



TO MAGNIFICO RETTORE OF UNIVERSITA' DEGLI STUDI DI MILANO

ID CODE \_\_ 5813□ \_\_\_\_\_

I the undersigned asks to participate in the public selection, for qualifications and examinations, for the awarding of a type A fellowship at Dipartimento di \_\_Dipartimento di Bioscienze\_\_\_\_\_

Scientist- in - charge: \_\_\_\_ Prof. Roberto Mantovani \_\_\_\_\_

[Khushboo]

## CURRICULUM VITAE

### PERSONAL INFORMATION

Surname	Sina
Name	Mohammad

### PRESENT OCCUPATION

Appointment	PhD student
Not Applicable	

### EDUCATION AND TRAINING

Degree	Course of studies	University	year of achievement of the degree
Degree	Bachelor of Cellular & Molecular Biology -Biotechnology,	Rabe Rashid University, Iran	2011
Specialization	Not Applicable	Not Applicable	Not Applicable
PhD	student of Molecular Genetics, Biotechnologies and Experimental Medicine, University of Brescia, Brescia	University of Brescia, Brescia	Thesis will be submitted on 25 st June, 2023
Master	Master of Science MSc in Medical Biotechnology,	Tabriz University of Medical Sciences,Iran	2015



Degree of medical specialization	Not Applicable	Not Applicable	Not Applicable
Degree of European specialization	Not Applicable	Not Applicable	Not Applicable
Other	Not Applicable	Not Applicable	Not Applicable

## REGISTRATION IN PROFESSIONAL ASSOCIATIONS

Date of registration	Association	City
Not Applicable	Not Applicable	Not Applicable

## FOREIGN LANGUAGES

Languages	level of knowledge
English	Proeficient

## AWARDS, ACKNOWLEDGEMENTS, SCHOLARSHIPS

Year	Description of award
2022	Travel grant award for US \$1700 by International society of Neurochemistry (ISN-APSN 2022) to participate in international conference in neurochemistry held at Honolulu, Hawaii during 28 August to 01 September 2022.

## TRAINING OR RESEARCH ACTIVITY

Moderate skill in R programming and Linux operating system
1) iPSC culture and EBV transfection 2) PBMCs isolation from peripheral blood 3) Flow cytometry 4) Sequencing Analysis 5) Variations filtering and interpretation using ANNOVAR, Samtools, Varaft, and IGV 6) Copy Number Variation Analysis (CNVs) using HMZDelFinder, ExomeDepth, cnmops.panel, and CoNVaDING, GATKcNV, 7) removing low mappability regions 8) Cytogenetics and karyotyping 9) RNA bulk sequencing, RUV III, PRPS



- 10) Flow cytometry
- 11) NGS library preparation, first-hand experience with PGM ion torrent chef
- 12) Fragment Analysis, including MSI
- 13) Humara Assay
- 14) DNA extractions from tissue and blood,
- 15) DNA purification from gel
- 16) PCR, touchdown PCR, Touch-up PCR, Long-range PCR
- 17) RNA extractions, RT-PCR, and Real-time PCR
- 18) Soluble expression of recombinant protein
- 19) Protein purification (pull-down)
- 20) SDS-PAGE
- 21) Cell culture
- 22) Bacterial culture
- 23) Mouse model management
- 24) Western blotting
- 25) Writing a systematic review
- 26) GWAS analysis
- 27) MLPA

## PROJECT ACTIVITY

Year	Project
2020	I am a member of the Middle-East Network on Hereditary Colorectal Cancer (HCCN-ME) ( <a href="https://www.hccn-me.com/">https://www.hccn-me.com/</a> )
2019	I have established lynchsyndrome.ir website to raise awareness among Iranian patients diagnosed with Lynch Syndrome
2019	I established a workshop on Genome-wide association study at Motamed cancer institute, Tehran, Iran
2019	I have established microsatellite instability testing at Motamed cancer institute, Tehran, Iran
2019	1) I have established KRAS and NRAS genetic testing at Motamed cancer institute, Tehran, Iran



