Asghar Aghamohammadi

List of Publications by Year in Descending Order

284 papers	6,302 citations	41 h-index	65 g-index
296 ext. papers	7,530 ext. citations	3.9 avg, IF	5.51 L-index

#	Paper	IF	Citations
284	Comprehensive Assessment of Skin Disorders in Patients with Common Variable Immunodeficiency´(CVID).. *Journal of Clinical Immunology*, **2022**, 1	5.7	0
283	Immunophenotypic and functional analysis of lymphocyte subsets in common variable immunodeficiency patients without monogenic defects.. *Scandinavian Journal of Immunology*, **2022**, e13164	3.4	2
282	Evaluation of Radiation Sensitivity in Patients with Hyper IgM Syndrome. *Immunological Investigations*, **2021**, 50, 580-596	2.9	2
281	Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature.. *Frontiers in Immunology*, **2021**, 12, 779502	8.4	2
280	T Cell Repertoire Abnormality in Immunodeficiency Patients with DNA Repair and Methylation Defects. *Journal of Clinical Immunology*, **2021**, 1	5.7	1
279	Genetic Risk Variants for Class Switching Recombination Defects in Ataxia-Telangiectasia Patients. *Journal of Clinical Immunology*, **2021**, 1	5.7	
278	Country Quarantine During COVID-19: Critical or Not?. *Disaster Medicine and Public Health Preparedness*, **2021**, 15, e24-e25	2.8	12
277	The spectrum of ATM gene mutations in Iranian patients with ataxia-telangiectasia. *Pediatric Allergy and Immunology*, **2021**, 32, 1316-1326	4.2	5
276	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. *Journal of Allergy and Clinical Immunology*, **2021**, 148, 1332-1341.e5	11.5	13
275	Application of Flow Cytometry in Predominantly Antibody Deficiencies. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2021**, 21, 647-663	2.2	0
274	Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. *Journal of Clinical Immunology*, **2021**, 41, 1339-1351	5.7	6
273	Clinical, immunological, and genetic features in 780 patients with autoimmune lymphoproliferative syndrome (ALPS) and ALPS-like diseases: A systematic review. *Pediatric Allergy and Immunology*, **2021**, 32, 1519-1532	4.2	7
272	Clinical, Immunologic and Molecular Spectrum of Patients with Immunodeficiency, Centromeric Instability, and Facial Anomalies (ICF) Syndrome: A Systematic Review. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2021**, 21, 664-672	2.2	3
271	International retrospective study of allogeneic hematopoietic cell transplantation for activated PI3K-delta syndrome. *Journal of Allergy and Clinical Immunology*, **2021**,	11.5	9
270	Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. *Pediatric Allergy and Immunology*, **2021**, 32, 1335-1348	4.2	5
269	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. *Immunological Investigations*, **2021**, 50, 201-215	2.9	18
268	Clinical, immunological and genetic findings in patients with UNC13D deficiency (FHL3): A systematic review. *Pediatric Allergy and Immunology*, **2021**, 32, 186-197	4.2	4

267	Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. *Journal of Clinical Immunology*, **2021**, 41, 76-88	5.7	8
266	Clinical complications and their management in a child with ataxia-telangiectasia (A-T): A case report study. *Clinical Case Reports (discontinued)*, **2021**, 9, 556-559	0.7	1
265	The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase-δ (PI3K) Syndrome (APDS). *Immunological Investigations*, **2021**, 1-16	2.9	1
264	Primary Immunodeficiency and Thrombocytopenia. *International Reviews of Immunology*, **2021**, 1-43	4.6	0
263	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. *Journal of Clinical Investigation*, **2021**, 131,	15.9	12
262	Primary Immunodeficiency Diseases in Iran: Past, Present and Future. *Archives of Iranian Medicine*, **2021**, 24, 118-124	2.4	0
261	Evaluation of miR-210 expression in common variable immunodeficiency: patients with unsolved genetic defect. *Allergologia Et Immunopathologia*, **2021**, 49, 84-93	1.9	2
260	Lymphocytes subsets in correlation with clinical profile in CVID patients without monogenic defects. *Expert Review of Clinical Immunology*, **2021**, 17, 1041-1051	5.1	2
259	Known and potential molecules associated with altered B cell development leading to predominantly antibody deficiencies. *Pediatric Allergy and Immunology*, **2021**, 32, 1601-1615	4.2	4
258	Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. *Clinical Immunology*, **2021**, 230, 108826	9	0
257	Predominantly antibody deficiencies **2021**, 93-123		
256	Management of inborn errors of immunity **2021**, 345-361		
255	Evaluation of MicroRNA-125b-5p and Transcription Factors BLIMP1 and IRF4 Expression in Unsolved Common Variable Immunodeficiency Patients.. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 700-710	1.1	
254	Clinical, Immunological, and Genetic Features in 49 Patients With ZAP-70 Deficiency: A Systematic Review. *Frontiers in Immunology*, **2020**, 11, 831	8.4	12
253	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. *Blood*, **2020**, 136, 2638-2655	2.2	32
252	Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. *International Archives of Allergy and Immunology*, **2020**, 181, 706-714	2.7	8
251	A new case of congenital ficolin-3 deficiency with primary immunodeficiency. *Expert Review of Clinical Immunology*, **2020**, 16, 733-738	5.1	5
250	Histocompatibility Complex Status and Mendelian Randomization Analysis in Unsolved Antibody Deficiency. *Frontiers in Immunology*, **2020**, 11, 14	8.4	3

