An intraocular lens (20) which is compressible to fit through a small incision into the eye and whose refractive power is adjustable once positioned in the eye. The lens (20) includes a deformable soft optic (22), a translucent collar (24) encircling the soft optic (22) and attached to it at fixed spaced points, and haptics (26) attached to the collar (24) for remedielly positioning the soft optic (22) in the eye. The collar (24) has a break or separation (42) to define two collar arms (43, 44). When the collar (24) is compressed the lens (20) can then be inserted in the eye. Once in the eye the collar (24) is released and the arms (43, 44) reformed together to form a rigid circle supporting the soft optic (22). The soft optic (22) is formed by a transparent bag (28) having a thickened perimeter and filled with a transparent fluid. Optic refractive power is adjusted by needle insertion into the bag perimeter to alter the amount of fluid as needed.
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SMALL INCISION INTRAOCULAR LENS WITH ADJUSTABLE POWER

Background of the Invention

The present invention relates to intraocular lenses, and more particularly to intraocular lenses which can be inserted through small incisions into the eye. It further relates to intraocular lenses whose refractive power can be altered while positioned in the eye.

Intraocular lenses have gained wide acceptance recently in the replacement of human crystalline lenses after a variety of cataract removal procedures. One current treatment of cataracts is to surgically remove them through ultrasonic emulsification, so that the light can once again reach the retina. When the natural lens is removed to eliminate the cataract it may be replaced by an artificial lens. The preferred method for restoring vision in an aphakic patient is to surgically implant a lens, a so-called intraocular lens, within the eye. Such a lens, however, need not and generally is not removed. Also, since the intraocular lens is positioned in approximately the same position as the natural lens, it provides vision correction without undue magnification of the image.

A problem associated with the proper implantation of intraocular lenses is the accurate determination of the precise refractive power required for them. Based on measurements of the prescriptive power of the patient's natural lens and measurement of the depth of the eye, a relatively accurate determination can be made of the proper refraction or power of the intraocular lens to be placed in the patient's eye. In most cases, the aphakic patient can
have an intraocular lens implanted which provides good
distance visual acuity even though spectacles may be
required for reading since the intraocular lens cannot
change its refraction or power like a natural lens.
However, in some cases the intraocular lens may not
provide good distance visual acuity. Since an
intraocular lens cannot be readily removed and a new
intraocular lens with a different power surgically
implanted without unduly jeopardizing the patient's
vision, the patient must rely on spectacles to provide
good distance visual acuity.

A disadvantage of conventional rigid
intraocular lenses is that implantation of the lens
requires a relatively large, often six to nine
millimeters, incision in the ocular tissue, and present
methods of cataract removal by phacoemulsification
require only a 3.5 mm incision which may be closed with
a single suture. This long incision surgical procedure
can lead to relatively high complication rates, such as
increased risks of infection, retinal detachment, and
lacerations of the ocular tissues, particularly with
respect to the pupil. Small incision intraocular
lenses do exist though and examples of them include
soft foldable silicone lenses, foldable silicone optics
with conventional non-foldable haptics, hydrogel lenses
which are inserted into the eye in a dry miniature
state and which then absorb ocular fluid to expand to
their full size, and rigid materials with an optic
divided into three sections wherein the outer two
nontransparent sections slide under the central
transparent one to compress the size of the optic.
Each has its disadvantages though including stability
problems. Thus, a need has arisen for an improved
small incision intraocular lens, whose refractive power
is adjustable after implantation.
Objects of the Invention

Accordingly, it is a principal object of the present invention to provide a novel intraocular lens construction.

Another object of the invention is to provide an improved intraocular lens design which can be inserted through a small incision in the ocular tissue.

A further object of the invention is to provide a novel intraocular lens designed to minimize the time required to surgically implant it.

A still further object of the invention is to provide an improved intraocular lens whose refractive power can be adjusted while in place in the eye.

Another object is to provide an improved fluid-filled intraocular lens which does not sag or deform under the effects of gravity.

