



**SPECIFIC PRACTICE INTERESTS**

**In Office Comprehensive sinus surgery with Balloon sinuplasty with image guidance.**

**Comprehensive management of chronic sinusitis**

**Allergic rhinitis**

**Rhinoplasty and facial plastic surgery including nose and eyelid surgery**

**Meniere's disease and sudden hearing loss**

**Sleep Medicine**

**Thyroid and parathyroid disorders**

**Hearing loss and balance disorders**

**RESEARCH FUNDING INFORMATION**

1992-1993

NIH grant T32 HL07627

**CURRENT MANUSCRIPTS IN PREPARATION**

- 1. In-office comprehensive balloon sinuplasty with subtotal ethmoidectomy (CBSSE) for stage III and IV CRSwNP and CRSsNP: feasibility, short term clinical and CT outcomes, challenges and pitfalls.**
- 2. Allergic fungal sinusitis (1-10 yr follow up): characteristics and intermediate to long term outcomes. Abstract for poster presentation, American Rhinology Society, Vancouver, Canada**
- 3. Allergy profile in chronic rhinosinusitis.**
- 4. A new comprehensive descriptive staging and scoring system for comparing outcomes of treatment in chronic rhinosinusitis.**

**CURRENT RESEARCH**

WESTERN INSTITUTIONAL REVIEW BOARD APPROVAL FOR "INVESTIGATIONS INTO CHRONIC RHINOSINUSITIS, ALLERGIC RHINITIS AND SINUS HEADACHES- CLINICAL AND MOLECULAR CORRELATES".

**PREVIOUS PUBLICATIONS**

**Khetarpal U:** DFNA9 is a progressive audiovestibular dysfunction with a microfibrillar deposit in the inner ear. *Laryngoscope* 2000;110:1379-1384.

Gacek RR, **Khetarpal U:** Neurotrophin 3, not Brain-Derived Neurotrophic factor or Neurotrophin 4, knockout mice have delay in vestibular compensation after unilateral labyrinthectomy. *Laryngoscope* 1998; 108: 671-678.

Gacek RR, Schoonmaker J, **Khetarpal U:** Morphological and neurochemical correlates of vestibular compensation. *Auris Nasus Larynx* 1998.

**Khetarpal U, Morton CC:** Inner ridge cells may be the source of type II collagen in human fetal tectorial membrane: a quantitative *in situ* hybridization study. *Acta Otolaryngol*, 1998.

Halpin C, **Khetarpal U**, McKenna MM: Autosomal dominant progressive audio-vestibular dysfunction in a large North American family. *Am J Audiol*, 1996; 5: 105.

Manolis E, ..., **Khetarpal U**, et al: A gene for non-syndromic autosomal dominant progressive postlingual sensorineural hearing loss maps to chromosome 14q12-13. *Hum Mol Genet* 1996; 5: 1047-1050.

Rehm HL, ..., **Khetarpal U**, et al.: Norrie disease gene mutation in a large Costa Rican kindred with a novel phenotype including venous insufficiency. *Human Mutation* 1996; 9: 402-408.

**Khetarpal U**, Morton CC: Expression of COL1A2 and COL2A1 mRNA in a temporal bone of lethal osteogenesis imperfecta. *Arch Otolaryngol Head Neck Surg*, 1993; 119: 1305-1314.

**Khetarpal U**, Robertson NG, Yoo TJ, Morton CC: Expression and localization of COL2A1 mRNA and type II collagen in human fetal cochlea. *Hear Res* 1994; 79, 59-73.

Magovcevic I, **Khetarpal U**, Bieber FR, Morton CC: GNAZ in human fetal cochlea: expression, localization, and potential role in inner ear function. *Hear Res* 1995; 90: 55-64.

Robertson NG, **Khetarpal U**, Guitierrez-Espelela G, Bieber FR, Morton CC: Isolation of novel and known genes from a human fetal cochlear cDNA library using subtractive hybridization and differential screening. *Genomics* 23: 42-50, 1994.

