

In the Name of God
Isfahan University of Medical Sciences
CURRICULUM VITAE



Personal Information:

Name: Silva Hovsepian

Nationality: Iranian

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University Education:

- **M.D. (General Practitioner)**

2003, Isfahan University of Medical Sciences, Isfahan, Iran

- **PhD of Research in Clinical Sciences**

2018, Isfahan University of Medical Sciences, Isfahan, Iran

Work Experience:

* 19 Years of experience in professional fields (as a physician, tuberculosis and AIDS coordinator in Falavarjan, researcher and as a physician for childhood obesity)

* Publication Manager of Journal of Research in Medical Sciences since 2003

* Research Assistant at Imam Hossein Children's Hospital, Isfahan University of Medical Sciences, Isfahan, Iran

Research Interests:

Thyroid

Nutrition

Diabetes

Metabolic syndrome

Childhood obesity

Publications:

In Iranian Medical Journals

1. Shams B, Hashemipour M, Saadat SH, Emami SMH, Abde-Yazdan Z, Hassan-Zadeh A, Khatibi Kh, Haghghi S, Hovsepian S. Prevalence of helicobacter pylori infection in type 1 diabetic children referring to Isfahan Endocrine & Metabolism Research Center. *Iranian Journal of Diabetes and Lipid Disorders* Fall 2003- Winter 2004; 3(1): 35-40.
2. Hashemipour M, Amini M, Iranpour R, Javadi AA, Sadri GH, Javaheri N, Sattari G, Haghghi S, Hovsepian S. High prevalence of congenital hypothyroidism in Isfahan, Iran. *Iranian Journal of Endocrinology and Metabolism* 2004; 6(1): 13-20.
3. Hashemipour M, Javanmard GhA, Hourfar H, Kelishadi R, Hovsepian S, Haghghi S. Relationship between pancreas echogenicity and insulin sensitivity during the course of OGTT in children with β-thalassemia major aged 10-20 years. *Iranian Journal of Diabetes and Lipid Disorders* 2004; 3(2): 161-8.
4. Hashemipour M, Iranpour R, Amini M, Hovsepian S, Haghghi S. Comparison of the recall rate using two different diagnostic criteria at Isfahan Screening Program for Congenital Hypothyroidism. *Journal of Rafsanjan University of Medical Sciences* 2004; 3(3): 134-40.
5. Hashemipour M, Taghavi A, Mosaiiebi Z, Karimi Dana M, Amini M, Iranpour R, Khatibi Kh, Hovsepian S, Haghghi S. Screening for congenital hypothyroidism in Kashan, Iran. *Journal of Mazandaran University of Medical Sciences* 2005, 15(45): 83-91.
6. Momenzadeh M, Amini M, Aminorroaya A, Hovsepian S, Haghghi S. The prevalence of antithyroperoxidase (TPO-Ab) and Antithyroglobuline (Tg-Ab) autoantibodies in healthy women and female patients with hyperthyroidism, hypothyroidism and simple goiter: a comparative study. *Iranian Journal of Endocrinology & Metabolism* 2005; 6(4): 283-90.
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8. Hashemipour M, Iranpour R, Amini M, Hovsepian S, Haghghi S, Ahmadi N. The prevalence of consanguineous marriages in parents of neonates with congenital hypothyroidism: the Isfahan Screening Program for Neonatal Hypothyroidism. *Urmia Medical Journal* 2005; 16(2): 104-8.
9. Kalantari F, Hovsepian S, Haghghi S, Amini M. The prevalence of cardiovascular risk factors in patients with type 1 diabetes in Isfahan, Iran. *Iranian Journal of Diabetes and Lipid Disorders* 2007; 6(3): 255-62.

