Curriculum Vitae

Dr Sima Mansoori Derakhshan, M.D, Ph.D
Associated Professor of Medical Genetics

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EDUCATION

Sep 2010–March 2011 — **Tabriz University of Medical Sciences**
Education and Development Center
Tabriz, Iran

Fellowship in Academic Skills
Consisting: Educational management and planning,educational counseling, program evaluation, teaching and research methodology; and student assessment

Jan 2004 – Feb 2008 — **University of Melbourne**
Faculty of Medicine, Dentistry and Health Sciences
Paediatrics Department
Murdoch Children's Research Institute
Cell and Gene Therapy Group
Melbourne_Australia

Ph.D in Medical Genetics
Thesis title: Antisense therapy for IVSI-110 mutation in β-thalassemia
This study aims to provide a gene therapy strategy for β-thalassemia.

Feb 2006 – Feb 2007 **University of Melbourne**
Faculty of Medicine, Dentistry and Health Sciences
Paediatrics Department
Murdoch Children's Research Institute
Melbourne_Australia
Graduate Diploma in Genetic Counceling

Oct 1991 – Jul 1998 **Tabriz University of Medical Sciences**
Tabriz_ IRAN
Medical Doctor (MD)

1. Thesis title: Retrospective study on side effects of cataract surgery in 50 diabetic patients in Nicoucari hospital in 1997

WORK HISTORY

2008-Now **Academic Staff**
Department of Medical Genetics
Faculty of Medicine
Tabriz University of Medical Sciences & Health Services
Attar Nishabori Road
Golgasht, Post code: 5166614711
Tabriz, Iran

2009-Now **Director of Ebne Sina Medical Genetic Centre**
Specialized and Sub specialized Outpatient Clinic of Tabriz Medical Science University
Baharan Clinic, Azadi Ave, Tabriz, Iran.

2007-2008 **Research assistant**
Murdoch Children's Research Institute
Cell and Gene Therapy Group
Melbourne, Australia

Responsibilities:
Working in gene therapy in β-thalassemia projects

2001-2004 **Head of Urban health center**
Tabriz, East Azerbaijan
IRAN

1999-2001 **Thalassaemia coordinator in health centre**
Bonab Health center
Bonab, East Azarbaijan
IRAN

1998-2001 **General physician in emergency unit**
Imam Khomeini Hospital- Bonab
East Azerbaijan
IRAN

1998-2001 **Head of Urban health center**
Bonab, East Azerbaijan
IRAN

PUBLICATIONS

1. The role of mir-151a-5p in tumorigenesis; A systematic review
   A Ebrahimi, SM Derakhshan, D Ghavi, Z Forouzandeh, S Hashemi

   Pathology-Research and Practice, 154576 2023

2. Evaluating the relationship between Ex3 vNTR and rs3758653 polymorphisms in DRD4 genes in children with attention deficit hyperactivity disorder with respect to the dose of …
   S Amiri, SM Negar, S Farhang, SM Derakhshan, MS Khaniani, LM Fanid

   Medical Journal of Tabriz University of Medical Sciences 44 (5), 390-402 2022

3. In Silico and Experimental Analysis of miR-125b-5 and miR-485-5p Expression in Serum of Patients with Breast Cancer.
   Z Bahmanpour, R Sheervalilou, MS Khaniani, A Poursheikhani, ...

   Microrna (Shariqah, United Arab Emirates) 2022

4. A novel missense variant in ESRRB gene causing autosomal recessive non-syndromic hearing loss: in silico analysis of a case
   T Ghasemnejad, M Shekari Khaniani, J Nouri Nojadeh, ...

   BMC Medical Genomics 15 (1), 18 2022

5. Analysis of Association Between the Effects of Methylphenidate and DRD4 Gene Polymorphisms in Patients with Attention Deficit Hyperactivity Disorder
   S Amiri, S Farhang, MS Khaniani, SM Derakhshan, A Zadfattah, ZM Bina, ...

   Journal of Comprehensive Pediatrics 12 (4) 2021

6. Identification of multi-exon deletion in the COL7A1 gene underlying dystrophic epidermolysis bullosa by whole-exome sequencing
   M Taghizadeh, SM Derakhshan, MS Khaniani, Y Eshaghkhani, ...

   Our Dermatology Online/Nasza Dermatologia Online 12 (4) 2021

7. Association between the rs8028440 polymorphism of CYFIP 1 gene in autism patients
   HG Almanghadim, M Assefi, S Masoumi, P Vakili, Z Shams, A Moradkhani, ...