**Khetarpal U**: Autosomal dominant sensorineural hearing loss - further temporal bone findings. *Arch Otolaryngol Head Neck Surg* 1993; 119: 106-108.

**Khetarpal U**: Investigations into the cause of vertigo in sudden sensorineural hearing loss. *Otolaryngol Head Neck Surg*. 1992; 105: 360-371.

**Khetarpal U**, Schuknecht HF, Gacek RR, Holmes LH: Autosomal dominant sensorineural hearing loss - Pedigrees, audiological and temporal bone findings in two kindreds. *Arch Otolaryngol Head Neck Surg*. 1991; 117: 1032-1042

**Khetarpal U**, Nadol JB, Glynn RJ: Idiopathic sensorineural hearing loss and postnatal viral labyrinthitis - a statistical comparison of temporal bone findings. *Ann Otol Rhinol Laryngol* 1990; 99: 969-976.

**Khetarpal U**, Schuknecht HF: In search of pathological correlate of hearing loss in Paget's disease. A study of 26 temporal bones. *Ann Otol Rhinol Laryngol* 1990; 145: 1-16.

Kumar L, **Khetarpal U**, et al. Intracranial Plasmacytoma. *J Ind Med Assoc* 1989; 20-24.

## ABSTRACTS

**Khetarpal U**, Deka RC, Kacker SK: Brainstem auditory potentials in cerebello-pontine angle tumors. Abstract, Sixth Asia Oceania Congress of Otorhinolaryngological Societies, New Delhi, India, Nov. 9-13, 1987.

Morton CC, ..., **Khetarpal U**, et al: Isolation of novel and known genes from a human fetal cochlear cDNA library using subtractive hybridization and differential screening. Abstract, Association for Research in Otolaryngology Meeting, Feb. 5-9, 1995, Fla.

**Khetarpal U**, et al: Dystrophin, cochlea and hearing-hearing loss. Abstract #45, Molecular Biology of Hearing and Deafness Meeting, October 6-8, 1995, Bethesda, MD.

**Khetarpal U**, Morton CC: Tissue specific expression of type II procollagen mRNAs and their role in chondrogenesis and inner ear morphogenesis. Abstract #28, Molecular Biology of Hearing and Deafness Meeting, October 6-8, 1995, Bethesda, MD.

**Khetarpal U**, Morton CC: The role of aggrecan and its alternatively spliced products in human fetal morphogenesis and chondrogenesis. Abstract # 10, Association for Research in Otolaryngology Meeting, Feb. 2-6, 1997, St. Petersburg, Fla.

**Khetarpal U**, Morton CC: Aggrecan is expressed in noncartilaginous human fetal tissue. Variable alternative splicing of EGF-like domains during development and mRNA localization in human fetal tissues.

**Khetarpal U**: Genetic mechanisms that determine tissue specificity during human development in general and cochlear development in particular. Abstract, International Otopathological Society Meeting, Boston, June 15-17, 1997.

### **BOOK CHAPTERS**

Krespi Y, **Khetarpal U**: Conservation Laryngeal Surgery. In *Complications of Head and Neck Surgery*, Eds. Krespi Y, Ossoff RJ. WB Saunders Co, Philadelphia, 1993, 215-232.

**Khetarpal U**, Lalwani A: Nonsyndromic hereditary hearing loss. In *Pediatric Otology and Neurotology*, Eds. Grundfast KG, Lalwani A. Lippincott-Raven Press, Philadelphia, 1998.

### **INVITED SPEAKER**

1) X-linked hearing loss. Mapping Genes for Deafness Workshop, NIH sponsored, New Delhi, India, Jan. 1992

2) Autosomal dominant sensorineural hearing loss. Mapping Genes for Deafness Workshop, NIH sponsored, New Delhi, India, Jan. 1992

3) Molecular correlates of audiovestibular development and disease. Department of Otolaryngology, University of Pennsylvania Medical Center, Jan 28, 1998.