PATENTS

Patent
Not Applicable

CONGRESSES AND SEMINARS

Date	Title	Place
2022	ACMG Annual Clinical Genetics Meeting – American College Of Medical Genetics and Genomics.	Nashville, Tennessee, USA
2018	Conference speaking engagements "Current health care of families with Lynch syndrome in the Middle East and North African countries, conference of prevention colorectal cancer and the first Iranian hereditary colorectal cancer meeting	Mashhad, Iran
2019	Attended health research strategies in EU (Horizon 2020) and Iran AND training for applicants to horizon 2020 project	Imam Khomeini hospital, Tehran University of Medical, Tehran, Iran
2019	Attended 9th conferences on the new approaches of diagnosis and the treatment of colorectal cancer	

PUBLICATIONS

Books (Not Applicable)
Sina M, Farajzadeh D, Dastmalchi S, 2015, Effects of Environmental Factors on Soluble Expression of a Humanized Anti-TNF- $\alpha$ scFv Antibody in Escherichia coli. Adv Pharm Bull, 5(4), 455-461. DOI: 10.15171/apb.2015.062
1) Sina, M., Ghorbanoghli, Z., Abedrabbo, A. et al. Identification and management of Lynch syndrome in the Middle East and North African countries: outcome of a survey in 12 countries. Familial Cancer (2020). <a href="https://doi.org/10.1007/s10689-020-00211-3">https://doi.org/10.1007/s10689-020-00211-3</a>
[title, place, publishing house, year ...]

Projects
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1) <b>Sina, M.</b> , Ghorbanoghli, Z., Abedrabbo, A. et al Current health care of families with Lynch syndrome in the Middle East and North African countries, ready to submit
2) <b>Sina, M.</b> , Ghorbanoghli, Z, H. F. A Vasen, et al Identification and surveillance of Four Iranian Families with Lynch Syndrome and a family with <i>PMS1</i> pathogenic variants, ready to submit.
3) <b>Sina, M.</b> Giliani S, A de novo novel pathogenic variant in <i>CYBB</i> gene caused exon skipping in a female patient with Chronic granulomatous disease: a case report study. ready to submit
4) <b>Sina, M.</b> Giliani S, et al Copy number variation analysis on exome sequencing data identified a 70 kb deletion in <i>MSH2</i> and <i>EPCAM</i> genes in two Iranian families. Ongoing
5) Julio, <b>Sina, M.</b> Giliani S, et al A de novo novel pathogenic variant in <i>MAGT1</i> gene caused exon skipping: a case report study. ongoing
6) <b>Sina M.</b> , Eidi M., Ferarro R. et al Copy number variant calling in a cohort of primary immunodeficiency disease using panel sequencing in diagnostic clinic: ongoing

Congress proceedings (Not Applicable)
[title, structure, place, year]
[title, structure, place, year]
[title, structure, place, year]

OTHER INFORMATION

<b>Broad Research Areas:</b> Medical genetics, DNA sequencing
<b>Title of Ph.D.Thesis:</b> “identification of human genetics mendian disorders”.
<b>Dissertation topic at M.Sc. level:</b> “Effects of Environmental Factors on Soluble Expression of a Humanized Anti-TNF- $\alpha$ scFv Antibody in Escherichia coli.

Declarations given in the present curriculum must be considered released according to art. 46 and 47 of DPR n. 445/2000.

The present curriculum does not contain confidential and legal information according to art. 4, paragraph 1, points d) and e) of D.Lgs. 30.06.2003 n. 196.

Please note that CV WILL BE PUBLISHED on the University website and It is recommended that personal and sensitive data should not be included. This template is realized to satisfy the need of publication without personal and sensitive data.

Please DO NOT SIGN this form.

Place and date:Sina, Brescia (15 JUN, 2023)