   Biomedical and Translational Science 1 (3), 1-6 2021

8. Molecular Analysis of STin2 (Intron 2) Variant of The SLC6A4 Gene in Children and Adolescents With Attention Deficit Hyperactivity Disorder
   S Amiri, M Asadian, MS Khaniani, SM Derakhshan, NP Rahmani, ...

   2021

9. A Novel Missense Variant in the ESRRB Gene Causing Nonsyndromic Hearing Loss: In Silico Analyses of a Case
   T Ghasemnejad, MS Khaniani, JN Nojadeh, SM Derakhshan

   2021

10. The worldwide frequency of MYO15A gene mutations in patients with non-syndromic hearing loss: A meta-analysis
    M Farjami, R Assadi, FA Javan, M Alimardani, S Eslami, SM Derakhshan, ...

    Iranian Journal of Basic Medical Sciences 23 (7), 841 2020

11. Investigating the prevalence of point mutations in the human mutL homolog 1 gene in colorectal cancer patients in the Northwest of Iran
    M Gholami, MS Khaniani, SM Derakhshan, A Esfahani

    Gaceta mexicana de oncología 18 (4), 258-264 2019

12. A new insight on serum microRNA expression as novel biomarkers in breast cancer patients
    Z Bahmanpour, R Sheervalilou, J Choupani, M Shekari Khaniani, ...

    Journal of Cellular Physiology 234 (11), 19199-19211 2019

13. The hopeful anticancer role of oleuropein in breast cancer through histone deacetylase modulation
    N Mansouri, MR Alivand, S Bayat, MS Khaniani, SM Derakhshan

    Journal of Cellular Biochemistry 120 (10), 17042-17049 2019

14. Differential Expression Pattern of Epithelial Mesenchymal Transition Gens: AXL, GAS6, Claudin-1, and Cofilin-1, in Different Stages of Epithelial Ovarian Cancer
    E Hassani, MS Khaniani, M Saffari, AE Razavi, R Shirkoohi, ...

    Iranian Journal of Public Health 48 (9), 1723 2019

15. Downregulation of HDAC2 and HDAC3 via oleuropein as a potent prevention and therapeutic agent in MCF-7 breast cancer cells
    S Bayat, S Mansoori Derakhshan, N Mansoori Derakhshan, ...

    Journal of cellular biochemistry 120 (6), 9172-9180 2019

16. Oleuropein inhibits migration ability through suppression of epithelial-mesenchymal transition and synergistically enhances doxorubicin-mediated apoptosis in MCF-7 cells
    J Choupani, MR Alivand, S M. Derakhshan, M Zaeifizadeh, M S. Khaniani

    Journal of cellular physiology 234 (6), 9093-9104 2019

17. Targeted mutation analysis of the SLC26A4, MYO6, PJVK and CDH23 genes in Iranian patients with AR nonsyndromic hearing loss
M Alimardani, SM Hosseini, MS Khaniani, MR Haghi, A Eslahi, M Farjami, ...

Fetal and Pediatric Pathology 38 (2), 93-102 2019

18. The associations between 5-HTTLPR polymorphism of SLC6A4 gene and autistic disorder in North West of Iran
SM Derakhshan, F Taheri, S Amiri, LS Khanian, MS Khanian

Medical Journal of Tabriz University of Medical Sciences 40 (5), 83-90 2018

19. Simultaneous downregulation of miR-21 and miR-155 through oleuropein for breast cancer prevention and therapy
M Abtin, MR Alivand, MS Khaniani, M Bastami, M Zaeifizadeh, ...

Journal of cellular biochemistry 119 (9), 7151-7165 2018

20. HLA-G regulatory variants and haplotypes with susceptibility to recurrent pregnancy loss
N Yazdani, M Shekari Khaniani, M Bastami, T Ghasemnejad, F Afkhami, ...

International Journal of Immunogenetics 45 (4), 181-189 2018

21. Narrower insight to SIRT1 role in cancer: A potential therapeutic target to control epithelial–mesenchymal transition in cancer cells
J Choupani, S Mansoori Derakhshan, S Bayat, MR Alivand, ...

