CURRICULUM VITAE

Name: VAHID REZA **Surname: YASSAEE**

Date of birth: 1962
Nationality: Iranian

Academic Rank: Associate Professor
School: Medicine
Education Department: Medical Genetic

Research Department: Genomic Research Center

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Genomic Research Center
Taleghani Hospital, Aarabi St., Yaman Ave.,
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High Education:

1986 -1993 Doctorate Degree in Medicine
Faculty of Medicine, Iran University of Medical Sciences
Tehran – Iran

General Practitioner (GP) license number: 43537

1999- 2002 Doctorate Degree of Philosophy in Medical Genetics
(1) Dept. of Molecular Biology and Biotechnology
The University of Sheffield, Sheffield, S10 2TN, UK

(2) North Trent Molecular Genetics Laboratory
Sheffield Children's Hospital – (NHS)
Western Bank Sheffield - S10 2TH, UK

Medical Genetic Laboratory license number: A-1751

Awards:

1) A prize for Medical Doctorate thesis by faculty of Medicine, The Iran Medical Sciences University, Tehran-Iran **(1993)**
Titled: Epidemiological study on common skin disorder in eight provinces of Iran

2) A scholarship for a PhD course by faculty of Medicine, Shahid Beheshti University of Medical Sciences,
Tehran-Iran **(1998)**

Experiences:

1) General:

1991-1998

Medical advisory units for Mayor-Tehran Municipality
Head of statistical unit for medical affairs

2) Professional:

1993- 1996

Iranian Ministry of Health and Medical Education
Employed as a General Physician in Primary Health Care (PHC), Roudbar city - Iran

3) Technical:

1988-present

1) Designing and directing of a molecular genetics laboratory; experiences on molecular analysis of numerous genetic disorders achieved from four years' experience at Molecular Genetics Laboratory at National Health Services (NHS), Sheffield Children's Hospital, Sheffield-UK

2) Genome preparation; DNA, RNA, cDNA

3) Gel based mutation detection techniques: PTT, Multiplex PTT, SSCP, CSGE, DGGE

4) Genome manual analysis; PCR (ARMS, TETRA ARMS, RFLP, MS-PCR …), RT PCR, etc.; Agarose, PAGE, SDS-PAGE analysis

5) Genome automated analysis, Sanger sequencing, RealTime-PCR

6) Gene Assembling, an experience on BRCA1/2 genes analysis

7) Human genome profile (Exome) analysis and signing clinical report - derived from Next Generation Sequencing (NGS) technology

Positioned Titles:

September 2015 –Aug.2020

Center for Comprehensive Genetic Services (CCGS)
Director General
Taleghani Hospital, Aarabi St., Yaman Ave.,
Evin, Velenjak
Shahid Beheshti University of Medical Sciences
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May 2013 onward

National Center for Genomic Excellence (NCGE)
Director General
Taleghani Hospital, Aarabi St., Yaman Ave.,
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Shahid Beheshti University of Medical Sciences
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June 2007 –June 2021

Genomic Research Center (GRC)
Director General
Taleghani Hospital, Aarabi St., Yaman Ave.,
Evin, Velenjak
Shahid Beheshti University of Medical Sciences
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May 2006- Dec. 2008

Head of Medical Genetic Department
Faculty of Medicine
Shahid Beheshti University of Medical Sciences
Koodakyar St., Daneshjoo Blvd., Velenjak Ave., Eveen
Tehran-1985717443
P.O.Box : 19395-4719
Iran

June 2002 onward

Associate Professor of Medical Genetics
Dept. of Medical Genetic
Faculty of Medicine
Shahid Beheshti University of Medical Sciences
Koodakyar St.,Daneshjoo Blvd., Velenjak Ave., Evin
Tehran-1985717443
P.O.Box : 19395-4719
IRAN
Tel: +98 21 23872572

Director of E-Learning courses in collaboration with ESGM/ESHG

Place	Date	Description
Genomic Research Center, Shahid Beheshti University of Medical Sciences (GRC-SBMU)	May 2015	28th course in Medical Genetics
GRC-SBMU	May 2014	27th course in Medical Genetics
YAZD University of Medical Sciences	November 2006	7th course in Genetic Counseling in Practice
GRC-SBMU	May 2006	19th course in Medical Genetics
GRC-SBMU	November 2005	Molecular Cytogenetics and DNA Microarrays
GRC-SBMU	October 2005	Cancer Genetics
GRC-SBMU	May 2005	18th Medical Genetics

