Recent Trends in Lamellar Keratoplasty

Theses

Submitted for Partial Fulfillment

Master Degree of Ophthalmology

BY

Eman Taher Alhwat M. B., B.Ch,

Under the supervision of

Prof. Dr. / Golzamin Ragheb EL-Hawary

Professor of Ophthalmology Faculty of Medicine, Ain shams University

Assist. Prof. Dr. /Mona Mohamed EL-Fiky

Assistant Professor of Ophthalmology Faculty of Medicine, Ain shams University

Faculty of Medicine

Ain Shams University

Cairo

2011

الطرق المستحدثه في الترقيع الطبقي للقرنيه

رساله

توطئه للحصول على درجه الماجستير في طب وجراحه العيون

مقدمه من

الطبيبه / إيمان طاهر محمود الحواط بكالوريوس الطب والجراحه

تحت اشراف

أ.د. جلزمين راغب الهوارى

أستاذ طب وجراحه العيون جامعه عين شمس

أ.د. منى محمد الفقى

أستاذ مساعد طب وجراحه العيون جامعه عين شمس

جامعه عين شمس

القاهر ه

7.11

Content

List of abbreviations

List of figures

List of tables

Introduction	1
Anatomy and Biochemisty	3
Indications of lamellar keratoplasty	18
Techniques of lamellar keratoplasty	34
Complications of lamellar keratoplasty	82
Summary	93
References	96

الملخص العربي

List of Tables

Table No.	Description
1	Hysteresis of normal, keratoconic and Fuch's subjects.
2	Indications of lamellar keratoplasty.
3	Techniques of lamellar keratoplasty.
4	Complications of lamellar keratoplasty.

List Of Figures

Figure No.	Description
1	A-Cornea cross-section, corneal layers, epithelium, bowman's, stroma, descemet's, endothelium. B- Layers of the cornea.
2	Reichert's Ocular Response Analyzer TM
3	Corneal hysteresis is the difference between the inward and outward applanation pressures
4	(1)Slit-lamp picture of an eye with advanced keratoconus showing gross inferior bowing (ectasia) of the cornea with associated thinning (2)vogt's strieas in keratocouns.
5	Corneal topography for Pellucid Marginal Degeneration.
6	Topography of post~LASIK keratoectasia
7	Juvenile Hereditary Epithelial dystrophy
8	Bowman's layer dystrophy (Reis-Bu"cklers corneal dystrophy
9	central corneal scar
10	posterior corneal vesicles and opacities in linear bands and other polymorphous configurations typical of posterior polymorphous corneal dystrophy
11	Corneal Guttata
12	Fuch's dystrophy
13	Congenital Hereditary endothelial dystrophy
14	Pseudophakic bullous keratopathy
15	Cogan-Reese syndrome
16	Chandler's syndrome
17	essential iris atrophy
18	Contact lens-induced endotheliopathy
19	Donor preparation of ALK
20	Anwar's big-bubble technique
21	DALK using both air and viscoelastic material
22	Microkeratome-assisted deep lamellar keratoplasty technique. Schematic diagram showing recipient bed preparation
23	Microkeratome-assisted deep lamellar keratoplasty technique.
24	Schematic diagram illustrating a femto-second LASER assist ALKP
25	Femtosecond laser-assisted anterior LKP
26	Drawings of the deep lamellar endothelial keratoplasty procedure
27	Slit-lamp photographs of the same eye one day and two weeks after posterior lamellar keratoplasty

1	
28	Drawings of the surgical procedure of DLEK through a 5.0 mm scleral tunnel incision
29	Surgical steps for small-incision deep lamellar endothelial keratoplasty (DLEK).
30	Barron Vacuum Punch is comprised of three pieces: base with tissue pedestal, tissue retainer and locking ring.
31	Surgical steps for small-incision deep lamellar endothelial keratoplasty (DLEK).
32	Slit-lamp photographs of the same eye one week and one month after DLEK
33	Removal of endothelium and Descemet's membrane from the recipient eye with the anterior chamber filled with air.
34	Inserting a folded donor graft into the eye with forceps
35	Inserting a donor graft into the eye through a Busin funnel glide.
36	Slit-lamp image showing the residual air bubble in the anterior chamber 2 days after Descemet stripping automated endothelial keratoplasty
37	Preparing the recipient cornea in DMEK.
38	John DMEK/DSAEK Dexatome spatula (ASICO)
39	Transplanting the donor cornea.
40	Trypan blue-stained donor Descemet's membrane in John DMEK block (ASICO)
41	Attachment of the donor Descemet's membrane to the patient's cornea using a big-bubble technique
42	Schematic diagram illustrating a femto-second LASER assist DLEK.
43	cross section view of lamellar &trephination cut (solid red line) in laserassisted DLEK
44	Pseudo-anterior hamber formation due to micro-perforation of the Descemets membrane.
45	Iris stromal atrophy following managed post-operative pupillary block.
46	Interface wrinkling following deep anterior lamellar keratoplasty.
47	Vascularization of the interface with protein extravasations and opacification
48	Suture-induced reaction, cheese-wiring and loose sutures after deep anterior lamellar keratoplasty
49	Surface scar from herpes simplex infection following deep lamellar endothelial keratoplasty (DLEK) surgery.
50	Folds in the lamellar graft are visible on retroillumination.
51	Folds are present in the center of the graft (A and B).

