As stated above a composite material is selected and a blank is formed there from. The blank is preferably laser etched and the etch work is generally verified for accuracy using visual recording microscopy. Dimensional measurements are taken to ensure strut thickness, segment angles, zone placement, etc. Moreover, the stent is preferably formed on a shaping tool that has substantially the desired contour of the external stent dimensions.

In the event the stent is to be shaped to the dimensions of a particular lumen, optical photography and/or optical videography of the target lumen may be conducted prior to stent formation. The geometry of corresponding zones and connector regions of the stent then can be etched and formed in accordance with the requirements of that target lumen. For example, if the stent were designed for the trachea, which has a substantially D shaped lumen and additionally the middle zones needed to be softer than the end zones, the stent could be designed to those specifications. With specific reference being made to FIG. 1, it can be seen that angles α , β , δ , ϵ and γ may be modified to provide different characteristics to different zones of the stent. In particular, if the topography of the trachea of a particular patient is captured optically and the appropriate dimension provided, a patient specific prosthesis could be engineered. These techniques can be adapted to other non-vascular lumen but is very well suited for vascular applications where patient specific topography is a function of a variety of factors such as genetics, lifestyle, etc.

It should be pointed out that unlike the use of differing shape memory materials to change regions of a stent, stents in accordance with the present invention can take on an infinite number of characteristic combinations as zones and segments within a zone can be modified by changing angles, segment lengths and segment thicknesses during the etching and forming stages of stent engineering or during post formation processing and polishing steps. Moreover, by modifying the geometry of the connectors between zones, additional functionality may be achieved.

Exemplary stents 10 are shown in FIGS. 1-3 showing the preferred interstice geometry. Not shown are a wide variety of interstice geometries that are also acceptable alternatives to the preferred, namely, U, V, W, Z, S and X geometries to name a few.

The stent 10 also is formed of memory metal and preferably has unique geometrical interstices that are laser etched therein. However, other conventional ways of forming interstices in unitary stents, though not equivalent are contemplated, may be employed and would be within the skill set of one in the art.

It cannot be overemphasized, however, that this does not mean the knowledge that changes in the geometry of interstices affect stent functionality is currently known in the art. To the contrary, the present inventors discovered the interrelation between interstice geometry, width, length and relative resistance to torsional stress and radial force. In fact, it can be said that the stent 10 has circumferential bands extending perpendicularly with respect to the luminal device's longitudinal axis. These bands are referred to generally as zones. A connector 50 connects these bands to one another; the connector 50 is an additional means for adjusting stent functionality. In particular, the connector 50 defines a substantially U shaped member, but could define other geometries such as U, V, W, Z, S and X to name a few. As shown specifically in FIG. 1, δ , ϵ and γ vary in shape and that the corresponding region of the stent differs in function. It can also be seen from FIG. 1, at least one but preferably a plurality of eyelets ϕ that allow a physician to purse string the stent with suture to facilitate removability. The eyelets are preferably between about 200 μ m and 300 μ m, however, the eyelets may be smaller or larger to accommodate the need of the target site. The preferred eyelet size is about 350 μ m as the preferred suture type is 4-0. The medical appliance may be pre-threaded with suture or the user may provide the suture after implantation.

In a standard orientation, as shown particularly in FIG. 2, the substantially U-shape connector comprises preferably two leg members 52 & 56 and a crossing member 54 that connects with and extends perpendicularly at preferably 90° angles with respect to the leg members 52 & 56. It must be noted that alternative angles may be provided without departing materially from the invention. The present inventors discovered that if you modify the length of the crossing member 54 and/or the leg members 52 & 56 and/or the angle γ at which the crossing member 54 and the leg members 52 & 56 intersect, the relative hardness/softness, radial force and/or flexibility of the stent 10 could be modified. The angles γ can be modified at varying acute angles short of 90° or varying obtuse angles greater than 90°. The incremental changes correspondingly change certain characteristics of the stent 10. As a result, different zones of the stent 10 can be given different rigidities to improve patient comfort and for example, in airway stents, to facilitate luminal patency. Moreover, various anatomical lumens may need different degrees of stent rigidity. As a result, stents 10 in accordance with the present invention can be manufactured to exacting specifications to contour properly to various lumens in a patient's anatomy, which may need varying levels of structural support from the medical appliance.