February 2007 : **Chairman of scientific committee for the 1st congress on IT development** in Iranian Medical Universities, 14th -15th Feb. 2007

Fields of interest:

1) Genome profiling of Iranian patients with **Breast Cancer**

2) **Molecular Epidemiology of Cancer** in IRAN (Breast Cancer, Prostate cancer, Bladder cancer)

3) Genome profiling of **Rare Genetic Disorders** with priority of children with metabolic disorders

4) **Public Health Genomics**

5) Genetic aspect of **Multifactorial Disorders** (Diabetes II, Coronary Artery Disease, Cancer)

Journal Publication (English papers)

https://scholar.google.com/citations?hl=en&user=0-rGlMcAAAAJ&view_op=list_works&sortby=pubdate

https://pubmed.ncbi.nlm.nih.gov/?term=yassaee+v&sort=pubdate

1) **Delineating the expanding phenotype of HERC2-related disorders: The impact of biallelic loss of function versus missense variation.**
Vincent KM, Eaton A, **Yassaee VR**, Miryounesi M, Hashemi-Gorji F, Rudichuk L, Goez H, Leonard N, Lazier
J.Clin Genet. **2021** Aug 9.
doi: 10.1111/cge.14039. Online ahead of PMID: 34370298

2) **Whole-Exome Sequencing Uncovers Novel Causative Variants and Additional Findings in Three Patients Affected by Glycogen Storage Disease Type VI and Fanconi-Bickel Syndrome.**
Eghbali M, Fatemi KS, Salehpour S, Abiri M, Saei H, Talebi S, Olyaei NA, **Yassaee VR**, Modarressi MH.
Front Genet. **2021** Jan 11;11:601566.
DOI: 10.3389/fgene.2020.601566. PMID: 33505429

3) **Novel long noncoding RNAs upregulation may have synergistic effects on the CYP24A1 and PFDN4 biomarker role in human colorectal cancer.**
Sadeghi H, Nazemalhosseini-Mojarad E, Sahebi U, Fazeli E, Azizi-Tabesh G, **Yassaee VR**, Savabkar S, Asadzadeh Aghdaei H, Zali MR, Mirfakhraie
R.J Cell Physiol**. 2021** Mar;236(3):2051-2057.
DOI: 10.1002/jcp.29992. PMID: 32743796

4) **A novel SRD5A2 mutation in an Iranian family with sex development disorder.**
Hashemi-Gorji F, Salehpour S, Miryounesi M, Mirfakhraie R, **Yassaee VR.**
Andrologia. **2021 Feb.** 53(1): e13847
DOI: 10.1111/and.13847. PMID: 33099786

5) **Association between single nucleotide polymorphisms rs12722489 and multiple sclerosis in Iranian patients with multiple sclerosis**
H Ahmadi, **VR Yassaee**, R Mirfakhraie, F Hashemi-Gorji
Current Journal of Neurology **2020**, 19 (1), 26-31

6) **Association of the P561T and C422F polymorphisms of the growth hormone receptor gene with facial dimensions.**
Dalaie, K., Behnaz, M., Banihashem, S., Motamedian, SR., **Yassaee, VR**. , Hashemi-Gorji, F., Khojasteh, A.
Journal of Oral Research, **2020**-8(6), pp. 499-504.
DOI: 10.17126/joralres.2019.073

7) **Could CYP24A1 promoter methylation status affect the gene expression in the colorectal cancer patients?**
Hossein Sadeghi, Ehsan Nazemalhosseini-Mojarad, **Vahid Reza Yassaee,** Sanaz Savabkar, Majid Ghasemian, Hamid Asadzadeh Aghdaei, Mohammad Reza Zali, Reza Mirfakhraie
Meta Gene, **2020**, Vol 24, 100656