Journal of cellular Physiology 233 (6), 4443-4457 2018

22. HDACis (class I), cancer stem cell, and phytochemicals: Cancer therapy and prevention implications
S Bayat, MS Khaniani, J Choupani, MR Alivand, SM Derakhshan

Biomedicine & Pharmacotherapy 97, 1445-1453 39 2018

23. Autistic phenotype of permutation and intermediate alleles of FMR1 gene
MS Khaniani, FA Yeganeh, S Amiri, SM Derakhshan

Iranian Journal of Pediatrics 27 (4) 2017

24. An update of common autosomal recessive non-syndromic hearing loss genes in Iranian population
T Ghasemnejad, MS Khaniani, F Zarei, M Farbodnia, SM Derakhshan

International journal of pediatric otorhinolaryngology 97, 113-126 2017

25. Restoration of correct splicing in IVSI-110 mutation of β-globin gene with antisense oligonucleotides: implications and applications in functional assay development
SM Derakhshan, MS Khaniani

Iranian Journal of Basic Medical Sciences 20 (6), 700 2017

26. Molecular analysis of glycogen storage disease type Ia in Iranian Azeri Turks: identification of a novel mutation

SK Mahmoud, A Khorrami, M Rafeey, R Ghergherehchi, MD Sima

Journal of genetics 96, 19-23 2017

27. Molecular analysis and prevalence of Huntington disease in northwestern Iran
MS Khaniani, P Aob, M Ranjouri, S DERAKHSHAN

Turkish Journal of Medical Sciences 47 (6), 1880-1884 2017

28. NUCB2/Nesfatin-1: a potent meal regulatory hormone and its role in diabetes
S Khalili, MS Khaniani, F Afkhami, SM Derakhshan

Egyptian Journal of Medical Human Genetics 18 (2), 105-109 2017

29. Screening for intermediate CGG alleles of FMR1 gene in male Iranian patients with Parkinsonism
A Entezari, MS Khaniani, T Bahrami, SM Derakhshan, H Darvish

Neurological Sciences 38, 123-128 2017

30. Genotyping of intron inversions and point mutations in exon 14 of the FVIII gene in Iranian Azeri Turkish families with hemophilia A
M Shekari Khaniani, A Ebrahimi, S Daraei, SM Derakhshan

Indian Journal of Hematology and Blood Transfusion 32, 475-480 2016

31. Extracts of Scrophularia frigida Boiss display potent antitumor effects in human breast cancer cells by inducing apoptosis and inhibition of expression of the human epidermal …
S Goldar, B Baradaran, MS Khaniani, A Azadmehr, SM Derakhshan, ...

Cellular and Molecular Biology 62 (9), 83-89 2016

32. Genetic Analysis of the ZNF512B, SLC41A1, and ALDH2 Polymorphisms in Parkinson's Disease in the Iranian Population
F Madadi, MS Khaniani, EE Shandiz, H Ayromlou, S Najmi, ...

Genetic testing and molecular biomarkers 20 (10), 629-632 2016

33. The Association of Nucleobindin 2 Gene (NUCB2) Variants with Type 2 Diabetes Mellitus Among Iranian Azeri-Turkish Population
MS Khaniani, N Aghamohammdzade, A Akbarzadeh, SM Derakhshan

Biosciences Biotechnology Research Asia 13 (3), 1821-1828 2016

34. Molecular study of deletional and nondeletional mutations on the α-globin locus in the Azeri population of Northwestern Iran
SM Derakhshan, MS Khaniani, F Afkhami, AH PourFeizi

Hemoglobin 40 (5), 319-322 2016

35. Cytogenetic findings in patients with intellectual disability and/or multiple congenital anomalies
S Mansoori Derakhshan, M Shekari Khaniani

Journal of Analytical Research in Clinical Medicine 4 (2), 97-103 2016

36. Evaluation of Thrombophilic genes in recurrent pregnancy loss: a case-control study in Iranian women
MS Khaniani, F Afkhami, F Abbasalizadeh, SM Derakhshan

International Journal of Human Genetics 16 (1-2), 48-52 2016

37. Sodium Butyrate and Valproic Acid as Splicing Restoring Agents in Erythroid Cells of β-Thalassemic Patients
MS Khaniani, M Tagizadeh, AH Feizi, SM Derakhshan

Iranian Journal of Biotechnology 14 (1), 9 2016

38. Association study of IL2RA and CTLA4 gene variants with Type I diabetes mellitus in children in the northwest of Iran
MR Ranjouri, P Aob, SM Derakhshan, MS Khaniani, H Chiti, A Ramazani