A further object is to provide an improved method for implanting intraocular lenses through small ocular incisions.

A still further object is to provide a novel method of altering the corrective power of an implanted intraocular lens which can be done quickly and with a minimum of trauma to the eye.

Another object is to provide an improved method of surgically implanting intraocular lenses through incisions generally less than 3.5 mm, which reduces the chances for induced astigmatism.

Other objects and advantages of the present invention will be more apparent to those skilled in the art from the accompanying description taken in conjunction with the accompanying drawings.

Brief Description of the Drawings

Figure 1 is a front perspective view of a small incision intraocular lens of the present invention.
Figure 2 is a perspective view of the lens of Figure 1.

Figure 3 is an end elevational view of the lens of Figure 1.

Figure 4 is a view similar to Figure 1 illustrating the lens in its compressed condition.

Figure 5 is a perspective view of the lens of Figure 1 being inserted in a manipulating instrument of the present invention.

Figure 6 is a front view of an insertion tool of the present invention wherein the lens of Figure 1 is illustrated being inserted therein.

Figure 7 is a fragmentary view of the forward portion of the insertion tool of Figure 6 illustrating the lens of the present invention inserted therein in its compressed condition.

Figure 8 is a view similar to Figure 7 illustrating the slider member engaging the compressed lens and pushing it for ejection from the insertion tool.

Figure 9 is a schematic of the needle system of the present invention provided for adjusting the amount of fluid in the lens of Figure 1.

**Detailed Description of the Preferred Embodiments**

Referring to the drawings, a small incision intraocular lens of the present invention is illustrated generally at 20. Lens 20 very simply comprises a soft optic 22, a generally rigid circular collar 24 attached to and surrounding the soft optic, and a pair of J-shaped haptic loops 26 attached to and extending out from the collar. Many other types of haptics are currently available and may be used in the present lens.

Soft optic 22 is formed from a transparent sac or bag 28 adapted to be filled with a transparent
fluid 30. Optic bag 28 is biocompatible with ocular fluids, transparent and can be formed of any suitable material such as polypropylene, polyethylene or silicone. Bag 28 is impermeable to fluids to prevent exchange of fluids between the bag and the eye. Also, since injected air bubbles in the bag would be troublesome, bag 28 should be constructed of a gas permeable material. Bag 28 is provided with a thickened peripheral portion 32 which serves two functions. First, greater support and stability for attachment of the soft optic to the collar is provided. Second, it forms a thicker edge providing for a self-sealing capability of small punctures, including punctures made by a fine needle, such as shown in Figure 9 at 34, for adjusting the amount of fluid 30 in bag 28. The presence of thicker perimeter 32 of the bag around its entire circumference allows the surgeon to pierce bag 28 at any point on the periphery of the lens to adjust the fluid therein, unlike a valving arrangement which provides limited accessibility. This gives greater flexibility to the adjustment procedure and provides accessibility after surgery. Further, the wound made by needle 34 does not require suturing and is similar to the wound made by a needle used to perform a secondary posterior capsulotomy.

When filled with fluid 30, bag 28 defines an artificial lens suitable for intraocular replacement of the natural lens. Either bag 28 or fluid 30 may have a higher refractive index than the fluids of the eye. If fluid 30 has the higher refractive power, the thickness of optic 22 dictates the power; if bag 28 has the power, fluid 30 separates the two lenses (front and back of bag 28) and specifies the power. In the latter case, fluid 30 may be a balanced salt solution similar to that used to irrigate the eye during surgery.
Silicone oil is a candidate for a refracting fluid. If fluid 30 comprises a gel, the gel should not polymerize or otherwise harden; it is to be a liquid so that some of it can be easily added or removed from the optic to change the refractive power of the lens.

Optic 22 preferably has a biconvex shape. This is the natural lens shape and can produce a very good optical resolution, minimize the effects of internal reflections, and minimize the problems of spherical aberration. For the present adjustable lens 20 this shape is desirable because the adding or removing of fluid 30 has its greatest effect on the central surfaces of optic 22, since the central surfaces of a lens are the locations of the first contact of incoming light rays.