Referring now to FIG. 3, there is an enhanced capability provided by stents. By adjusting the distance between the connector 50 and the zones between which connector 50 resides, the way in which the stent reacts to strain can be modified. By way of non-limiting example, if the connector 40 is oriented closer to one zone than to another zone, the stent will be less flexible and be able to withstand greater radial force. Alternatively, if the connector is equidistant between zones, the stent will be more flexible and be able to withstand less radial force. Please note that these differences are relative to a neutrally located connector 40. The behavior is a function of distance and as a result varies along a continuum with respect to the connector's orientation between the medium between zones and the tip of each zone. Moreover, by varying the number of connectors 40 that connect the zones to one another, functionality can be impacted. In particular, the fewer the number of connectors connecting the zones the more torsional flexibility the stent will have. The converse will generally hold true with respect to a greater number of connectors.

Connector 40, which serves a similar purpose as connector 50 also has a crossing member 44 that connects leg members 42 & 46 at a predetermined angle δ . As discussed above, since form follows function for stents prepared through this novel method, by changing the degrees of angles α , β , δ & γ , stent characteristics can be modified. Moreover, by changing the leg lengths of all the previously discussed legs or individual legs separately, additional stent characteristics can be obtained. The beauty of this system is that the desired characteristics can be determined prior to forming the stent and by staying within certain forming parameters, the stent can be formed, crimped, delivered and deployed with confidence that the desired functionality will result. This is important in light of the fact that both vascular and nonvascular lumen have unique topography. As a result, methods and devices in accordance with the present invention usher in the ability to tailor prosthesis to anatomical tissue in general and particular patient anatomy in particular.

The U shaped connectors 40 & 50 have a crossing member and at least two leg members, respectively. The present inventors discovered that if you increase/decrease the length of leg members and/or increase/decrease the length of crossing members and/or vary the angle at which crossing members and leg members intersect, you affect the functionality of the stent. In particular, the shorter the length of the leg members, the less flexibility available in that portion of the stent. Taking particular note of FIG. 3, by way of example only, if you want to decrease the amount of torsional flexibility of the stent 10, you would have to modify the connector 40 so that the legs 42 & 46 are longer than shown and that the angle δ formed by legs 42 & 46 and crossing member 44, respectively, is slightly greater than 90°. Alternatively, the length of the crossing member 44 can impact the functionality of the stent as well. The stent can be made more rigid by shortening crossing member 44 or the stent may be made more flexible by lengthening crossing member 44. It should be noted that the combination of the changes of leg lengths, crossing member lengths, angle variations, shapes and number of connectors provide the stent with the ability to conform to specific lumens in the anatomy of a patient. The result is a form fitting medical prosthesis that may be tailored to specific anatomical lumens in general and to the anatomical lumens of an individual patient in particular.

The modification of interstice geometries and manipulation of the U shaped connection member to achieve variable stent functionality is provided. The rigidity of the stent scaffolding or interstice matrix along with the torsionality of the stent itself is principally a function of these modifications. In an exemplary embodiment, the stents relative flexibility can be rated soft, medium or hard based on the degree of flex and torsionality. The less torsionality and flex in the stent the harder the stent is rated.

An exemplary stent in accordance with the present invention with relatively great torsionality and radial flexibility would be rated soft. An exemplary soft rated stent comprises distance between U shaped connectors of about 4.5 μ m in the compressed state (i.e., contracted in the 3mm tube subject to laser etching). Moreover, the length of the crossing member is preferably about 1.0 μ m. The lengths of the leg members are preferably about 1.5 μ m long. Additionally the leg members may further comprise feet attachable to the remainder of the stent scaffolding. The feet can be adjusted from a standard length of about 0.25 μ m to further adjust the characteristics of the stent. There is additionally a substantially rectangular member incorporated in the U shaped connector with similar capacity for variability. The variability factors and results of modifying the dimensions of the substantially rectangular members are similar to those evinced by leg length

dimensional modifications.

