

## COMPLETE LIST of PUBLICATIONS (Impact factor 368.822)

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65. Kanwal A, Pardo JV\*, **Naz S\*** [<sup>\*</sup>corresponding authors] (2022) RGS3 and IL1RAPL1 missense variants implicate defective neurotransmission in early-onset inherited schizophrenia **J Psychiatry Neurosci** 47:E379-E390
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67. Manzoor H, Brüggemann N, Hussain HMJ, Bäumer T Hinrichs F, Wajid M, Münchau A, Lohmann K\*, **Naz S\*** [<sup>\*</sup>corresponding authors] (2023) A novel ECEL1 variant associated with a congenital contracture disorder. **Pak J Zool** 55: 391-395
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71. Shafique, A, Sultan T, Alzahrani F, Seo GH, Alkuraya FS, **Naz S** (2023) Genomic analysis of multiplex consanguineous families reveals causes of neurodevelopmental disorders with epilepsy **Gene** 879:147599 1-8
72. Ishaq T, Loid P, Ishaq HA, Seo GH, Mäkitie O\*, **Naz S\*** [<sup>\*</sup>corresponding authors] (2023) Clinical, radiographic and molecular characterization of two unrelated families with Multicentric Osteolysis, Nodulosis, and Arthropathy **BMC Musculo** 24:735 1-11
73. Kanwal A , Sheikh SA, Aslam F, Yasin S, Beethem Z, Pankratz N, Clabots CR, **Naz S\***, Pardo JV\* [<sup>\*</sup>corresponding authors] (2023) Genome sequencing of consanguineous family implicates ubiquitin specific protease 53 (USP53) variant in psychosis/schizophrenia: Wild type expression in murine hippocampal CA 1-3 and granular dentate with AMPA synapse interactions **Genes** 14:1921 1-23
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## **EDITED BOOKS WITH CHAPTERS**

**Naz S** (2012). "Hearing Loss" ed **Naz S**, Intech, Croatia. (ISBN 979-953-307-271-4)

**Naz S** (2022) "Auditory System - Function and Disorders", Intech, Croatia (ISBN 978-1-80355-190-6)

## **CHAPTERS IN BOOKS**

1. Naz S (2012) Genetics of nonsyndromic recessively inherited moderate to severe and progressive deafness in humans. In: "Hearing Loss" ed Naz S, Chapter 12, pp 247-274. Intech, Croatia. (ISBN 979-953-307-271-4)

2. Naz S (2022) Nonreceptor protein kinases and phosphatases necessary for auditory function. In: "Auditory System - Function and Disorders" ed Naz S, Chapter 7, pp 1-18. Intech, Croatia (ISBN 978-1-80355-190-6)
3. Naz S, Makitie O (2023). Acromesomelic Dysplasia I, Maroteaux Type (AMD1). In: Rezaei, N. (eds) *Genetic Syndromes*. Springer, Cham. [https://doi.org/10.1007/978-3-319-66816-1\\_1382-1](https://doi.org/10.1007/978-3-319-66816-1_1382-1)
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## **CONFERENCES AND SEMINARS**

32 abstracts published in national and international proceedings [4 won ASHG Developing Countries Awards]

## **RESEARCH AWARDS FOR RESEARCH ABSTRACTS**

[2017 ASHG, USA, Developing Countries award \(as first author\)](#)

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