8) **A candidate intronic *CYP24A1* gene variant affects the risk of colorectal cancer.**
Sadeghi H, Nazemalhosseini-Mojarad E, Piltan S, Fazeli E, Moradi Y, Amin-Beidokhti M, **Yassaee VR**, Aghdaei HA, Zali MR, Mirfakhraie R.
Biomark Med. **2020** Jan;14(1):23-29.
DOI: 10.2217/bmm-2019-0189. PMID: 31802707

9) **Relationship of the rs10850110 and rs11611277 polymorphisms of the MYO1H gene with non-syndromic mandibular prognathism in the Iranian population.**
Dalaie K, **Yassaee VR**, Behnaz M, Yazdanian M, Jafari F, Farimani RM.
Dent Med Probl. **2020** Oct-Dec;57(4):433-440.
DOI: 10.17219/dmp/122004.PMID: 33448167

10) **Mutations in the VPS13B Gene in Iranian Patients with Different Phenotypes of Cohen Syndrome.**
Alipour N, Salehpour S, Tonekaboni SH, Rostami M, Bahari S, **Yassaee V**, Miryounesi M, Ghafouri-Fard S. J Mol Neurosci. **2020** Jan;70(1):21-25.
DOI: 10.1007/s12031-019-01394-w. PMID: 31444703

11) **Next generation sequencing elucidated a clinically undiagnosed metabolic disorder - An Iranian family with hereditary orotic aciduria**
Zeinab Ravesh, **Vahid Reza Yassaee**, Seyed Hasan Tonekaboni, Maryam Razzaghy-Azar, Feyzollah Hashemi-Gorji, Shadab Salehpour, Mohammad Miryounesi, Soudeh Ghafouri-Fard.
Gene Reports, Volume 16, **2019**, 100457
DOI:10.1016/j.genrep.2019.100457

12) **Cloning, Expression, and Purification of Recombinant CEL I Endonuclease in HEK293T Cell Line**
Askari S., Hasannia S., Hassan Sajedi R., **Yassaee V.R.**
Modares Journal of Biotechnology. **2019**;10(3):473-481 ISSN: 2476-6917

13) **Investigation of CEBPA and CEBPA-AS Genes Expression in Acute Myeloid Leukemia**
Milad Gholami, Sahar Bayat, Saba Manoochehrabadi, Hossein Pashaiefar, Mir Davood Omrani, Hasan Jalaeikhoo, **Vahid Reza Yassaee**, Mohammad Reza Ebrahimpour, Farkhondeh Behjati, Reza Mirfakhraie.
Reports of Biochemistry and Molecular Biology, **2019**, 136-141. PMID:30805392

14) **Advanced molecular approaches pave the road to a clear-cut diagnosis of hereditary retinal dystrophies**
Ravesh Z, Dianatpour M, Fardaei M, Taghdiri M, Hashemi-Gorji F, **Yassaee VR**, Miryounesi M.
Mol Vis. **2018** Oct 19;24:679-689. PMID: 30416334

15) **An intron variant in the FLT1 gene increases the risk of preeclampsia in Iranian women**
Amin-Beidokhti M, Gholami M, Abedin-Do A, Pirjani R, Sadeghi H, Karamoddin F, **Yassaee VR,** Mirfakhraie R.
Clin Exp Hypertens. 2019;41(8):697-701.
doi: 10.1080/10641963.2018.1539097. Epub 2018 Nov 8. PMID: 30409050

16) **miR-30a promoter variation contributes to the increased risk of colorectal cancer in an Iranian population**
Sadeghi H, Nazemalhosseini-Mojarad E, Yaghoob-Taleghani M, Amin-Beidokhti M, **Yassaee VR**, Aghdaei HA, Zali MR, Mirfakhraie
R. J Cell Biochem. **2018** Nov 1.
PMID: 30387187, DOI: 10.1002/jcb.28047.