It is also within the scope of the present invention to provide a fluid filled bag similarly constructed as bag 28, which acts on its own as a lens within a nearly intact, natural capsular bag. The bag would be inserted empty and filled during surgery, and post-operative adjustments can then be made. This type of lens can also be used in focusable optical systems such as in microscopes, cameras, and binoculars.

Collar 24 is generally rigid thereby stabilizing fluid filled optic 22 so it does not sag or deform with gravity. It is attached to optic 22, for example, by a plurality of spaced spot welds, preferably four welds 36. Collar 24 is translucent so that it will not transmit light and create a possible double image on the patient's retina. This translucent property also allows the surgeon to easily identify the edge of optic 22 should changes in its liquid volume be required. Collar 24 can be made, for example, of polypropylene or PMMA (polymethyl methacrylate).

The haptics of lens 20 are shown constructed as conventional J-shaped loops 26 and typically are
constructed of either polypropylene or PMMA. Loops 26 stabilize and center optic 22 within the eye with respect to the pupil. Loops 26 are anchored at their proximal ends in the collar 24, as shown in Figure 2. By the orientation of loops 26 it is possible under one construction of the lens, by vaulting optic 22 slightly from the posterior capsule due to the configuration of the haptics angling from the collar towards the posterior capsule, to keep the optic from resting on the posterior capsule of the eye. In other words, referring to Figure 3, both of the haptics would angle downwardly generally below the bottom surface of the optic. This vaulted design would make secondary capsulotomy easier. This lens closely resembles current posterior chamber intraocular lenses, but the haptics are more stable than soft haptics of the prior art lenses. In an alternative construction depicted in Figure 3 the haptics extend generally radially from the collar. The surfaces of the optic protrude above and below the collar and thus the collar does not contact the posterior capsule when the lens is implanted in the posterior chamber. The optic in contact with the capsule may prevent cell migration to the central posterior capsule. If this construction is thereby effective in preventing posterior capsule opacification, secondary capsulotomy by YAG laser or invasive surgery would not be required.

Bag 28 may be filled at the factory for a specified refractive power, or partially filled at the factory and finished in surgery, or inserted empty and completely filled in the operating room. It is further within the scope of the present invention to provide an automated filling system, as illustrated schematically in Figure 9, which allows the surgeon to concentrate on positioning needle 34 in the intraocularly positioned lens through the thickened lens perimeter 32 while a
foot control pump 38 slowly delivers the fluid through the needle into it. Further, a minicomputer 40 can be provided to take the refractive data of lens 20 and determine the amount of fluid to be injected into bag 28. Minicomputer 40 (or other control means) can also control the fluid injection rate and amount so that the surgeon can then concentrate on placing and holding needle 34 in bag 28 without applying excessive pressure, to insure a more accurate system and a reduction of the surgical time.

Collar 24 has a separation or break, as shown in Figure 1 at 42, to define first and second collar arms or sides 43, 44. This allows the surgeon to compress lens 20 by depressing one side 43 of collar 24 at the separation and sliding it under the opposite side 44, as best illustrated in Figure 4. Collar 24 and optic 22 are thereby narrowed to enable them to pass through smaller ocular incisions. Soft optic 22 will thereby be wrinkled temporarily but otherwise not affected. By spot welding bag 28 to collar 24 and not attaching it completely around the circumference of the bag, soft optic 22 can be compressed, and the depressed collar side 43 will slide to the point of the first spot weld 36. After being compressed and inserted in the eye by instrumentation, which is described later, lens 20 is released in the eye and the collar sides 43, 44 repositioned to reform the rigid circular configuration of the collar. In this configuration a locking mechanism, such as is shown in Figures 2 and 4 at 46, locks the arms together in place. This locking mechanism can take the form of a dovetail relationship of the collar sides. In lieu of a locking mechanism, a straight split in the collar is sufficient according to one embodiment of the invention. It further is within the scope of the present invention to provide positioning holes 48 in collar 26 in which tools can be inserted for handling and manipulating lens 20.
The present invention further defines a specially configured instrument, for handling and/or compressing the lens, as best shown in Figure 5, generally at 50. Referring thereto instrument 50 is shown to have a forcep type of construction with first and second arms 52, 54 joined at one end 56 and their opposite ends 58, 60 being provided, respectively, with oppositely disposed box-shaped channels 62, 64 configured for holding collar 24 between them. When the collar is positioned between channels 62, 64, arms 52, 54 can be compressed together to compress the lens into its narrowed shape, as shown in Figure 4, and then inserted as illustrated by the arrow in Figure 6 into an insertion tool 66 and positioned therein as shown in Figure 7 to be positioned entirely within insertion tool 66 or with its tip extending out from it. In lieu of the pin attachment at end 56 the arms 52 and 54 can be welded together as in conventional forceps.

