## **ORIGINAL RESEARCH**

# Clinical profile of keratoconus in central India

<sup>1</sup>Dr. Rishi Gupta, <sup>2</sup>Dr. Ritika Agrawal, <sup>3</sup>Dr. Mita Joshi, <sup>4</sup>Dr. Teena Agrawal, <sup>5</sup>Dr. Abha Verma

<sup>1</sup>Assistant Professor, <sup>2</sup>Senior Resident, <sup>3,4</sup>Associate Professor, <sup>5</sup>Professor, School of Excellence for Eye, MGM Medical College, Indore, MP, India

### **Corresponding Author**

Dr. Rishi Gupta

Assistant Professor, School of Excellence for Eye, MGM Medical College, Indore, MP, India Email: grishi09@gmail.com

Revised Date: 18 March, 2024 Acceptance Date: 10 April, 2024

#### ABSTRACT

**Aim:** The aim this study to describe the demography and clinical profile of keratoconus in central India. **Methods:** This is single centre, cross-sectional hospital-based study conducted on68 patients presenting between April 2023 to September 2023. All Patients of Keratoconus without any previous interventions were included in this study. **Results:** 129 eyes of 68 patients diagnosed with Keratoconusare used for the analysis. The mean age of patients was 19.01 +/- 6.64years.The majority of patients were male (58.82%) Out of 68patients 61 had bilateral keratoconus (89.7%) and 7 patients had unilateral presentation (10.29%). Common signs noted were prominent corneal nerves (58.13%), Fleischer ring (50.38%), corneal thinning(45.73%), Vogt's striae (24.80%), and corneal hydrops (2.32%). Allergic eye disease like Vernal kerato conjunctivitis(VKC) was most common association noted. **Conclusion:** Study concludes that bilateral nature of disease was more preponderant. Gender dominance was seen in males with commonest presentation in second and third decades. Common Mode treatment included corneal collagen cross linking, Scleral contact lenses and keratoplasty. **Key word:** Keratoconus, Corneal Ectasia, Collagen Cross Linking

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution- Non Commercial-Share Alike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

#### **INTRODUCTION**

Keratoconus isa non-inflammatory, progressive ectasia affecting the cornea.[1] It is commonly a bilateral but asymmetric[2,3] disease. It is characterized by the protrusion of the cornea caused by localized and central corneal thinning.[4] Keratoconus causes irregular astigmatism, leading to various degree of visual impairment.[5] Existing literature indicates that keratoconus tends to occur in adolescence and most likely the course of progression extends to the age of 30 or 40 years.[3] Many risk factors of keratoconus have been described previously, including rubbing one's eyes, family history of keratoconus, atopy, allergy, asthma, and eczema.[6] Clinically, the disease ranges from subclinical "formefruste" keratoconus (ffKC) to the more severe progressive form, resulting in corneal scarring, hydrops, and blindness. The prevalence of keratoconus is highly variable, ranging from 0.2/100,000 in Russia[7] to 3,300/100,000 in Iran.[8]The aim of this study is to describe the prevalence and clinical profile of keratoconus in central India.

#### **METHOD**

This is single centre, cross-sectional hospital-based study conducted on 68 patients presented between April 2023 to September 2023. All the patients of Keratoconus without any previous interventions were included in this study. Any patient with a history of ophthalmic surgery was excluded. Informed consent from was obtained the patient or the parents/guardians of the patient. Clinical records from the Cornea Clinic of School of Excellence for Eye, MGM Medical College Indore were reviewed. The parameters noted were patient's demographics, associated conditions, presenting visual acuity (uncorrected or best corrected with spectacle), Slit lamp biomicroscopic findings( Clinical Finding ), keratometric reading on auto keratometer and topography. Based on presenting visual acuity (logMar) patients were classified as no visual impairment (0.00-0.18), mild visual impairment (0.30-0.48), moderate visual impairment (0.60-1.00)and severe visual impairment (< 1.00). Keratoconus was categorized according to the Amsler classification with corneal Krumeich (AK) topography data[9]

commonly with male to female ratio1.42. Out of 68

patients 61 had bilateral keratoconus (89.7%) and 7

patients had unilateral presentation (10.29%). (Table

#### RESULT

Diagnosis was made clinically and confirmed by topography. 129 eyes of 68 patients were diagnosed with keratoconus. The mean age of keratoconus is 19.01 + -6.64 (9-45) years. Male were affected more