52	Toluidine blue–stained specimen of the excised donor graft demonstrating a fixed fold
----	---

List of Abbreviation.

Abb.	Description
ALK	Anterior Lamellar Keratoplasty.
ATP	Adenosine Ttriphosphate.
BSCVA	Best Spectacle Corrected Visual Acuity.
BSS	Balanced Salt Solution.
СН	Corneal Hysteresis.
CS	Chondroitin Sulphate.
DALK	Deep Anterior Lamellar Keratoplasty.
DLEK	Deep Lamellar Endothelial Keratoplasty.
DM	Descemet's Membrane.
DMEK	Descemet's Membrane Endothelial Keratoplasty.
DNA	Deoxy Nucleic Acid.
DS	Dermatan Sulphate.
DSAEK	Descemet's Stripping Automated Endothelial Keratoplasty.
DS~PGS	Dermatan Sulphate-Proteoglicans.
FED	Fuch's Endothelial Dystrophy.
FS Laser	Femto Seconed Laser.
GAGS	GlycosAminGlycans.
ICE	Irido-Corneal Endothelial Syndrome.
IOP	Intra Ocular Pressure.
KS	Keratain Sulphate.
KS~PGS	Keratain Sulphate-Proteoglycans.
LASIK	Laser in Situ keratomalesis.
LK	Lamellar Keratoplasty.
NADPH	Nicotinamide adenine Dinucleotide Phosphate.
OVD	Ophthalmic Viscoelastic Device.
PAS	Periodic Acid Shift.
PBK	Pseudophakic Bullous Keratopathy.
Pk	Penetrating Keratoplasty.
PMD	Pellucid Marginal Degeneration.
PPLK	Progressive Post Laser Keratoectasia.
PPMD	Posterior Polymorphous Destrophy.
PPV	Pars Plana Vitrectomy.
UV	Ultra Violet.

Introduction

The history of lamellar keratoplasty (LK) spans over 100 years, and the advantages of lamellar surgery have long been known. The surgery was usually used for tectonic purposes; however, new techniques have expanded the applications of LK in optical rehabilitation. Instrumentation such as viscoelastics, diamond knives, ultrasonic pachymetry, artificial anterior chambers, advanced microkeratomes, and the excimer laser have enhanced our ability to work more safely in the tedious microsurgical environment of the lamellar procedure (Terry; 2000).

Microsurgical techniques have vastly improved the technique of lamellar keratoplasty, as they have also substantially improved the results with penetrating keratoplasty.

Lamellar keratoplasty involves replacement of the damaged or diseased part of the cornea or even the stroma (middle layer of the cornea), Bowman's membrane (second layer of the cornea) or the endothelium with donor material. Most of the other layers of the cornea can be preserved. The donor corneal disc becomes repopulated with host cells, and the recipient epithelium usually covers the anterior corneal surface. This procedure is technically more difficult than penetrating keratoplasty. Anterior Lamellar keratoplasty has the advantage of being primarily extraocular, making it a procedure that preserves the endothelium. The risk of endothelial rejection is abolished or eliminated. The risks of wound leaks or flat anterior chambers associated with an intraocular procedure may be eliminated (Synder;2005).

Lamellar keratoplasty is an exciting new form of surgery that provides selective lamellar transplantation. This surgery allows restoration of the normal corneal topography, with very little change from the preoperative corneal curvature. This in turn provides more accurate IOL calculations for combined cataract and transplantation procedures, as well as avoiding the surprises of high astigmatism and high refractive errors that are so common after standard PK surgery (Busin et al;2005). DLEK can even be performed suturless through a lamellar wound as small as a modern cataract surgery wound (Melles;2002).

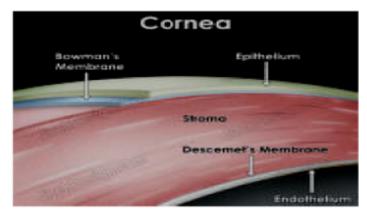
Whereas the surgical techniques of LK are still in accelerated development, it appears that LK is moving forward quickly to become the preferred, if not the ideal method of corneal transplantation for the treatment of endothelial dysfunction.

The disadvantages of LK are the technical challenge of the procedure, the risk of perforation, and the need for special designed blades for dissection. Another source of frustration for surgeons is the haze that may form in the interface between the host and donor corneas (*Leccisotti;2007*).

Anatomy

The cornea is the most densely innervated tissue in the body. The sensory supply is via the first division of the trigeminal nerve. There is a subepithelial and a stromal plexus of nerves (Kanski, 2003).

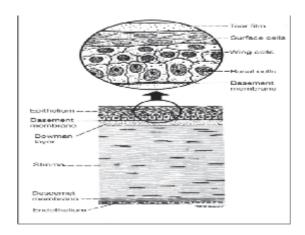
The average corneal diameter is 11.5 millimeter (mm) (vertical) and 12mm (horizontal). It is made up of 5 distinct layers, fig. (1) (Snell and lemb; 1998). Starting from the outer layer and moving inward, they are:



(Fig.1 A): Cornea cross-section, corneal layers, epithelium, Bowman's, stroma, Descemet's, endothelium (Gipson and Nancy; 1995).