By way of example, but not to be construed in any way as limiting, the softness index or relative flexibility can be increased by increasing the various lengths discussed above. For example, by increasing the length of the legs and crossing members of the U shaped connector, flexibility increases. However, with respect to the distance between U shaped members and distance between interstices in a preferred stent embodiment, there is an inverse correlation between length and softness. This relative softness/hardness indexing as a corollary of interstice dimensions is a novel aspect of preferred embodiments of the present invention. As a practical rule of thumb, longer leg lengths coupled with acute angles provide for greater flexibility. Conversely, shorter leg lengths and more obtuse angles provide more rigidity. By way of non-limiting example, a U shaped connector with short legs deviating from the crossing member at angles greater than 90°, will be extremely rigid and resistant to torsional strain as compared to a U shaped connector with longer legs diverging from the crossing member at angles less than 90°.

In addition to the length and spacing differences, the interstices themselves may define various shapes that by their very nature afford novel functionality to the stent. The changes of functionality, however, are more a function of the dimensional differences of the various shapes rather than a function of the shapes themselves. Therefore, it is important to keep in mind that the dimensional differences discussed in the previous paragraph are determinative of the functionality accorded the stent by the varying interstice geometries. It is for this reason that one of ordinary skill in the art, after being apprised of the present invention, would be able to conceive of a number of interstice geometries to satisfy certain functionality criteria by keeping certain dimensional parameters constant.

FIGS. 1-3 also show the cover provided in select embodiments in accordance with the present invention. The cover 100 preferably comprises a stable polymeric material such as polyurethane that can be deposited on a stent to form a thin film. The film preferably forms layers when annealed to the stent such that the hydrophobic moieties within the polymer are predominately oriented outward and the hydrophilic moieties are predominately oriented inward. It should be noted that depending on the characteristics desired by the user, the relative hydroaffinity may be altered. For example, in the event the implant was placed with the intention of collecting mucous in the respiratory system, the cover 100 would more suitably have a predominately hydrophobic or partially hydrophilic outer surface. Moreover, by manipulating the hydroaffinity of the cover 100, the physicochemical parameters such as surface-free energy, charge density provide a substantial barrier to biofilm formation in general and ligand-binding events mediated by microbial adhesions and extracellular polymers. However, additional anti-adherents known in the art may be applied to provide lubricity as well as an additional barrier for microbes. For example, a preferred cover 100 in accordance with the present invention would be hydrophilic and hygroscopic to ensure the surface would always appear to be wet which prevents mucostasis as well as microbial adherence.

Regardless of desired covering surface characteristics, preferred stents in accordance with the present invention are covered from the interior of the stent lumen such that the stent scaffolding is raised about between 1 Å to 10⁶ Å above the surface of the covering 100 as shown in FIG. 3 as indicia 200. One of the principal functions of such architecture is to facilitate cilia action by allowing cilia movement between stent struts.

The stent is preferably coated in a multi-step process, which comprises providing a stent and initially spraying the stent with a polymeric material to coat the struts. Though the steps may be reversed it is preferable to follow the spraying step with the interior covering step. In particular, the stent is placed into a hollow mold to retain the stent shape as the internal diameter of the stent is covered with the polymeric material to form a non-porous covering 100. The covering 100 can be provided in sheets or additional spray applications, however, the preferred embodiment is the sheets. Sheets are generally preferred to facilitate the proper orientation of the polymer side chains to ensure that the desired moiety (e.g., hydrophilic and/or hydrophobic) is facing the lumen. Once the layer of polymer is introduced into the inner diameter of the stent, a balloon or other device in which temperature can be regulated is implanted to sandwich the layer of polymer between the stent inner diameter and the balloon. The balloon is expanded and heated to a temperature of about between 93.3°C (200°F) and 204.4°C (400°F) to anneal the polymer to the stent. Preferred polymers such as various designer polyurethanes (e.g., Cronoflex® manufactured by Cardiotech International) are suitable for such applications but other polymers are acceptable.

The present invention may be embodied in other specific forms. The described embodiments are to be considered in all respects only as illustrative, and not restrictive. The scope of the invention is, therefore, indicated by the appended claims, rather than by the foregoing description. All changes, which come within the meaning and range of equivalency of the claims, are to be embraced within their scope.