Insertion tool 66 comprises a rectangular shaped tubular member 68 having oppositely facing channels 70, 72 in which collar 24 is inserted. A slider member 74 of the tool slides in the channels and is configured with a low profile, as best shown in Figure 8, to preferably pass entirely underneath haptic loop 26 and directly engage collar 24 for accurate control of the ejection of lens 20 from the tool into the eye. Although depicted in the drawings as being tubular, the preferred configuration of slider member 74 is flat; it is a flat sheet with proximal thumb loop or ring 80 attached to it to fit under haptic 26 and push against collar 24 to thereby move the lens. Insertion tool 66 includes a pair of oppositely placed finger rings 76, 78 on the sides of tubular member 68 and a third finger ring 80 attached to the end of slider member 74 for controlling its sliding movement as best shown by the hand illustrated with phantom
lines in Figure 6. These rings are similar to the rings on certain hypodermic syringes provided to ease manipulations of them.

Tubular member 68 is a rectangular tube approximately 3.0 mm wide and 0.5 mm high with a central portion of one flat surface 82 (the top) missing. The internal rectangle defined by tubular member 68 may have its sharp corners rounded. Lens 20 held in the surgeon's finger, or in the compressing instrument 50, is lubricated with VISCOAT or other visco-elastic substance and inserted into the distal (or proximal) end of the tube. (Alternatively the lens can be inserted in an uncompressed state into a modified insertion tool and the tool compresses it, as well as controllably inserts it, as by pushing it through and out a narrowing channel.) The surgeon slides lens 20 to the desired point, either completely within tubular member 68 or with the forward haptic part of the lens extended. The central portion of the top of the tubular member is omitted so that the surgeon can then use an instrument to position lens 20 in the tubular member from above. The upper opening also allows the surgeon to lubricate the lens from above, to position it within the tube, and to lift the haptic of the lens so that the slider member can pass under it.

By ejecting the compressed intraocular lens 20 slowly and controllably into the eye as is possible with insertion tool 66, the lens resumes its natural shape without "springing open" within the eye and possibly causing injury. Tool 66 can place the lens in the same position as the manual technique employed with current lenses. The lens is manipulated into its remedial position in either the anterior or posterior chambers of the eye. Fluid 30 then is added or withdrawn from bag 28 through needle 34 inserted into
the periphery 32 of the bag. Lens 20 is thus quickly, safely and easily inserted through a small incision into the eye and its refractive power adjusted during the implantation surgery and/or at a later time as needed.

From the foregoing detailed description, it will be evident that there are a number of changes, adaptations, and modifications of the present invention which come within the province of those skilled the art. However, it is intended that all such variations not departing from the spirit of the invention be considered as within the scope thereof as limited solely by the claims appended hereto.
WHAT IS CLAIMED IS:

1. A small incision intraocular lens comprising:
   a deformable soft optic,
   a collar positioned generally about said soft optic and compressible about an axis thereof to fit through a small incision into the eye and then expandable about said axis when in the eye,
   an attaching means for attaching said collar to said soft optic, and
   a haptic means attached to said collar for remedially positioning said soft optic in the eye after having been inserted therein.