1-The epithelium

The epithelium is stratified squamous and non-keratinized, and consists of a single layer of basal columnar cells attached by hemidesmosomes to the underlying basement membrane, 2 to 3 rows of wing cells and two layers of squamous epithelial cells (Snell and lemb, 1998). The surface area of the outermost cells is increased by microplicae and microvilli that facilitate the attachment of mucin and the tear film. After a lifespan of a few days the superficial cells are shed into the tear film (Kanski, 2003).



(Fig. 1 B) layers of the cornea (Kanski, 2007).

The epithelial stem cells are principally located at the superior and inferior limbus, possibly in the palisades of Vogt, and are indispensable for the maintenance of healthy corneal epithelium. They also act as a junctional barrier, preventing conjunctival tissue from growing onto the cornea (kanski, 2007).

2-The Bowman's layer

The Bowman's layer is the acellular superficial layer of the stroma (Roberts, 2000).

3-The substantia Propria

The substantia propria makes up 90% of corneal thickness. It is principally composed of regularly orientated layers of collagen fibrils whose spacing is maintained by proteoglycan ground substance {chondroitin sulphate (CS) and keratin(KS)} with interspersed modified fibroblasts (keratocytes) (Kanski, 2007).

4-The Descemet's membrane

The Descmet's membrane; is composed of a fine latticework of collagen fibrils. It is periodic acid schiff (PAS) positive glassy basement membrane of the endothelial cells and does regenerate after injury. It forms at the periphery the line of Schwalbe, referred to clinically when assessing the anterior chamber by gonioscopy. It consists of an anterior banded zone that is deposited in-utero and a posterior non-banded zone laid down throughout life by the endothelium (Kanski, 2003).

5-The endothelium

The endothelium consists of a single layer of hexagonal cells that cannot regenerate. It plays a vital role in maintaining corneal deturgescence (*Kanski*, 2007).

The corneal structural and reparative properties are essential to its function as a resilient, yet transparent, barrier to intraocular injury. Because the cornea is also the scaffold for the major refractive surface of the eye, any mechanical or biological response to injury will also influence optical performance. Consequently, the same mechanisms responsible for preserving ocular integrity can undermine the goals of achieving predictable and stable visual outcomes after keratorefractive surgery (Roberts, 2000).

Biochemistry

The cornea is not only designed for light transmission but is the main light refracting element in the eye. Its cellular and extracellular components are of the same chemical basic composition as other cells and tissues in the body that normally scatter light extensively, thus rendering them opaque. Light scattering in opaque tissues is due to the large disparity in refractive index (RI) between matrix components such as collagen, glycosaminglycans (GAGs) and cells. The cornea's ability to transmit light is a function of how the cells and matrix components are organized within the tissue to reduce this refractive index disparity (Forrester et al, 2003).

Corneal transparency

1-The epithelium

All the thick stratified squamous epithelium presents the first refracting interface to transmitted light. Most of the light absorbing properties take place in this layer, mainly for short wave-length light. However the majority of light of visible spectrum is transmitted through the epithelium (*David*, 2004).

The cells are typical keratin expressing epithelial cells, containing integrin receptors for basement membrane components such as fibronectin & laminin. Corneal epithelial cells express a particular combination of keratin 3 and keratin 12. Keratin 12 may be important for corneal epithelial junctions (Berman, 1991). Hemidesmosomes affect the adhesion between the basal epithelium and the basement membrane. They bound to the corneal stroma through a band of anchoring fibrils, which pass through the lamina

densa in a woven network. These fibers are composed of type VII collagen. In addition type XVI collagen supports firm adhesions in theses basal cells (*David*, 2004).

The epithelium presents an effective barrier against fluid transport, which is achieved by extensive close contacts and junctional complexes between the basal cells. Spot desmosomes are numerous and studies have revealed that there are differences in the content of desmosomal proteins depending on the site (Forrester et al, 2003).

2~The stroma

A-collagen

Several different types of collagen are present in the cornea. In addition to the normal basement membrane, type IV & VII are in contact with the epithelial and endothelial layers, the two specialized corneal regions. Bowman's layer is a condensation of type I and IV with high proportion of type III collagen in a matrix containing proportions of chondoitin sulphate (CS) & dermatan sulphate (DS), while Descmet's membrane contains high levels of novel collagens (types IV, VIII, IX & XII) organized in lattice arrangement. This provides elasticity and deformability to the cornea while maintaining high levels of light transmission (Kenney et al, 2005).

The regular arrangement of the stromal type I collagen (which accounts 50 - 55 % of stromal collagen) fibrils is considered to be an important factor in corneal transparency. Transparency was initially attributed to destructive interference in which light is scattered by neighboring fibrils in predictable and opposing directions, which tends to cancel each other out except along the primary visual axis. However, this concept cannot be applied to