Patentkrav

1. Medicinsk anordning til anbringelse i en del af anatomien hos en patient, hvilken anordning omfatter:

5 et gittersystem (10), hvilket gittersystem er konfigureret til at definere et i det væsentlige cylindrisk element, der har en distal ende og en proksimal ende og strækker sig på langs derimellem, idet det danner et hulrum derigennem, hvor gittersystemet langs den længdegående udstrækning af
10 anordningen har en indre og en ydre overflade, der omfatter tværstivere med geometriske mønstre, der dannes af vinkler, hvor vinklerne bestemmer anordningens relative fleksibilitet, således at anordningen tilpasser sig til topografien i et målrum, og anordningen, når der påføres et tryk på
15 forskellige punkter i den længdegående udstrækning af denne, ikke forkortes eller forlænges uønsket, hvilken anordning yderligere omfatter:

en polymer coating (100), der dækker i det mindste en del af både tværstiverne og områder mellem tværstiverne, hvor
20 coatingen enten er: (i) en indre coating, der er koblet til anordningen fra den indre overflade af gittersystemet udefter, hvor gittersystemet er yderligere coatet, og overfladerne af de coatede tværstivere, der berører hulrummet hos patienten, er hævet over den indre coating; eller (ii) en ydre coating,
25 der er koblet til anordningen fra den ydre overflade af gittersystemet indefter, hvor de dækkede tværstivere på den ydre overflade af gittersystemet er hævet i forhold til det dækkede område mellem tværstiverne på anordningen.

30 2. Medicinsk anordning ifølge krav 1, hvor coatingen er af en tilstrækkelig tykkelse til at forhindre, at den medicinske anordning bliver epitelialiseret, når den installeres i den ønskede del af patientens anatomi.

35 3. Medicinsk anordning ifølge krav 1 eller 2, hvor coatingen er i det væsentlige hydrofob.

4. Medicinsk anordning ifølge krav 1 eller 2, hvor coatingen

er i det væsentlige hydrofil.

5. Medicinsk anordning ifølge et hvilket som helst af ovennævnte krav, hvor coatingen er hygroskopisk.

5

6. Medicinsk anordning ifølge et hvilket som helst af kavene 1 til 5, hvor coatingen ikke hæmmer bøjning eller radial ekspansion af den medicinske anordning.

10 7. Medicinsk anordning ifølge et hvilket som helst af ovennævnte krav, hvor coatingen er den indre coating, idet de hævede tværstivere er hævet mellem 1 Å og 106 Å i forhold til det coatede område mellem tværstiverne på den medicinske anordning.

15

8. Medicinsk anordning ifølge et hvilket som helst af ovennævnte krav, hvor coatingen er den ydre coating, idet den relative grad, hvormed de coatede tværstivere er hævet i forhold til de coatede områder mellem tværstiverne, er tilstrækkelig til at muliggøre ciliers funktion på implantationsstedet.

9. Medicinsk anordning ifølge et hvilket som helst af ovennævnte krav,
25 hvor dimensionerne i gittersystemets geometri bestemmer vridningsevnen for den medicinske anordning; eller hvor gittersystemet er fremstillet af en hukommelseslegering; eller hvor gittersystemet er elektropoleret; eller hvor den medicinske anordning langs gittersystemets
30 længdegående flade yderligere omfatter et antal kraver, der definerer åbninger derigennem, eller hvor den medicinske anordning i nærheden af den distale og proksimale ende af gittersystemet yderligere omfatter et antal flanger, der definerer åbninger derigennem.

35

10. Medicinsk anordning ifølge et hvilket som helst af ovennævnte krav, der yderligere omfatter en konektor (50), der er koblet til dele af de geometriske mønstre, hvilken

konnektor omfatter et krydselement (54) og et antal benelementer (52, 56), der udgår fra krydselementet.

11. Medicinsk anordning ifølge krav 10, hvor konnektoren
5 yderligere omfatter en rektangulær pal, der udgår fra et ben derpå.

12. Medicinsk anordning ifølge krav 10 eller 11, hvor længden
af benelementerne og størrelsen af vinklen, hvormed benene
10 udgår fra krydselementet, bestemmer den relative fleksibilitet af den medicinske anordning.

13. Medicinsk anordning ifølge krav 12, hvor vinklen, hvormed
benelementerne udgår fra krydselementet, er større end 90° , og
15 fortrinsvis hvor den medicinske anordning er relativt stiv.