2. The intraocular lens of Claim 1 wherein said haptic means comprises at least two spaced haptics extending out from said collar.

3. The intraocular lens of Claim 2 wherein said haptics comprise at least two J-shaped loops.

4. The intraocular lens of Claim 2 wherein said haptics comprise at least two flexible, resilient loops attached at their proximal ends to said collar.

5. The intraocular lens of Claim 1 wherein said collar is rigid, generally circular and positioned about the periphery of said soft optic.

6. The intraocular lens of Claim 1 wherein said collar includes two collar ends which overlap when said collar is compressed and are in generally abutting relation when said collar is in a relaxed state.
7. The intraocular lens of Claim 1 wherein said attaching means attaches the inner surface of said collar directly to the outer surface of said soft optic.

8. The intraocular lens of Claim 1 wherein said attaching means comprises a welding means.

9. The intraocular lens of Claim 1 wherein said attaching means comprises a plurality of attachments spaced about said collar.

10. The intraocular lens of Claim 9 wherein said attachments comprise spot welds.

11. The intraocular lens of Claim 1 wherein said soft optic has a biconvex configuration.

12. The intraocular lens of Claim 1 wherein said soft optic has a thickened perimeter.

13. The intraocular lens of Claim 1 wherein said collar is translucent.

14. The intraocular lens of Claim 1 further comprising a means for changing the power of said soft optic when in the eye.

15. The intraocular lens of Claim 1 further comprising a means for changing the thickness of said soft optic when in the eye.

16. The intraocular lens of Claim 1 further comprising a means for altering the corrective power of said soft optic when said haptic means is positioning said soft optic in the eye.
17. The intraocular lens of Claim 1 wherein said collar is sufficiently compressible so that said intraocular lens will pass through an ocular incision having a length generally not greater than three and one half millimeters.

18. The intraocular lens of Claim 1 wherein said soft optic is constructed of polypropylene, polyethylene or silicone.

19. The intraocular lens of Claim 1 wherein said soft optic comprises a fluid-filled sac.

20. The intraocular lens of Claim 1 wherein said soft optic comprises a bag filled with a pliable substance.

21. The intraocular lens of Claim 20 wherein said pliable substance is inert and optically transparent.

22. The intraocular lens of Claim 20 wherein said pliable substance is a liquid.

23. The intraocular lens of Claim 20 wherein said pliable substance is a liquid gel which does not polymerize or otherwise harden.

24. The intraocular lens of Claim 20 wherein said pliable substance comprises a polymeric liquid material of gel-like consistency.

25. The intraocular lens of Claim 20 wherein said bag is formed of a fluid-impervious, flexible, transparent material.
26. The intraocular lens of Claim 1 further comprising a device including a needle adapted to pass through the eye and into said soft optic when remediably positioned in the eye for varying the amount of fluid in said soft optic to thereby alter its corrective power.

27. The intraocular lens of Claim 26 wherein said soft optic includes a portion through which said needle passes and is adapted to self seal about the puncture opening created by said needle, as said needle is removed from said soft optic.

28. The intraocular lens of Claim 27 wherein said portion is thicker than adjacent portions of said soft optic.

29. The intraocular lens of Claim 27 wherein said portion is about the periphery of said soft optic, and adjacent to said collar.

30. The intraocular lens of Claim 27 wherein said soft optic is filled with a fluid which is transparent, biocompatible with the fluids of the eye, and has a specific gravity generally not greater than aqueous.

31. The intraocular lens of Claim 1 wherein said haptic means is adapted to position said soft optic off of the posterior capsule of the eye when said soft optic is remediably positioned in the posterior chamber of the eye.

32. The intraocular lens of Claim 1 further comprising a compressing means engageable with opposite sides of said collar for compressing said collar so
that said lens can be compressed and inserted through a small incision into the eye.