## Table 1: Demography

SN	Parameters	No	%
1	Total patients	68	
2	Bilateral cases	61	89.70
3	Unilateral cases	7	10.29
4	Mean age +/- SD(range )	19.01+/- 6.64(9-45)	
5	Male female Ratio	1.42	

1)

Common clinical findings included prominent nerves, Fleischer ring, central thinning, Vogt's striae, corneal scarring and hydrops.(Table 2)

#### Table 2: Clinical findings

SN	Findings	No of eyes	%
1	Prominent corneal nerves	75	58.13
2	Fleischer ring	65	50.38
3	Central thinning	59	45.73
4	Vogtsstriae	32	24.80
5	Corneal scarring	14	10.85
6	Hydrops	3	2.32

Among the 68 patients, history of eye rubbing and allergic eye disease like vernal keratoconjunctivitis(VKC) were the most common association noted in this study. Family history of keratoconus were noted in 3 (4.41%) patients. (Table-3)

#### **Table 3: Associated Conditions**

5	SN	Associated conditions	No of patients	%
	4	Eye rubbing history	12	17.6
	1	Allergic eye disease like Vernal keratoconjuctivitis	11	16.17
	2	Allergic Rhinitis/Asthma	4	5.88
	3	Family history	3	4.41
	5	Marfan's syndrome	1	1.47

At the time of diagnosis most of the eyes; 58 (44.96%) had moderate visual impairment (0.60-1.00). Nil visual impairment (0.00-0.18), mild visual impairment (0.30-0.48) and severe visual impairment (Worse than 1.00) were seen in 33 (25.58%), 26 (20.15%), and 12 (18.38%) eyes, respectively.(Table-4)

#### Table 4: Presenting Visual Acuity

SN	Grade of visual impairment	Visual acuity (Log Mar)	No of eyes	%
1	No impairment	0.00-0.18	33	25.58
2	Mild	0.30-0.48	26	20.15
3	Moderate	0.60-1.00	58	44.96
4	Severe	< 1.00	12	9.30
	Total eyes		129	

Corneal topography was done in all the patients. Severity of all the eyes were graded according to the Amsler-Krumeich(AK) criteria. Stage 2 keratoconus was seen in nearly 66 (51.16%) eyes followed by stage 3(24.03%), stage 1(17.82%) and stage 4(6.6%).(Table-5)

#### Table 5: Staging of Keratoconus-According to the Amsler-Krumeich(AK) criteria

SN	Stage	No. of eyes	%
1	Stage 1	23	17.82
2	Stage 2	66	51.16
3	Stage 3	31	24.03
4	Stage 4	9	6.60

In our study corneal collagen cross linking(37.20%) was the most commonly performed procedure in the

management of keratoconus. Scleral lens with or without prior corneal collagen cross linking was used

in 30 eyes (23.25%). Keratoplasty (Full thickness/DALK) was done in 11 eyes (8.52%).

#### DISCUSSION

In our study mean age of presentation was 19.01 +/-6.64 (9-45) years, which is more and less similar to mean age reported in others studies from Asia.[12,14] A study in United Kingdom has shown that Asians develop Keratoconus at a significantly younger age compared to the Caucasian population.[13] A study from Saudi Arabia reported the mean age (17.7 y) of presentation is nearly similar to our study.[15]. The correlation between Keratoconusand gender is unclear. In our study male were affected more commonly than female with male to female ratio 1.42. Many studies have shown Keratoconus to be more prevalent in males.[5,16,17] Others have found females to be more commonly affected.[10,11] Ziaei et al. [18] found a male-to-female ratio of 1.02 among the Iranian population. A multicentric study from India showed significantly higher prevalence in male patients.[21]In our study most of the cases were bilateral (89.70%) and similar results werenoted in other studies also (19, 20). We have noted that patients presented with severe visual impairment had unilateral presentation. In our study most common clinical findings noted were prominent corneal nerves (58.13%) followed by Fleischer's ring (50.38%), CLEK study [22] showed fleischer's ring as the most common finding followed by prominent corneal nerves. Other finding noted in our study were central thinning (45.73%), Vogt's striae (24.80%), and hydrops (2.32%).Eye rubbing (17.6%) and allergic disease like vernal keratoconjunctivitis eve (VKC)(16.17%) were the most common association noted in this study. Allergic rhinitis, asthma and marfan's syndrome were the other associations observed in the study. Family history of keratoconus were noted in 4.41% patients. Similar associations were noted in other studies[6, 23-25] In present study most patients (65.11%) presented with mild to moderate visual impairment in the affected eye. This is contrary to another study of India[21], where most of the patients(61.42%) had nil to mild visual impairment. Most of the patients in this study were diagnosed in stage 2-3, might be a reason for mild to moderate visual impairment. According to Amsler-Krumeich staging Stage 2 keratoconus was seen in nearly 66 (51.16%) eyes followed by stage 3(24.03%), stage 1(17.82%) and stage 4(6.6%). Previous studies[26,27] have reported a high percentage of patients presenting with severe Keratoconus based on the CLEK classification. Mahadevan et al (2009) revealed that most Keratoconus patients presenting to a tertiary eye care hospital had advanced KC with corneal curvatures of greater than 52D (27). We have noted that spectacle intolerance was the commonest reason for seeking medical intervention in the in the later stages of Keratoconus. A study from Iran reported that over