14. Medicinsk anordning ifølge krav 12, hvor vinklen, hvormed
benelementerne udgår fra krydselementet er 90° eller derunder,
og fortrinsvis hvor den medicinske anordning er relativt
20 fleksibel.

15. Medicinsk anordning ifølge et hvilket som helst af
ovennævnte krav, der yderligere omfatter en ekstra distal
ende, hvor den medicinske anordning danner en i det væsentlige
25 Y-form.

16. Medicinsk anordning ifølge et hvilket som helst af
ovennævnte krav, hvor mindst én tværstiver definerer en åbning
derigennem.
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17. Medicinsk anordning ifølge krav 16, hvor den mindst ene
åbning definerer et øje med en tilstrækkelig diameter til at
modtage en sutur.

35 18. Medicinsk anordning ifølge krav 16 eller 17, hvor øjediameteren er mindst 300 μm .

19. Medicinsk anordning ifølge et hvilket som helst af

ovenstående krav, hvor gittersystemet langs anordningens længdegående udstrækning danner geometriske mønstre.

20. Fremgangsmåde til coating af en medicinsk anordning, der omfatter trinene:

tilvejebringelse af en form med en indre og en ydre diameter;

tilvejebringelse af en medicinsk anordning, der omfatter et gittersystem (10), hvilket gittersystem er konfigureret til at definere et i det væsentlige cylindrisk element, der har en

distal ende og en proksimal ende og strækker sig på langs derimellem, idet det danner et hulrum derigennem, hvor gittersystemet langs den længdegående udstrækning af anordningen har en indre og en ydre overflade, der omfatter tværstiver med geometriske mønstre, der dannes af vinkler, hvor vinklerne bestemmer anordningens relative fleksibilitet, således at anordningen tilpasser sig til topografien i et målrum, og anordningen, når der påføres et tryk på forskellige punkter i den længdegående udstrækning af denne, ikke forkortes eller forlænges uønsket;

sprøjtning af stenten med et polymert materiale til coating af tværstiverne;

indsættelse af anordningen i den indre diameter af formen;

påføring af en polymer på den indre overflade af anordningen; og

afhærdning af polymeren til stenten ved påføring af varme til polymeren, således at en polymer coating (100) dækker i det mindste en del af både tværstiverne og områderne mellem tværstiverne, hvor coatingen er en indre coating, der kobles til anordningen fra den indre overflade af gittersystemet udefter, hvor gittersystemet coates yderligere, og overfladerne af de coatede tværstiver, der berører hulrummet hos patienten, er hævet over den indre coating.

