Keraring Implantation Using Keraring Nomogram Versus the Q Value Modified Nomogram of a Single Keraring Segment Implantation for Treatment of Keratoconus: a Comparative Study

Protocol study

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INTRODUCTION

Keratoconus (KC) is a disorder of the eye which results in progressive thinning of

the cornea. 1-3 This may result in blurry vision, double vision, near sightedness, astigmatism,

and light sensitivity. Usually both eyes are affected. In more severe cases a scarring or

a circle may be seen within the cornea.⁴

While the cause is unknown, it is believed to occur due to a combination of genetic,

environmental, and hormonal factors.³ About seven percent of those affected have a family

history of the condition. ³ Proposed environmental factors include rubbing the eyes

and allergies. The underlying mechanism involves changes of the cornea to a cone

shape.³ Diagnosis is by examination with a slit lamp.³

Initially the condition can typically be corrected with glasses or soft contact lenses.³ As the

disease worsens special contact lenses may be required.³ In most people the disease stabilizes

after a few years without severe vision problems.³ In a small number of people scarring of the

cornea occurs and a corneal transplantation is required.³

Keratoconus affects about 1 in 2,000 people. 3,4 It occurs most commonly in late childhood

to early adulthood. 3 While it occurs in all populations it may be more frequent in certain ethnic

groups such as those of Asian descent. The word is from the Greek kéras meaning cornea and

the Latin conus meaning cone.⁵

Improvement in visual acuity and refraction after ICRS implantation is accomplished by a

shortening of collagen lamellae along the arc length of the ring. There is a redistribution of

corneal stress due to the change in the shape of the cornea after implantation of ICRS. 11,12

Keraring is an orthosis implanted in the corneal stroma. It acts upon corneal tissue by altering

its central curvature and shape, thus reducing or eliminating morphological irregularities and

existing myopia and astigmatism. They are available as 160° segments made of PMMA. They

are triangular in cross-section with a 600 micron base and an apical diameter of 5 mm. They

come in variable thickness (0.15, 0.20, 0.25, 0.30 and 0.35 mm) in 0.05 mm steps.⁵

The change in the corneal structure induced by any ICRS can be roughly estimated by the

Barraquer thickness law; therefore, the outcome achieved is directly proportional to the thickness

of the ICRS and inversely proportional to its diameter. ^{1, 7, 11}

PURPOSE

To compare the outcomes of implantation of the kerarings according to kera nomogram

versus the implantation of a single keraring segment (210 arc) according to the modified Q value

nomogram.

PATIENTS AND METHODS

Design: A prospective comparative clinical study.

The author will obtain the approval of the ethical committee in faculty of medeicine in

Sohag University Hospital and informed consent will obtained from all patients.

This study will be conducted on 150 eyes with type I, II and III keratoconus. The patients will be divided into 3 groups. Group A will include 50 eyes to be treated with keraring implantation according to keraring nomogram. While group B will include 50 eyes to be treated with single (210 arc) keraring implantation according to the modified Q value nomogram. The patient will followed up for 6 months after surgery. Group C will include KC 50 eyes as control group with matching the same age group and subjected to surgery.

All patients were subjected to complete ophthalmologic examinations that included measurement of the uncorrected visual acuity (UCVA), best spectacle-corrected visual acuity (BSCVA), manifest refraction, slit lamp examination of anterior segment, intraocular pressure measurement and a detailed fundus examination.

patients were subjected to preoperative and postoperative UCVA, BCVA, refraction, Pentacam pachymetry and keratometry examinations at 1, 3 and 6 months follow up period.

The ring segments were chosen based on a nomogram from the manufacturer.

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