50% of patients presented with moderate Keratoconus<sup>[17]</sup> Most patients in the Collaborative Longitudinal Evaluation of Keratoconus (CLEK) presented with moderate to advanced study stages.[26]Another study from India showed half of the patients presented with stage 1 disease and over a quarter presented in stage 2.[21] In our study corneal collagen cross linking (37.20%) was the most commonly performed procedure to manage keratoconus which is more than other studies.[17] reasons beingadvancement The probable in technology that offers treatment for wide range of patients and more number of stage 2 - 3 patients with progressive keratoconus. Scleral lens with or with prior corneal collagen cross linking was used in 30 eyes (23.25%). A reasonable contact lens service, noninvasive and better visual outcome are the reason for such number of scleral contact lens. Keratoplasty (Fullthickness/DALK) was done in 11 eyes (8.52%). Earlier studies have reported that approximately 12-Keratoconusrequire 20% of patients with keratoplasty.[3,28-29] S Rafati et al. [17] also reported that 10% of their patients needed keratoplasty.

#### CONCLUSION

Keratoconus is usually bilateral and predominantly affects males. Keratoconus commonly presents in the second and third decade of life with majority diagnosed in stage 2/3 with mild to moderate visual impairment. Presently Collagen cross linking and sclera lenses are the commonly used modalities to treat keratoconus.

#### Financial Interest - Nil Conflicts of Interest - Nil

#### REFERENCES

- Gokul A, Patel DV, McGhee CNJ. Dr John Nottingham's 1854 landmark treatise on conical cornea considered in the context of the current knowledge of keratoconus. Cornea 2016;35:673-8.
- 2. Chopra I, Jain AK. Between eye asymmetry in keratoconus in an Indian population. ClinExpOptom 2005;88:146-52.
- Kennedy RH, Bourne WM, Dyer JA. A 48-year clinical and epidemiologic study of keratoconus. Am J Ophthalmol 1986;101:267-73.
- Auffarth GU, Wang L, Völcker HE. Keratoconus evaluation using the Orbscan Topography System. J Cataract Refract Surg 2000;26:222-8.
- Godefrooij DA, de Wit GA, Uiterwaal CS, Imhof SM, Wisse RPL. Age-specific incidence and prevalence of keratoconus: A nationwide registration study. Am J Ophthalmol 2017;175:169-72.
- Hashemi H, HeydarianS, HooshmandE, SaatchiM, YektaA, AghamirsalimM, et al. The prevalence and risk factors for keratoconus: A systematic review and meta-analysis. Cornea 2020;39:263-70.
- 7. Gorskova EN, Sevost'ianov EN. [Epidemiology of keratoconus in the Urals]. VestnOftalmol 1998;114:38-40.