FIG. 1

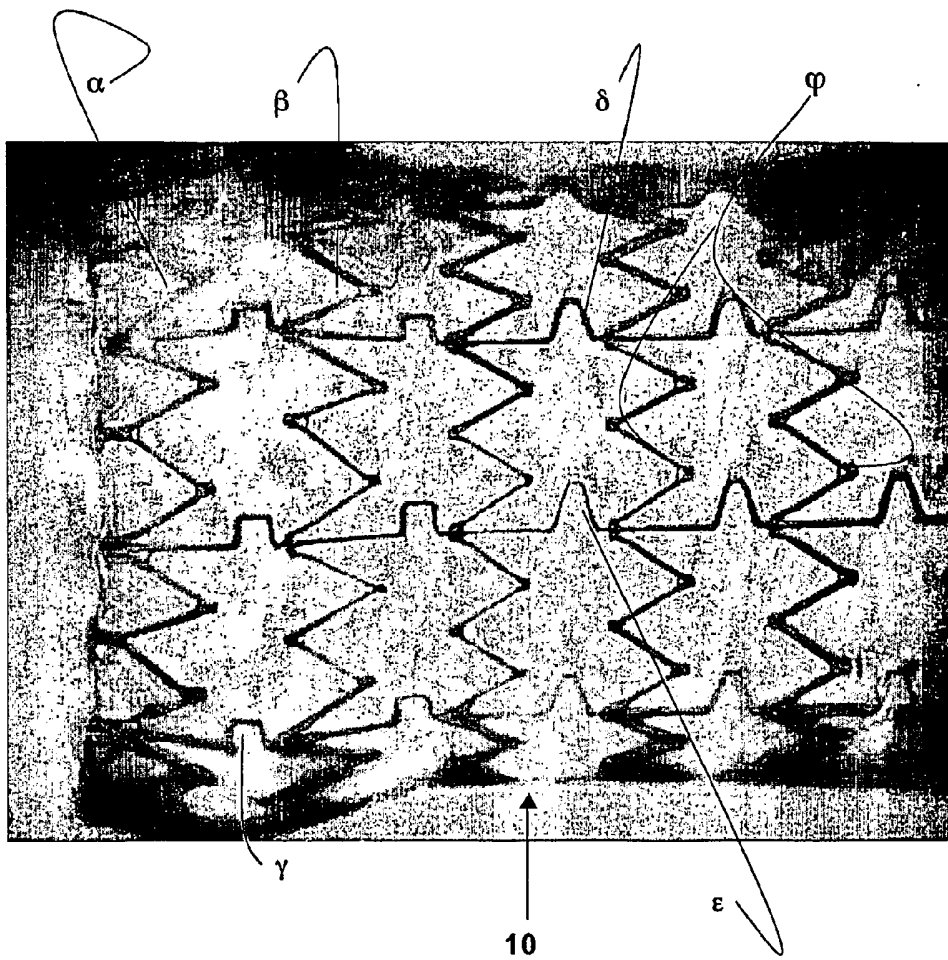


FIG. 2

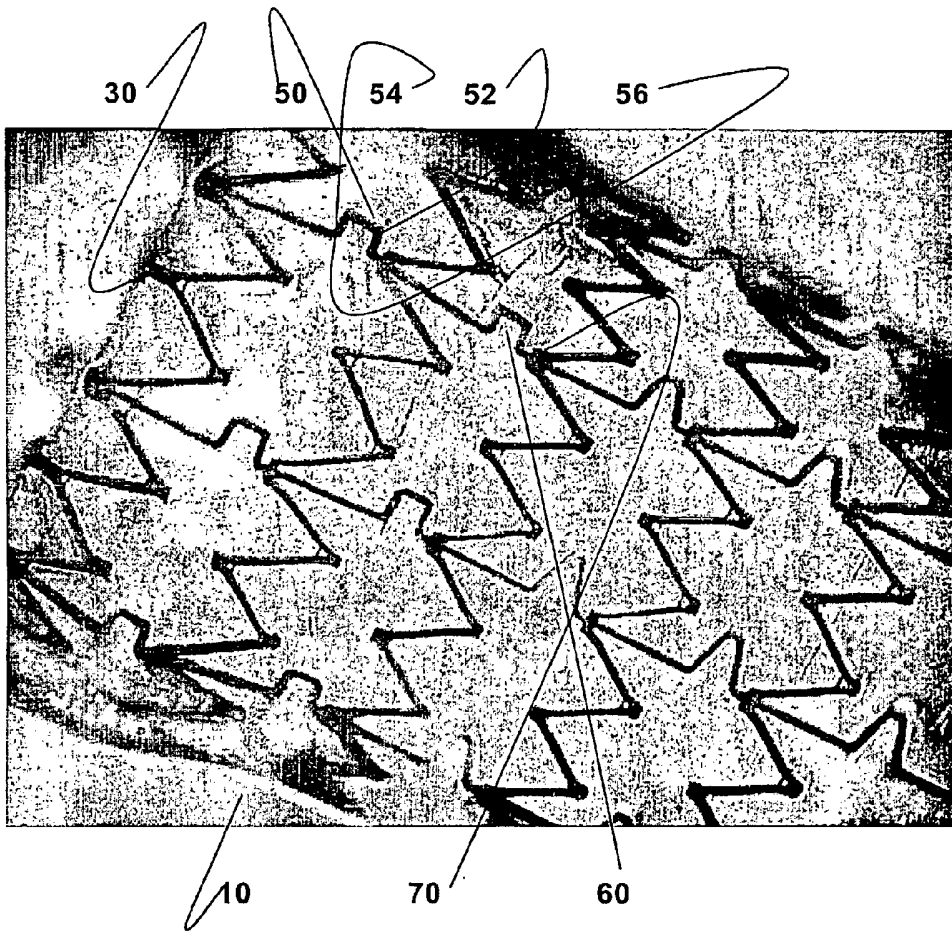


FIG. 3