33. The intraocular lens of Claim 1 further comprising an injecting means for inserting said lens when compressed into the eye and releasing it once in the eye.

34. The intraocular lens of Claim 33 further comprising a lens inserting means for inserting said lens in a compressed condition into said injecting means.

35. The intraocular lens of Claim 34 wherein said lens inserting means comprises an instrument including a pair of opposed channels adapted to engage opposite sides of said collar and a moving means for moving said opposed channels towards each other to compress said collar held therebetween.

36. The intraocular lens of Claim 35 wherein said moving means comprises manually operated forceps.

37. The intraocular lens of Claim 35 wherein said opposed channels are box-shaped.

38. The intraocular lens of Claim 34 further comprising a lubricant lubricating said collar while held by said inserting means and before being inserted into said injecting means.

39. The intraocular lens of Claim 33 wherein said injecting means releases said compressed collar into the anterior chamber of the eye.
40. The intraocular lens of Claim 34 further comprising a lubricant lubricating said collar before said inserting means inserts said collar into the eye.

41. The intraocular lens of Claim 40 wherein said lubricant comprises a visco-elastic substance.

42. The intraocular lens of Claim 33 wherein said injecting means includes a tube configured to hold said collar in its compressed condition and a sliding member positioned in said tube for sliding said compressed collar out of said tube.

43. The intraocular lens of Claim 42 wherein said injecting means includes a pair of finger rings attached to said tube and said intraocular lens further comprises a finger ring attached to said sliding member.

44. The intraocular lens of Claim 42 wherein said sliding member passes underneath said haptic means to directly engage said collar.

45. The intraocular lens of Claim 42 wherein said tube has an upper opening providing access to a lens positioned in said tube.

46. The intraocular lens of Claim 33 wherein said haptic means comprises a pair of opposed haptic loops, and said injecting means positions said loops along the axis of lens ejection.

47. The intraocular lens of Claim 1 wherein said collar is rigid and formed from polypropylene or polymethyl methacrylate, and said haptic means is anchored in said rigid collar.
48. The intraocular lens of Claim 1 wherein said soft optic comprises a bag and fluid at least partially filling said bag.

49. The intraocular lens of Claim 48 wherein said collar stabilizes said soft optic so that said soft optic is not significantly deformed by gravity and so that the edges of said soft optic do not waver in the fluid of the eye.

50. The intraocular lens of Claim 1 wherein said collar is broken so that a first collar side can slide relative to a second collar slide when said collar is compressed.

51. The intraocular lens of Claim 50 wherein said collar includes a locking means for locking said first and second collar sides together when said collar is in its expanded state.

52. The intraocular lens of Claim 51 wherein said locking means includes a dovetail arrangement of said first and second collar sides.

53. A small incision intraocular lens comprising:
   a transparent bag forming an artificial lens when filled with a transparent fluid,
   a compressible stabilizing collar attached to and encircling the periphery of said bag, and
   at least one haptic loop attached to and extending out from said collar for positioning said lens in the eye.

54. The intraocular lens of Claim 53 wherein said bag includes a bag portion generally adjacent the
periphery thereof which is self-sealing after the
removal therefrom of a fluid-adjusting needle.

55. The intraocular lens of Claim 54 wherein
said bag portion is thicker than adjacent portions of
said bag.

56. The intraocular lens of Claim 54 wherein
said bag portion extends about the periphery of said
bag.

57. A method of inserting an intraocular
lens into the eye comprising:
providing an intraocular lens including a
soft optic, a compressible collar attached to said soft
optic, and a haptic attached to said collar,
compressing said collar to thereby compress
said intraocular lens,
thereafter, inserting said compressed
intraocular lens through a small incision into the eye,
thereafter, releasing said compressed
intraocular lens, and
manipulating said released intraocular lens
to its remedial position in the eye.

58. The method of Claim 57 further
comprising adjusting the amount of fluid in said soft
optic to adjust its refractive power.