17) **Novel LAMA2 Gene Mutations Associated with Merosin-Deficient Congenital Muscular Dystrophy**
Hashemi-Gorji F, **Yassaee VR**, Dashti P, Miryounesi M.
Iran Biomed J. **2018**. PMID:29707938

18) **Neurodegeneration with brain iron accumulation 2A: Report of four independent cases**
M Miryounesi, S Salehpour, SH Tonekaboni, **VR Yassaee**, M Nejabat, F Hashemi-Gorji, Majid Fardaei, Soudeh Ghafouri-Fard.
Meta Gene 15, **2018**; 87–89.
DOI:10.1016/j.mgene.2017.12.006

19) **A new mutation in steroidogenic acute regulatory protein (StAR) is segregated in an Iranian family**.
S Ghafouri-Fard, **VR Yassaee**, N Alipour, Z Ravesh, M Miryounesi.
Meta Gene 16 ,**2018**; 196–198

20) **Whole exome sequencing unraveled the mystery of neurodevelopmental disorders in three Iranian families**
ZeinabRavesh, SoudehGhafouri-Fard, MasoumehRostami, NasrinAlipour, **Vahid Reza Yassaee**, Mohammad Miryounesi.
Gene Reports, Volume 13, **2018**, Pages 141-145;
DOI:10.1016/j.genrep.2018.10.002

21) **Genetic Diagnosis if a Lethal Form of Autosomal Recessive Polycystic Kidney Disease** S Mirzajani, M Mohebi, M Miryounesi, **VR Yassaee**, S Ghafouri-Fard.
International Journal of Pediatrics 6 (2), **2018;** 7033-7037

22) **Autosomal Recessive Hypohidrotic Ectodermal Dysplasia Caused by a Novel Mutation in EDAR Gene**
N Ebadi, S Javadi, TA Salmani, M Miryounesi, **VR Yassaee**, S Ghafouri-Frad.
International Journal of Pediatrics. **2018;** 6 (1), 6899-6902

23) **Clinical, biochemical and molecular features of Iranian families with mucopolysaccharidosis: A case series**
VR Yassaee, Hashemi-Gorji F, Miryounesi M, Rezayi A, Ravesh Z, Yassaee F, Salehpour S. Clin Chim Acta. **2017**.
DOI:10.1016/j.cca. PMID: 28844463

24) **Multidisciplinary management of a patient with van der Woude syndrome: A case report**
Tehranchi A, Behnia H, Nadjmi N, **VR Yassaee**, Ravesh Z, Mina M.
Int J Surg Case Rep. **2017**; 30:142-147.
DOI: 10.1016/j.ijscr.2016.11.032. PMID: 28012331

25) **Mutational Analysis of *FLT3* Internal Tandem Duplication and D835 in *De novo* Adult Acute Myeloid Leukemia**
Milad Gholami, Hossein Pashaiefar, Mohammad Reza Ebrahimpour, Sahar Bayat, Marzieh Hosseini, Ahmad Monabati, Mir Davood Omrani, **VR Yassaee**, Parvin Yavari, Farkhondeh Behjati and Reza Mirfakhraie. JAMMR.36370, **2017;** 24(5), 1-9

26) **Autosomal Recessive Hypohidrotic Ectodermal Dysplasia Caused by a Novel Mutation in EDAR Gene**
N Ebadi, S Javadi, TA Salmani, M Miryounesi, **VR Yassaee**.
International Journal of Pediatrics. **2017**; 6 (1), 6899-6902.

27) **A Novel Nonsense Mutation in *PANK2* Gene in Two Patients with Pantothenate Kinase-Associated Neurodegeneration**
Ghafouri-Fard S, **Yassaee VR**, Rezayi A, Hashemi-Gorji F, Alipour N, Miryounesi M.
Int J Mol Cell Med. **2016**; 5(4):255-259
PMID: 28357202; WOS: 000394500700006

28) **A Novel Splice Site Mutation in *HPS1* Gene is Associated with Hermansky-Pudlak Syndrome-1 (HPS1) in an Iranian Family**
Ghafouri-Fard S, Hashemi-Gorji F, **Yassaee VR**, Alipour N, Miryounesi M.
Int J Mol Cell Med. **2016**;5(3):192-195;
PMID: 27942505; WOS: 000391134900008

29) **Gene expression profiling of the 8q22-24 position in human breast cancer: *TSPYL5, MTDH, ATAD2*and *CCNE2* genes are implicated in oncogenesis, while *WISP1* and *EXT1* genes may predict a risk of metastasis.**
Taghavi A, Akbari ME, Hashemi-Bahremani M, Nafissi N, Khalilnezhad A, Poorhosseini SM, Hashemi-Gorji F, **Yassaee VR**.
Oncol Lett. 12(5), **2016**; 3845-3855;
PMID: 27895739; WOS: 000388838900118