10. Salek M, Hashemipour M, Hashemi M, Hajrahimi M, Sadeghi S, Farajzadegan Z, Hovsepian S, Hadian R. Does the Prevalence of Hearing Impairment Differ in Children with Congenital Hypothyroidism and Healthy Children in Isfahan? *JIMS* 2007;25(87):53-39
11. Esfandiary E, Rabiei AA, Sobhan Ardekani A, Raeisifar M, Ahmadian Moghaddam S, Ramezanladeh Yazdi M, Mousavi SM, Mahdinezhad Gorgi H, Haghghi S, Hovsepian S, Amini M. The prevalence of thyroid pathological lesions in 202 autopsic samples obtained at Isfahan Forensic Medicine Center. *Iranian Journal of Endocrinology & Metabolism* 2008; 10(3):211-8.
12. Nasri P, Hashemipour M, Hovsepian S, Shahkarami M, Mehrabi A, Hadian R, Amini M. Usefulness of Ultrasonography in The Diagnosis of The Etiology of Congenital Hypothyroidism Comparing with Radioisotope Scanning. *Iranian Journal of Endocrinology and Metabolism* 2009;10(6):615-21.
13. Nasri P, Hashemipour M, Hovsepian S, Amini M, Heidari K, Sajjadi A, Ajami A, Movahedian Attar H, Dastanpoor M, Hadian R, Mohhebat. Comparison of Urine and Milk Iodine Concentration among Congenitally Hypothyroid Neonates and Their Mothers and a Control Group. *Iranian Journal of Endocrinology and Metabolism* 2008;11(3):265-272.
14. Hasani N, Dehghan B, Amini M, Heidari K, Sajjadi A, Ajami A, Dastanpoor M, Hadian R, Aminoroaya A, Pournaghshband Z, Hovsepian S, Hashemipour M. Congenital Hypothyroidism; is There any Familial Component? *JIMS* 2009;27(94):135-142
15. Rostampour N, Tajaddini M, Hashemipour M, Salehi M, Feizi A, Haghjooy Javanmard SH, Kelishadi R, Saneian H, Hovsepian S, Amini M. The Mutation of Dual Oxidase 2 (DUOX2) Gene among Patients with Permanent and Transient Congenital Hypothyroidism. *JIMS* 2011;29(139):588-599.
16. Feizi A, Hashemipour M, Hovsepian S, Amirkhani Z, Klishadi R, Rafiee Al Hosseini M et al. Study of the Efficacy of Therapeutic Interventions in Growth Normalization of Children with Congenital Hypothyroidism Detected By Neonatal Screening. *Iranian Journal of Endocrinology and Metabolism*. 2011; 13 (6) :681-689
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18. Mahjoubi F, Hashemipour M, Iranpour R, Amini M, Hovsepian S. TTF2 Gene Mutation in Neonates with Congenital Hypothyroidism Caused by Thyroid Dysgenesis. *JIMS* 2012;30 (189):657-666.
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20. Feizi A, Hashemipour M, Hovsepian S, Amirkhani Z, Kelishadi R, Rafee Al Hosseini M, Amini M. Study of the efficacy of therapeutic interventions in growth normalization of children with congenital hypothyroidism detected by neonatal screening. *IJEM* 2012;13(6 suppl):681-89.
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In Peer-Reviewed International Journals

1. Hashemipour M, Amini M, Iranpour R, Sadri GhH, Javaheri N, Haghghi S, Hovsepian S, Javadi AA, Nematbakhsh M, Sattari G. Prevalence of congenital hypothyroidism in Isfahan, Iran: results of a survey on 20000 neonates. *Hormone Research* 2004; 62: 79-83.
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10. Aminorroaya A, Amini M, Hovsepian S. PREVALENCE OF HYPERTHYROIDISM IN ISFAHAN-IRAN, IN THE YEAR 2006, FIFTEEN YEARS AFTER UNIVERSAL SALT IODIZATION: A COMMUNITY BASED STUDY. *Acta Endocrinologica (Buc)*, 2008;4(3): 273-285.
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29. Mostofizadeh N, Nikpour P, Javanmard SH, Emadi-Baygi M, Miranzadeh-Mahabadi H, Hovsepian S, Hashemipour M. The G395R Mutation of the Sodium/Iodide Symporter (NIS) Gene in Patients with Dyshormonogenetic Congenital Hypothyroidism.Int J Prev Med. 2013 Jan;4(1):57-62.
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