BioImpacts: BI 6 (4), 187 2016

39. Role of 14-bp HLA-G, INDEL polymorphism in recurrent miscarriage
F Afkhami, N Yazdani, MS Khaniani, SM Derakhshan

Global Journal of Health Science 8 (12), 1-45 2016

40. Evaluation of a newly discovered breast cancer susceptibility locus at 6q25. 1 in Iranian Azari-Turkish women
Z Garehdaghchi, SM Derakhshan, MS Khaniani

Contemporary Oncology/Współczesna Onkologia 20 (4), 308-310 2016

41. Detection of Hemophilia A Carriers in Azeri Turkish Population of Iran: Usefulness of HindIII and BclI markers
T Moharrami, SM Derakhshan, AAH Pourfeizi, MS Khaniani

Clinical and Applied Thrombosis/Hemostasis 21 (8), 755-759 2015

42. The association between human leukocyte antigen class II DR3–DQ2 haplotype and type 1 diabetes in children of the East Azerbaijan state of Iran
SM Derakhshan, FZ Sehrig, N Sohrabi, S Shiva, B Baradaran, ...

Iranian Red Crescent Medical Journal 17 (9) 2015

43. One novel frameshift mutation on exon 64 of COL7A1 gene in an Iranian individual suffering recessive dystrophic epidermolysis bullosa
MS Khaniani, N Sohrabi, NM Derakhshan, SM Derakhshan

Annals of Clinical & Laboratory Science 45 (5), 582-584 2015

44. Evaluation of association between HLA class II DR4–DQ8 haplotype and type I diabetes mellitus in children of East Azerbaijan state of Iran
N Sohrabi, MS Khaniani, SM Derakhshan

Advanced pharmaceutical bulletin 5 (1), 137 2015

45. Spectrum of β-globin gene mutations and β-thalassemia Haplotype Analysis among the Iranian Azeri Turkish population
S Derakhshan, A Khorrami, A Feizi, M Khaniani

Epidemiology (sunnyvale) 5 (210), 2161-1165.1000210 2015

46. Molecular mechanisms of apoptosis and roles in cancer development and treatment
S Goldar, MS Khaniani, SM Derakhshan, B Baradaran

Asian Pacific journal of cancer prevention 16 (6), 2129-2144 2015

47. Cloning of soluble human stem cell factor in pet-26b (+) vector
S Asghari, MS Khaniani, M Darabi, SM Derakhshan

Advanced Pharmaceutical Bulletin 4 (1), 91 2014

48. Minor role of BRCA2 mutation (Exon2 and Exon11) in patients with early-onset breast cancer amongst Iranian Azeri-Turkish women
NK Fathi, MS Khaniani, V Montazeri, SM Derakhshan

Iranian journal of basic medical sciences 17 (2), 108 2014

49. The HLA-G 14-bp insertion/deletion polymorphism in recurrent spontaneous abortion among Iranian women
F Afkhami, MS Khaniani, L Farzadi, Z Paknejad, SM Derakhshan

Iranian Journal of Allergy, Asthma and Immunology, 364-369 2014

50. Construction of pPIC9 Recombinant Vector Containing Human Stem Cell Factor
B Farhadi, MS Khaniani, SM Derakhshan

Advanced Pharmaceutical Bulletin 3 (2), 303 2013

51. Construction of yeast recombinant expression vector containing human epidermal growth factor (hEGF)
J Mohammadian, S Mansoori-Derakhshan, M Mohammadian, ...

Advanced Pharmaceutical Bulletin 3 (2), 473 2013

52. Molecular approaches to assessing the genetic risk of stroke
MS Khanian, SM Derakhshan, FA Yeganeh

6th national & first international Iranian stroke congress 2013

53. Evaluating the Genes causes to risk for common forms of stroke
N Sohrabi, Z Sohrabi, MS Khaniani, SM Derakhshan

6th national & first international Iranian stroke congress 2013

54. NPHS1 gene mutations in children with Nephrotic Syndrome in northwest Iran.
AG Behbahan, B Poorshiri, F Mortazavi, MS Khaniani, SM Derakhshan

Pakistan journal of biological sciences: PJBS 16 (17), 882-886 2013

55. Molecular Characterization and Assessment of Clinical Significance of Small Fragile X Alleles: Molecular and Clinical Significance of Small Fragile X Alleles
M Shekari Khaniani, S Mansoori Derakhshan

