Farheena Iqbal Awan

Publications in chronologically desending order:

AHMED, Z., BABAR, M. E., HUSSAIN, T., NADEEM, A., AWAN, F. I., WAJID, A., SHAH, S. A. & ALI, M. M. 2014. Genetic diversity analysis of kail sheep by using microsatellite markers. *Journal of Animal and Plant Sciences,* 24**,** 1329-1333.

SABAR, M. F., AWAN, F. I., SHAHID, M., GHANI, M. U. & YAQUB, M. 2013. Synthesis and Bioactivity Study of 30KDa Linear PEG-Interferon and its Comparison with Tri-Branched PEG-Interferon. *Journal Chemical Society of Pakistan,* 35**,** 119-124.

ZIL, E. H. B., NOREEN, L., INNA, A. B., FARHEENA, I., SHEIKH AMER, R., SHAHEEN, N. K., THOMAS, B. F., SHEIKH, R. & SAIMA, R. 2012. Phenotypic variability of CLDN14 mutations causing DFNB29 hearing loss in the Pakistani population. *Journal of Human Genetics,* 58**,** 102.

SABIR, N., RIAZUDDIN, S. A., BUTT, T., IQBAL, F., NASIR, I. A., ZAFAR, A. U., QAZI, Z. A., BUTT, N. H., KHAN, S. N., HUSNAIN, T., HEJTMANCIK, J. F. & RIAZUDDIN, S. 2010a. Mapping of a new locus associated with autosomal recessive congenital cataract to chromosome 3q. *Molecular vision,* 16**,** 2634.

SABIR, N., RIAZUDDIN, S. A., KAUL, H., IQBAL, F., NASIR, I. A., ZAFAR, A. U., QAZI, Z. A., BUTT, N. H., KHAN, S. N., HUSNAIN, T., HEJTMANCIK, J. F. & RIAZUDDIN, S. 2010b. Mapping of a novel locus associated with autosomal recessive congenital cataract to chromosome 8p. *Molecular vision,* 16**,** 2911.