In the Name of The Most Beautiful

Curriculum Vitae

Newsha Molavi

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CONTACT & PROFILES

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Educational Background

M.Sc., Human Genetics

(2016-2020)

School of Medicine, Isfahan University of Medical Sciences (MUI), Isfahan, Iran

- **Thesis:** Genetics and Bioinformatics Investigation of The *CYP21A2* Gene Mutations among Patients with Congenital Adrenal Hyperplasia Referred to Imam Hossein Hospital
- **Total CGPA:** 3.93/4 (17.55/20 in Iranian Scale)

B.Sc., Molecular and Cellular Biology – Genetics

(2011-2015)

Faculty of Biological Science and Technology, University of Isfahan, Isfahan, Iran

• **Total CGPA:** 3.82/4 (17.39/20 in Iranian Scale)



Publications

1- Aghaei S, Parvizpour S, Farrokhi E, <u>Molavi N</u>, Hoseinzadeh M, Tabatabaiefar MA. Characterization of a novel androgen receptor gene variant identified in an Iranian family with complete androgen insensitivity syndrome (CAIS): a molecular dynamics simulation study. J Biomol Struct Dyn. 2022 Nov 21;0(0):1–15.

DOI: https://doi.org/10.1080/07391102.2022.2148125

2- Abbasi B, Molavi N, Tavalaee M, Abbasi H, Nasr-Esfahani MH. Alpha-lipoic acid improves sperm motility in infertile men after varicocelectomy: a triple-blind randomized controlled trial. Reprod Biomed Online. 2020 Dec 1;41(6):1084–91.

DOI: https://doi.org/10.1016/j.rbmo.2020.08.013

3- Hassanzadeh S, Sadeghi S, Jafari M, Najafi S, <u>Molavi N</u>, Sherkat R. Ciliary and immune dysfunctions and their genetic background in patients with non-cystic fibrosis bronchiectasis in Central Iran. Ir J Med Sci 1971. 2022 Apr 7

DOI: https://doi.org/10.1007/s11845-022-02994-z

- 4- Shahhoseini M, <u>Molavi N</u>, Tabatabaiefar MA, Sehhati M. Implementation and Optimization of Annotation and Interpretation Step of Next-Generation Sequencing Data for Non-Syndromic Autosomal Recessive Hearing Loss. J Health Biomed Inform. 2021 Mar 10;7(4):435–44.
- 5- Hoseinzadeh M, **Molavi N**, Norouzi M, Aghaei S, Zeinalian M, Hashemipour M, et al. A Novel Homozygous Pathogenic Variant in CYP11B1 in a Female Iranian Patient with 11B Hydroxylase Deficiency. Lab Med. 2023 Jul 1;54(4):439–46.

DOI: https://doi.org/10.1093/labmed/lmac141

6- Aghaei S, Farrokhi E, Saffari-Chaleshtori J, Hoseinzadeh M, **Molavi N**, Hashemipour M, et al. New molecular insights into the A218V variant impact on the steroidogenic acute regulatory protein (STAR) associated with 46, XY disorders of sexual development. Mol Genet Genomics. 2023 May;298(3):693–708.

DOI: https://doi.org/10.1007/s00438-023-02006-4

- 7- Khoshnevisan R, Hassanzadeh Sh, Klein Ch, Rohlfs M, Grimbacher B, Khoshnevisan A, Zamanifar A, Molavi N, Jafari M; Behnam M, Sherkat R. Predominant Antibody Deficiency and Absence of B-cell in a Cohort Study of Iranian Patients with Inborn Error of Immunity between 1976 and 2019 (Submitted).
- 8- Molavi N, Hoseinzadeh M, Aghaei Sh, KhanAhmad H, Hashemipour M, Tabatabaiefar MA. A novel homozygous frameshift mutation in the *CYP21A2* gene leads to classic salt-wasting congenital adrenal hyperplasia in Iran. (Under Review)



Professional Experience

Research Assistant

(2021-present)

 Acquired Immunodeficiency Research Center, Isfahan University of Medical Sciences, Isfahan, Iran

Teaching Assistant for 'Practical Methods of Molecular Genetics'

(2021-2022)

- Presented by Dr. Mohammad Amin Tabatabaiefar
- Department of Genetics and Molecular Biology, School of Medicine, Isfahan University of Medical Sciences, Isfahan, Iran

Instructor for 'Next-Generation-Sequencing data analysis workshop'

(2022)

• Isfahan University of Medical Sciences, Isfahan, Iran

Research Assistant (2018-2019)

• Department of Animal Biotechnology, Reproductive Biomedicine Research Center, Royan Institute for Biotechnology, ACECR, Isfahan, Iran



Technical Skills

Laboratory skills:

- o Autozygosity mapping with STR markers
- General Molecular biology techniques
- Karyotyping
- o Fundamentals of cell culture
- Laboratory animal handling
- o SDS page electrophoresis and western-blotting
- o Basics of gene editing tools

Bioinformatics Skills:

- o Primer designing software: Oligo 7, AlleleID, Gene-runner
- o Interpretation of Sanger and Next-Generation Sequencing data

Scripting Language: UNIX in big data analysis

Language Proficiency

English: Fluent (TOEFL: total score 85)

Persian: Native

Research Interests

- Bioinformatics - Medical Genetics - Molecular Genetics

- Mechanisms of human diseases - Molecular Diagnostics

All Degrees and Certificates are available on request

Update on Nov, 2023.