Journal of Analytical Research in Clinical Medicine 1 (1) 2013

56. Prenatal diagnosis of spinal muscular atrophy: clinical experience and molecular genetics of SMN gene analysis in 36 cases
MS Khaniani, SM Derakhshan, S Abasalizadeh

Journal of prenatal medicine 7 (3), 32 2013

57. Small MINOR ROLE FOR BRCA2 (EXON 2 AND EXON11) AMONG PATIENTS WITH EARLY ONSET BREAST CANCER IN NORTHWEST OF IRAN
FN KARIMIAN, KM SHEKARI, V MONTAZERI, DS MANSOORI, 2013

58. ANALYSIS OF HLA DR-DQ HAPLOTYPES AND GENETIC SUSCEPTIBILITY TO TYPE 1 DIABETES MELLITUS IN NORTHWEST OF IRAN
N SOHRABI, SF ZEINALI, KM SHEKARI, S SHIVA, B BARADARAN, 2013

59. Association between a new polymorphism (RS2046210) of the 6Q25. 1 locus and breast cancer risk in Iran
Z GHREHDAGHCHI, DS MANSOORI, M GHOJAZADEH, KM SHEKARI, 2013

60. Antisense Therapy for IVSI-110 ß-thalassaemia
SM Derakhshan

University of Melbourne, Department of Paediatrics 2008

61. Construction and Expression of a New Modified Coagulation FVIII cDNA In NIH3T3, CHO, and HepG2 Cell Lines
M Sima, KS Mahmoud, E Ammar, M Tamouchin

The Journal of Coagulation Disorders 2 (3), 1

62. A novel cryptic splice site in IVSI-110 B-thalassemia
SM Derakhshan, H Wardan, M Kleanthous, S Christou, J Vadolas

JOURNAL OF GENE MEDICINE 9 (6), 535-535 2007

63. The associations Between 5-HTTLPR Pplymorphism of SLC6A4 Gene and Autistic disorder in North West of IRAN
DS Mansoori, F Taheri, S Amiri, KL Shekari, KM Shekari

Medical Journal of Tabriz University of Medical Sciences 40 (5), 84-90

64. Informativeness of St14 VNTR Marker For Carrier Detection And Prenatal Diagnosis of Hemophilia A in Iranian Families
A Ebrahimi, SM Derakhshan

65. Humanised mouse model containting the common IVS1-110 splicing mutation
J Vadales, M nefedov, H wardan, S Mansoori derakhshan, L Voullaire, ...

The journal of Biological Chemistry 281 (11), 7399-7405

CONGRESS PRESENTATIONS: **A**bstract **P**ublished

1. 3th International and 18th Natinal congress of Iraniani Socity for Reporductive Medicine, Tabriz, Iran, Khaniani M, Derakhshan S, Taghizadeh S, Khoramei A, Mohrramei, T., 2012 April

2. First International congress of Molecular and cellular Medicine. Bahan 1389, Shiraz, Iran, Ebrahimi A, Shekari Khaniani M, Mansoori Derakhshan S, Moharramei T

3. M. S. Khaniani, S.M. Derakhshan, L Mohammadnejad. Controversy in the mode of inheritance in familial Mediterranean fever diseases, Molecular analysis of MEFV gene in patients. European Human Genetics conference, Nirenberg, Germany. 2012, June.

4. S.M. Derakhshan, M. S. Khaniani, Tagizadeh, Sodium Butyrate and Valproic acid as splicing restoring agents in erythroid cells of b-thalassemia patients. European Human Genetics conference, Nirenberg, Germany. 2012, June.

5. M. S. Khaniani, S.M. Derakhshan. To characterize and analyze the haplotype of Normal and "at risk of expansion" FMR1 CGG repeat alleles. European Human Genetics conference, Nirenberg, Germany. 2012, June.