30) **Novel FKBP10 Mutation in a Patient with Osteogenesis Imperfecta Type XI**
Seyedhassani SM, Hashemi-Gorji F, Yavari M, Harazi F, **Yassaee VR**.
Fetal Pediatr Pathol. **2016**; 35(5):353-358;
PMID:27362741; WOS: 000386453100009

31) **A novel splice site mutation in the GNPTAB gene in an Iranian patient with mucolipidosis II α/β**
Hashemi-Gorji F, Ghafouri-Fard S, Salehpour S, **Yassaee VR**, Miryounesi M.
J Pediatr Endocrinol Metab. **2016**; 1;29(8):991-3.
PMID: 271803, DOI: 10.1515/jpem-2016-0032. 37; WOS:000380756900017

32) **First Case Report of EX3del4765 Mutation in PAH Gene in Asian Population**
Soltani Z, Karami F, **Yassaee V**, Hashemi Gorji F, Talebzadeh M, Miryounesi M.
Iran Red Crescent Med J. **2016** Jan 1;18(2):e21633.
PMID: 27175306, DOI: 10.5812/ircmj.21633.

33) **New Gene Profiling in Determination of Breast Cancer Recurrence and Prognosis in Iranian Women**
Poorhosseini SM, Hashemi M, Alipour Olyaei N, Izadi A, Moslemi E, Ravesh Z, Hashemi-Gorji F, Kheiri HR, **Yassaee VR**.
Asian Pac J Cancer Prev. **2016**; 17(S3):155-60. PMID: 27165221

34) **Mutation Spectra of BRCA Genes in Iranian Women with Early Onset Breast Cancer - 15 Years Experience**
Yassaee VR, Ravesh Z, Soltani Z, Hashemi-Gorji F, Poorhosseini SM, Anbiaee R, Joulaee A. Asian Pac J Cancer Prev. **2016**;17(S3):149-53. PMID: 27165220

35) **Mutation spectra of the ITGB2 gene in Iranian families with leukocyte adhesion deficiency type 1**
Yassaee VR, Hashemi-Gorji F, Boosaliki S, Parvaneh N.
Hum Immunol. **2016** Feb;77(2):191-5.
PMID: 26639818, DOI: 10.1016/j.humimm.2015.11.019.; WOS: 000371191800007

36) **A novel homozygous LMNA mutation (p.Met540Ile) causes mandibuloacral dysplasia type A.**
Yassaee VR, Khojaste A, Hashemi-Gorji F, Ravesh Z, Toosi P.
Gene. **2016**; 10;577(1):8-13.
DOI: 10.1016/j.gene.2015.08.071; PMID: 26602028; WOS: 000368308600002

37) **Comparison of Insulin Expression Levels in White Blood Cells of infants with and without Family History of Type II Diabetes.**
Seyyed Reza Mazhari, Reza Mirfakhraie, Mojgan Asadi, Nasrin Alipour Olyaei, Hamidreza Kheiri, Elham Moslemi, Mahnaz Khanmohamadi, Elham Tohidnejad, **Vahid Reza Yassaee** NBM. Autumn **2016**; Vol.4 No.4

38) **Prognosticating Metastasis Risk in Early Breast Cancer with EXT1 and WISP1 Genes in 8q22-24 Position**.
Afsoon Taghavi, **Vahid Reza Yassaee,** Mohammad Esmaeil Akbari.
International Journal of Biology, Pharmacy and Allied Sciences(IJBPAS), **2016**, 5(8): 1996-2007

39) **A New Nonsense Mutation in CDKL5 Gene in a Male Patient with Early Onset Refractory Epilepsy.**
Soudeh Ghafouri-Fard, Shadab Salehpour, **Vahid Reza Yassaee**, Mohammad Miryounesi. a Case Report. IJP, Case Report. Vol. 4 (2) **2016** pp: 1315-1318

40) **A new approach for molecular diagnosis of TAR syndrome.**
Yassaee VR, Hashemi-Gorji F, Soltani Z, Poorhosseini SM.
Clin Biochem. **2014** Jun;47(9):835-9.
DOI: 10.1016/j.clinbiochem. 2014; 04.018.; PMID: 24769264; WOS: 000337992500106