59. The method of Claim 58 wherein said
adjusting step is after said releasing step.

60. The method of Claim 58 wherein said
adjusting step is after said manipulating step.
61. The method of Claim 58 wherein said adjusting step includes inserting a needle into said soft optic and adjusting the amount of fluid in said soft optic through said needle.

62. The method of Claim 57 wherein said compressing includes placing said intraocular lens into a compressing tool and thereafter actuating said compressing tool.

63. The method of Claim 62 further comprising, before said inserting, transferring said compressed intraocular lens from said compressing tool to an insertion tool, and said inserting including inserting said insertion tool holding said intraocular lens through the small incision into the eye.

64. The method of Claim 62 wherein said actuating includes squeezing said compressing tool.

65. The method of Claim 57 wherein said inserting includes holding said compressed intraocular lens in a tubular instrument and ejecting said compressed intraocular lens from said tubular instrument when at least the tip of said tubular instrument is in the eye.

66. The method of Claim 65 wherein said ejecting includes engaging a sliding member of said tubular instrument against said held intraocular lens and pushing it out of said tubular instrument.

67. The method of Claim 66 wherein said engaging includes engaging said sliding member directly against said collar.
68. The method of Claim 65 further comprising, before said inserting, lubricating said intraocular lens.

69. The method of Claim 57 wherein said releasing includes releasing said intraocular lens into the anterior chamber of the eye.

70. The method of Claim 57 further comprising, filling said soft optic at least partially with fluid before said inserting step.

71. The method of Claim 57 further comprising, filling said soft optic at least partially with fluid after said inserting step.

72. The method of Claim 57 further comprising, filling said soft optic with fluid after said manipulating step.

73. The method of Claim 57 wherein said manipulating includes manipulating said haptic into position against ocular tissue.

74. The method of Claim 73 further comprising, after said manipulating step, altering the amount of fluid in said soft optic to achieve the desired refractive power thereof.

75. The method of Claim 57 further comprising, filling said soft optic with a fluid having a higher refractive index than that of the fluids of the eye.

76. The method of Claim 57 further comprising, before said inserting, forming said small
incision, in the ocular tissue to have a length of
about three and one half millimeters.

77. A method for inserting an intraocular
lens into the eye comprising:
providing an intraocular lens including a
transparent bag defining an artificial lens when filled
with fluid, and a plurality of haptic loops attached to
said bag,
inserting said intraocular lens through a
small incision into the eye,
thereafter positioning said haptic loops to
remedially position said intraocular lens in the eye,
and
thereafter, inserting a needle directly into
the periphery of said transparent bag and changing the
amount of fluid in said bag through said inserted
needle to adjust the refractive power of said
artificial lens.

78. An instrument for compressing an
intraocular lens having a compressible stabilizing
collar surrounding a soft optic comprising:
a pair of arms connected together at one end
and having opposite ends, and
a pair of box-shaped channels attached at
said opposite ends and configured and oriented in
opposed relation to hold the stabilizing collar in and
between them and to compress the collar when said pair
of arms are compressed towards one another.

79. An instrument for inserting a
compressible intraocular lens into the eye
comprising:
a tubular member having a distal member end
and defining a channel in which the compressible
intraocular lens can slide,
said channel having a distal channel opening,
a pushing means for pushing, when said distal
member end is in the eye, a compressed intraocular lens
out of said distal channel opening and into the eye,
and
a handle attached to said tubular member at a
location spaced from said distal member end adapted to
be grasped for manipulating said tubular member.

80. The instrument of Claim 79 wherein said
channel has a rectangular cross section.

81. The instrument of Claim 79 wherein said
tubular member defines in cross-section a rectangle
approximately 3 mm wide and 0.5 mm high.

82. The instrument of Claim 79 wherein said
pushing means comprises an elongated sliding member
adapted to slide in said channel.

83. The instrument of Claim 82 further
comprising a finger ring attached to the outer end of
said sliding member.

84. The instrument of Claim 79 wherein said
handle comprises a pair of finger rings attached to the
sides of said tubular member.

