Saba Tasneem

Publications

1. Anwar, S., Riazuddin, S., Ahmed, Z. M., **Tasneem, S**., Ateeq ul, J., Khan, S. Y., . . . Riazuddin, S. (2009). SLC26A4 mutation spectrum associated with DFNB4 deafness and Pendred's syndrome in Pakistanis. *J Hum Genet, 54*(5), 266-270. doi: 10.1038/jhg.2009.21. (IF: 2.487)

2. Khan, S. Y., Ahmed, Z. M., Shabbir, M. I., Kitajiri, S., Kalsoom, S., **Tasneem, S**., . . . Riazuddin, S. (2007)v. Mutations of the RDX gene cause nonsyndromic hearing loss at the DFNB24 locus. *Hum Mutat, 28*(5), 417-423. doi: 10.1002/humu.20469. (IF: 5.089)

3. Nal, N., Ahmed, Z. M., Erkal, E., Alper, O. M., Luleci, G., Dinc, O., . . . Friedman, T. B. (2007). Mutational spectrum of MYO15A: the large N-terminal extension of myosin XVA is required for hearing. *Hum Mutat, 28*(10), 1014-1019. doi: 10.1002/humu.20556. (IF: 5.089)

International Symposia/ Meetings Attended

1. Satellite Symposium on **"Reprogramming of Somatic Cells for the Therapy of Heart Disorders"** October 9-10, 2007 Hilton Hotel Düsseldorf, Germany. Funded by BMBF Program "Cell-Based Regenerative Medicine".

2. **4th International Meeting Stem Cell Network North Rhine-Westphalia**, October 8-9, 2007, Düsseldorf, Germany.

3. First Symposium of the BMBF-sponsored Research Alliance on **"Induction of Immunologic Tolerance by Mesenchymal Stem Cells"** September 12-13, 2007, Düsseldorf, Germany.