41) **Association between SLC4A7 and COX11 variants and breast cancer in an Iranian population.**
VR Yassaee, Z Soltani, M Movahedi,
European journal of Cancer, Volume 50, Supplement 4, May **2014**, Pages e45,
DOI: 10.1016/j.ejca.2014.03.172; WOS: 000351589200126

42) **Cloning and Expression of Influenza H1N1 NS1 Protein in *Escherichia Coli* BL21.**
Marzieh Sadeghi; Mojgan Bandehpour; Fatemeh Yarian; **Vahidreza Yassaee**; Elham Torbati; Bahram Kazemi.
Iran J. Biotech. **2014** April; 12(1): e12625. DOI: 10.5812/ijb.12625

43) **Screening for genomic rearrangements at BRCA1 locus in Iranian women with breast cancer using multiplex ligation-dependent probe amplification.**
Yassaee VR, Emamalizadeh B, Omrani MD.
J Genet. **2013** Apr;92(1):131-4; PMID: 23640417; WOS: 000318868500016

44) **TAR Syndrome, a Rare Case Report with Cleft Lip/Palate.**
A. Naseh , A. Hafizi , F. Malek , H. Mozdarani , **V.R. Yassaee.**
The Internet Journal of Pediatrics and Neonatology. **2012** Vol.14 Number 1. DOI:10.5580/2c2a

45) **Mutation spectra of the AAAS gene in Iranian families with Allgrove Syndrome.**
Yassaee VR, Soltani Z, Ardakani BM.
Arch Med Res. **2011** Feb;42(2):163-8. doi: 10.1016/j.arcmed.2011.02.006. PMID: 21565631; WOS: 000290827100013

46) **BRCA1 and BRCA2 Genetic Testing in Breast and/or Ovarian Cancer Families in Iran.** F. Keshavarzi, G. R. Javadi, Nahid Nafissi, M. E. Akbari, **V.R. Yassaee**, M. Sharafi Farzad, S. Zeinali.
Yakhteh Medical Journal, Vol. 12, No 3, Autumn **2010**, Pages: 329-340; WOS: 000285526600003

47) **Ethical Issues in prevention of genetic diseases in Iran.**
Jamaldini, SH; **Yassaee, VR**; Ghaderian, SMH.
JOURNAL OF MEDICAL GENETICS, **2009**, Vol.: 46; WOS: 000270705500241

48) **No Association between Gadolinium-Based Contrast Agents and Development of Nephrogenic Systemic Fibrosis: a Case Study.**
B. Malekafzali A., S. Pirozi, **Vahid R. Yassaee.**
Iranian Journal of Dermatology, Autumn **2009**, Vol. 12, No 3(Suppl.)

49) **Comparison of CEL I gene expression and mismatch-cleavage activity in some Apiaceae plants.**
J. Zolala, A.R. Bahrami, M. Farsi, M. M. Matin, **Vahid R. Yassaee.**
Mol. Breeding, **2009**,
DOI: 10.1007/s11032-009-9267-x; WOS: 000267785700002

50) **Gene assembling, a new approach in molecular diagnosis of hereditary breast cancer. Vahid R. Yassaee,** A. Dalton
Medical Journal of Islamic Republic of Iran, **2007**, Vol.21, Issue.1; WOS: 000187166101115

51) **The New Genetically Mutations in the Breast Cancer's main Genes (BRCA1, BRCA2) in Iranian women affected by Unripe Cancer.**
Yassaee, V. R., A. Dalton, and D. P. Hornby.
Research Journal of Medical Science. **2006**: Vol. 28, p101-108.

52) **Novel mutations in the BRCA1 and BRCA2 genes in Iranian women with early-onset breast cancer.**
Yassaee VR, Zeinali S, Harirchi I, Jarvandi S, Mohagheghi MA, Hornby DP, Dalton A Breast Cancer Res. **2002**;4(4):R6;
PMID: 12100744; WOS: 000176674800001

GenBank accession number for Nucleotide (*search GenBank for Yassaee, V.R.)*
https://www.ncbi.nlm.nih.gov/nuccore?term=yassaee

Items: 252

Data has been attached as a separate profile

GenBank accession number for Protein (*search GenBank for Yassaee, V.R.)*
https://www.ncbi.nlm.nih.gov/protein/?term=yassaee