- Hashemi H, Khabazkhoob M, Fotouhi A. Topographic keratoconus is not rare in an Iranian population: The Tehran Eye Study. Ophthalmic Epidemiol 2013;20:385-91
- 9. Amsler M. [The "formefruste" of keratoconus]. Wien KlinWochenschr 1961;73:842-3.
- Shilpy N, Shah Z, Singh S, Purohit D. Prevalence of keratoconus in refractive surgery cases in Western India. Middle East Afr J Ophthalmol 2020;27:156-9
- Jonas JB, Nangia V, Matin A, Kulkarni M, Bhojwani K. Prevalence and associations of keratoconus in rural Maharashtra in central India: The central India eye and medical study. Am J Ophthalmol 2009;148:760-5.
- Sharma R, Titiyal JS, Prakash G, Sharma N, Tandon R, Vajpayee RB. Clinical profile and risk factors for keratoplasty and development of hydrops in north Indian patients with keratoconus. Cornea 2009;28:367-70.
- Georgiou T, Funnell CL, Cassels-Brown A, O'Conor R. Influence of ethnic origin on the incidence of keratoconus and associated atopic disease in Asians and white patients. Eye LondEngl 2004;18:379-83.
- Ertan A, Muftuoglu O. Keratoconus clinical findings according to different age and gender groups. Cornea 2008;27:1109-13.
- Assiri AA, Yousuf BI, Quantock AJ, Murphy PJ. Incidence and severity of keratoconus in Asir province, Saudi Arabia. Br J Ophthalmol 2005;89:1403-6.
- Hwang S, Lim DH, Chung TY. Prevalence and incidence of keratoconus in South Korea: A nationwide population-based study. Am J Ophthalmol 2018;192:56-64.
- Rafati S, Hashemi H, Nabovati P, DoostdarA, YektaA, Aghamirsalim M, et al. Demographic profile, clinical, and topographic characteristics of keratoconus patients attending at a teriary eye center. J CurrOphthalmol 2019;31:268-74
- Ziaei H, Jafarinasab MR, Javadi MA, Karimian F, Poorsalman H, Mahdavi M, et al. Epidemiology of keratoconus in an Iranian population. Cornea 2012;31:1044-7.

- 19. Khor WB, Wei RH, Lim L, Chan CM and Tan DTH: Keratoconus in Asians: Demographics, clinical characteristics and visual function in a hospital-based population. ClinExpOphthalmol 39: 299-307, 2011.
- 20. Mohd-Ali B, Abdu M, Das S and Mohidin N: Ethnicity related to keratoconus: A study with clinical implications. Int Med J 18: 237-240, 2011.
- 21. Das AV, Deshmukh RS, Reddy JC, Joshi VP et al. Keratoconus in India: Clinical presentation and demographic distribution based on big data analytics. Indian J Ophthalmol. 2024 Jan 1;72(1):105-110.
- Wagner H, Barr JT, Zadnik K. Collaborative Longitudinal Evaluation of Keratoconus (CLEK) Study: methods and findings to date. Cont Lens Anterior Eye. 2007;30:223-32. doi: 10.1016/j.clae.2007.03.001. Epub 2007 May 3.
- Almusawi LA, Hamied FM. Risk factors for development of keratoconus: a matched pair casecontrol study. ClinOphthalmol 2021;15:3473–9.
- Ahuja P, Dadachanji Z, Shetty R, Nagarajan SA, Khamar P, Sethu S, et al. Relevance of IgE, allergy and eye rubbing in the pathogenesis and management of Keratoconus. Indian J Ophthalmol 2020;68:2067– 74.
- 25. Claessens JLJ, Godefrooij DA, Vink G, Frank LE, Wisse RPL. Nationwide epidemiological approach to identify associations between keratoconus and immune-mediated diseases. Br J Ophthalmol 2021.
- 26. Serdarogullari H, Tetikoglu M, Karahan H, Altin F and Elcioglu M: Prevalence of keratoconus and subclinical keratoconus in subjects with astigmatism using pentacam derived parameters. J Ophthalmic Vis Res 8: 213-219, 2013.
- 27. Mahadevan R, Arumugam AO, Arunachalam V and Kumaresan B: Keratoconus-a review from a Tertiary Eye-care center. J Optom 2: 166-172, 2009
- Gordon MO, Steger-May K, Szczotka-Flynn L, Riley C, Joslin CE, Weissman BA, et al. Baseline factors predictive of incident penetrating keratoplasty in keratoconus. Am J Ophthalmol 2006;142:923-30.
- 29. Tuft SJ, Moodaley LC, Gregory WM, Davison CR, Buckley RJ. Prognostic factors for the progression of keratoconus. Ophthalmology 1994;101:439-47