6. Sima Mansoori Derakhshan & Jim Vadoles. 10th Iranian of Biochestry $ 3th international congress of Biochemistry and Molecular Biology, 2009, Nov

7. Sima Mansoori Derakhshan & Jim Vadoles. 10th Iranian of Biochestry $ 3th international congress of Biochemistry and Molecular Biology, 2009, Nov

8. M. Shekari Khaniani **, S Mansoori Derakhshan**. 6th International Congress of Multiple Sclerosis in Iran, Tabriz, Mehr 1388

9. M. S. Khaniani, **S.M. Derakhshan**, S. Tagezadeh, M, Zyadei, M, Nyosha. New aspect of fragile X mental retardation 1 (FMR1) gene. The Second Iranian Medical Genetics Congress. Khordad, 1390

10. M. S. Khaniani, **S.M. Derakhshan**. Inhibition of aberrant splicing of IVS-110 β-globin pre-mRNA by antisese oligonucleotide. The Second Iranian Medical Genetics Congress. Khordad, 1390

11. M. S. Khaniani**, S.M. Derakhshan**, L. Mohmmadnejade, M, Hajalilo, A. Mallekiyan, Y. Moaddabe. Molecular analysis of MEFVgene in Iranian patients with famimlial meditaranian fevere. The Second Iranian Medical Genetics Congress. Khordad, 1390

12. M. S. Khaniani**, S.M. Derakhshan** A. pourfeizi, A. Kazemei. An Improved diagnostic PCR assay for detection of α-thalassemia detection and α-triplication by multiplex PCR.The Second Iranian Medical Genetics Congress. Khordad, 1390

13. M. S. Khaniani**, S.M. Derakhshan,** A. pourfeizi A, Kazemei. Eritroid progenitor cells expanded from bone marrow: molecular characterisation and fuctional competence. The Second Iranian Medical Genetics Congress. Khordad, 1390

14. S Mansoori Derakhshan, M. ziadi, A. Hosseinpour Feizi, M. Shekari khaniani. Singapor. Molecular study of alpha thalassemia deletion and non deletional mutation frquency in alpha globin locus among potential carriers in North West of Iran. 2012 Apri

15. Nahid karimian fahi**,** mahmoud Shekari Khaniani, Vahid Motazeri, **Sima Mansoori derakhshan .** Minor role of BRCA2 mutation (Exon2 and Exon11) in patients with early-onset breast cancer amongst Iranian Azeri-Turkish women. 8th international congress of cancer**. Tehran, Esfand 1391**

16. M. S. Khaniani, S.M. Derakhshan, Z. Garadagchei, 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

17. S.Mansoori Derakhshan, Akbar Jalili, M. Shekari Khaniani, Sorayya Shiri torkamani, Soghra Taghizadeh. Frequency of G6PC gene mutations in GSD1a patients from North West of Iran. 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

18. M. Shekari. Khaniani, S.Mansoori Derakhshan, Akbar Jalili, Sorayya Shiri torkamani, Soghra Taghizadeh. GJB2 mutation spectrum in 450 Iranian Azeri Turkish patients with nonsyndromic hearing impairment. 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

19. M. Shekari. Khaniani, S.Mansoori Derakhshan, Akbar Jalili, Sorayya Shiri torkamani, Soghra Taghizadeh. The spectrum of mutations in the phenylalanine hydroxylase gene in patients with phenylketonuria North West Iran. 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

20. M. Shekari Khaniani S.Mansoori Derakhshan, Soghra Taghizadeh, A Jalili, Sorayya Shiri torkamani. β-globin Frameworks in β-Thalassemia carriers from Northwest of Iran 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

21. S.Mansoori Derakhshan, M. Shekari Khaniani, F. Zynali Sohrag. Evaluation of Association between HLA Class II DR3–DQ2 Haplotype and Type 1 Diabetes in Children of East Azerbaijan State of Iran. 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

22. S.Mansoori Derakhshan, M. Shekari Khaniani, S Daraii. β-globin Frameworks in β-Thalassemia carriers from Northwest of Iran. 3rd Iranian Medical Genetics Congress. Ordibehesht, 1391

23. Z. Garadagchei, S.Mansoori Derakhshan M. Ghojazadeh, M. Shekari. Khaniani. The 21th Iranian Congress of Physiology and Pharmacology, 23-27 August 2013

24. N. Sohrabei, Zeinali Sehrig, M. Shekari Khaniani, S. Shiva, B Baradaran, S.Mansoori Derakhshan. The 21th Iranian Congress of Physiology and Pharmacology, 23-27 August 2013

25. European Human Genetics conference: 2007/Nice/france

26. 11th international congress of Human Genetics: 2006/ Brisbane/ Australia

27. 10th International conference on Thalassaemia and Haemoglobinopathi and12th Intenernational TIF conference for thalassaemia Patients and Parents: 2006/ Dobai/ UAE

28. 5th Australian Gene therapy Society Meeting: 2007/ Melbourne/ Australia