Items: 222

Data has been attached as a separate profile

Poster Presentations

1. Clinical, biochemical and molecular features of Iranian families with mucopolysaccharidosis: **Yassaee VR**, Hashemi-Gorji F, Miryounesi M, Rezayi A, Ravesh Z, Yassaee F, Salehpour S. International Congress of Inborn Errors of Metabolism (ICIEM) 2017, (poster #598) , 4th-8th Sep. **2017**, Brazil, Rio de Janeiro

2. Mutation spectra of *BRCA* genes in Iranian women with early onset breast cancer, 15 years experiences. **Yassaee V.R.**, Soltani Z., Ravesh Z., Hashemi-Gorgi F., Poorhosseini S.M., Anbiaee R., Joulaee A. (poster #2049F). The 65th Annual Meeting of The American Society of Human Genetics, 7th-10th Oct. **2015**, USA, Baltimore

3. Impact of reasonable genetic testing in prevention of rare genetic disorders. F. Hashemi-Gorji, **V.R. Yassaee**, A. Khojasteh, P. Toossi, Z. Ravesh. (poster #2049F). The 65th Annual Meeting of The American Society of Human Genetics, 7th-10th Oct. **2015**, USA, Baltimore

4. Novel FKBP10 mutation induces osteogenesis imperfecta type XI. S. Seyedhassani, M. Yavari1, F. Harazi1, F. Hashemi-Gorji, **V. Yassaee**. (poster #3015T). The 65th Annual Meeting of The American Society of Human Genetics, 7th-10th Oct. **2015**, USA, Baltimore

5. Association between SLC4A7 and COX11 variants and breast cancer in an Iranian population, **VR Yassaee**, Z Soltani, M Movahedi, European journal of Cancer, Volume 50, Supplement 4, May **2014**, Pages e45, DOI: 10.1016/j.ejca.2014.03.172,

6. A novel mutation in Iranian family with Phenylketonuria. Z. Soltani, M. Miryounesi, **V. R. Yassaee**. European Journal of Human Genetics, Volume 22 Supplement 1, May **2014**

7. A novel homozygote p.Met540Ile LMNA mutation causes mandibuloacral dysplasia type A. **V. R. Yassaee**, A. Khojasteh, F. Hashemi-Gorji, P. Toossi, S. M. Poorhosseini, S. R. Mazhari. European Journal of Human Genetics, Volume 22 Supplemenr 1, May **2014**

8. Association study of promoter polymorphisms of *nucb2* gene in Iranian patients with type 2 diabetes. S. Mosammami, **V. R. Yassaee**, Z. Soltani, K. Roohi Gilani. European Journal of Human Genetics, Volume 22 Supplemenr 1, May **2014**

9. Mutation spectra of the ITGB2 gene in Iranian families with Leukocyte Adhesion Deficiency type1. **V. R. Yassaee**, S. Boosaliki, F. Hashemi-Gorji, P. Dashti, N. Parvaneh*3*, M. Piryaei*2*;, European Journal of Human Genetics, Volume 22 Supplemenr 1, May **2014**

10. Quantitative real-time PCR technique for rapid diagnosis of TAR Syndrome. **V.R. Yassaee,** F. Hashemi-Gorji. ESHG congress. 8th -11th June **2013**, Paris, FRANCE

11. C/ebpbeta, e2f-1 and maz are predicted as a putative transcription factor for regulating rbm8a gene expression in TAR syndrome. **V.R. Yassaee**, F. Hashemi-Gorji, R. Mirfakhraie. HGM 13th -18th April **2013**, Singapore

12. Mutation Spectra of the AAAS Gene in Iranian Families with Allgrove Syndrome. **Vahid R. Yassaee**, Z. Soltani , B. Malekafzali A. ESHG congress, 28th-31th May **2011**, Amsterdam, The Netherlands

13. Association study of a new ARLTS1 polymorphism (Pro127Leu) in familial breast cancer. Fakhri M, **Yassaee V.R.**, Ohadi M, Karimloo M, Heshmati Y, Kamali K,*Khorram Khorshid HR.

The 8th International Congress on Obstetrics and Genecology, Nov.**2009**, Tehran-Iran

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