85. The instrument of Claim 79 further
comprising lubricant on the lens contact surfaces of
said channel.

86. The instrument of Claim 79 wherein said
channel is configured to hold the intraocular lens in
its compressed condition.
87. The instrument of Claim 79 wherein said pushing means passes underneath the haptic of the intraocular lens positioned in said channel when pushing the intraocular lens.

88. The instrument of Claim 87 wherein said pushing means pushes directly against the collar of the intraocular lens.

89. The instrument of Claim 79 wherein said tubular member has a top side and an upper opening passing through said top side, communicating with said channel and through which an intraocular lens can be manipulated when in said channel.
### I. CLASSIFICATION OF SUBJECT MATTER

According to International Patent Classification (IPC) or to both National Classification and IPC:

IPC(4): A61F 2/16, A61B 17/00
U.S. Cl.: 623/6, 128/303R

### II. FIELDS SEARCHED

Minimum Documentation Searched * 

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Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *

### III. DOCUMENTS CONSIDERED TO BE RELEVANT

<table>
<thead>
<tr>
<th>Category</th>
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<td>X</td>
<td>US, A, 4,615,702 (KOZIOL ET AL) 07 OCTOBER 1986, See abstract, column 2, lines 45-68 and column 3, lines 1-37</td>
<td>1-5, 11, 17, 18, 31, 12, 14-16, 19-30, 32-49, 57-76</td>
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<td>Y</td>
<td>US, A, 4,608,049 (KELMAN) 26 AUGUST 1986 See Figs. 7 and 8, column 4, lines 52-57, column 6, lines 61-68 and column 7, lines 1 and 19-68</td>
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<td>US, A, 4,596,578 (KELMAN) 24 JUNE 1986 See Figs. 1-4 and 7, column 2, lines 1-25, and column 4, lines 4-12</td>
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<td>GB, A, 2,124,500A (MAZZOCCO) 22 FEBRUARY 1984 See page 4, lines 5-43</td>
<td>6-10, 47, 50, 57-76</td>
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<td>US, A, 4,073,014 (POLER) 14 FEBRUARY 1978 See Figs. 1-10, column 3, lines 20-29</td>
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<td>US, A, 4,573,998 (MAZZOCCO) 04 MARCH 1986 See Figs. 50-53, Fig. 35, Figs. 4 and 7</td>
<td>78, 12, 32, 79-89</td>
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* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
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**"A" document member of the same patent family**

### IV. CERTIFICATION

Date of the Actual Completion of the International Search:

11 FEBRUARY 1988

Date of Mailing of this International Search Report:

10 MAR 1988

International Searching Authority:

ISA/US

Signature of Authorized Official:

[Signature]

Ronald L. Frinks
<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of Document, with indication, where appropriate, of the relevant passages</th>
<th>Relevant to Claim No.</th>
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<td><strong>Y</strong></td>
<td>EP, A, 0,190,056 (DAVENPORT) 06 AUGUST 1986</td>
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<td>US, A, 4,585,457 (KALB) 29 APRIL 1986</td>
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<td>US, A, 3,919,724 (SANDERS ET AL) 18 NOVEMBER 1975, See abstract and column 2,</td>
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<td>US, A, 4,619,662 (JUERGENS, JR.) 28 OCTOBER 1986, See column 1, lines 47-54</td>
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<td>IOL Implantation&quot; by Evan D. Jones, M.D., pages 28 and 29 See entire document</td>
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<td>See column 3, lines 31-42</td>
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<td>US, A, 4,619,256 (HORN) 28 OCTOBER 1986</td>
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<td>US, A, 2,887,110 (ROESCHMANN) 19 MAY 1959</td>
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<td>See Figs. 1-5</td>
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<td><strong>Y</strong></td>
<td>EP, A, 0,151,020 (SHUTE) 07 AUGUST 1985</td>
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<td>See elements 18 and 22 in Fig. 9</td>
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