249	Evaluation of respiratory complications in patients with X-linked and autosomal recessive agammaglobulinemia. *Pediatric Allergy and Immunology*, **2020**, 31, 405-417	4.2	3
248	Primary Immunodeficiency Diseases and Bacillus Calmette-Guérin (BCG)-Vaccine-Derived Complications: A Systematic Review. *Journal of Allergy and Clinical Immunology: in Practice*, **2020**, 8, 1371-1386	5.4	27
247	Comprehensive assessment of respiratory complications in patients with common variable immunodeficiency. *Annals of Allergy, Asthma and Immunology*, **2020**, 124, 505-511.e3	3.2	6
246	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. *Journal of Allergy and Clinical Immunology*, **2020**, 146, 901-911	11.5	29
245	Inborn Errors of Immunity and Cancers **2020**, 545-583		
244	Infectious Complications Reporting in Common Variable Immunodeficiency: A Systematic Review and Meta-analysis. *Oman Medical Journal*, **2020**, 35, e157	1.4	2
243	Leishmaniasis and Autoimmunity in Patient with LPS-Responsive Beige-Like Anchor Protein (LRBA) Deficiency. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 479-484	2.2	0
242	Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 157-171	2.2	6
241	Agammaglobulinemia: Epidemiology, Pathogenesis, Clinical Phenotype, Diagnosis, Prognosis and Management. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 1434-1447	2.2	1
240	Dystonia in Ataxia Telangiectasia: A Case Report with Novel Mutations. *Oman Medical Journal*, **2020**, 35, e93	1.4	3
239	Are asthma and allergic diseases phenotypic markers for patients with common variable immunodeficiency?. *Annals of Allergy, Asthma and Immunology*, **2020**, 124, 636	3.2	1
238	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. *Journal of Allergy and Clinical Immunology*, **2020**, 145, 1452-1463	11.5	61
237	Bronchiectasis in common variable immunodeficiency: A systematic review and meta-analysis. *Pediatric Pulmonology*, **2020**, 55, 292-299	3.5	16
236	Cutaneous Granulomatosis and Class Switching Defect as a Presenting Sign in Ataxia-Telangiectasia: First Case from the National Iranian Registry and Review of the Literature. *Immunological Investigations*, **2020**, 49, 597-610	2.9	9
235	Expanding Clinical Phenotype and Novel Insights into the Pathogenesis of ICOS Deficiency. *Journal of Clinical Immunology*, **2020**, 40, 277-288	5.7	13
234	Protein Kinase C-Delta Defect in Autoimmune Lymphoproliferative Syndrome-Like Disease: First Case from the National Iranian Registry and Review of the Literature. *Immunological Investigations*, **2020**, 1-12	2.9	3
233	Serum sickness-like reactions in Iranian children: a registry-based study in a referral center. *Allergologia Et Immunopathologia*, **2020**, 48, 424-429	1.9	
232	Autoimmunity in common variable immunodeficiency: a systematic review and meta-analysis. *Expert Review of Clinical Immunology*, **2020**, 16, 1227-1235	5.1	6

231	Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. *Immunological Investigations*, **2020**, 1-14	2.9	1
230	Global systematic review of primary immunodeficiency registries. *Expert Review of Clinical Immunology*, **2020**, 16, 717-732	5.1	29
229	Ataxia-telangiectasia: epidemiology, pathogenesis, clinical phenotype, diagnosis, prognosis and management. *Expert Review of Clinical Immunology*, **2020**, 16, 859-871	5.1	12
228	Clinical, Immunological, and Genetic Features in Patients with Activated PI3K Syndrome (APDS): a Systematic Review. *Clinical Reviews in Allergy and Immunology*, **2020**, 59, 323-333	12.3	38
227	Newborn Screening for Presymptomatic Diagnosis of Complement and Phagocyte Deficiencies. *Frontiers in Immunology*, **2020**, 11, 455	8.4	12
226	Malignancy in common variable immunodeficiency: a systematic review and meta-analysis. *Expert Review of Clinical Immunology*, **2019**, 15, 1105-1113	5.1	26
225	Graft versus host disease and microchimerism in a patient. *Allergy, Asthma and Clinical Immunology*, **2019**, 15, 47	3.2	2
224	Ataxia-telangiectasia: A review of clinical features and molecular pathology. *Pediatric Allergy and Immunology*, **2019**, 30, 277-288	4.2	70
223	G2-lymphocyte chromosomal radiosensitivity in patients with LPS responsive beige-like anchor protein (LRBA) deficiency. *International Journal of Radiation Biology*, **2019**, 95, 680-690	2.9	5
222	Respiratory Complications in Patients with Hyper IgM Syndrome. *Journal of Clinical Immunology*, **2019**, 39, 557-568	5.7	3
221	Disturbed Transcription of TLRs, Negative Regulators and Cytokines Secretion among TLR4- and 9-Activated PBMCs of Agammaglobulinemic Patients. *Immunological Investigations*, **2019**, 48, 860-874	2.9	4
220	Clinical Manifestations, Immunological Characteristics and Genetic Analysis of Patients with Hyper-Immunoglobulin M Syndrome in Iran. *International Archives of Allergy and Immunology*, **2019**, 180, 52-63	3.7	2
219	Outcomes and Treatment Strategies for Autoimmunity and Hyperinflammation in Patients with RAG Deficiency. *Journal of Allergy and Clinical Immunology: in Practice*, **2019**, 7, 1970-1985.e4	5.4	41
218	Candidiasis associated with very early onset inflammatory bowel disease: First IL10RB deficient case from the National Iranian Registry and review of the literature. *Clinical Immunology*, **2019**, 205, 35-42	3	4
217	The Heterogeneous Pathogenesis of Selective Immunoglobulin A Deficiency. *International Archives of Allergy and Immunology*, **2019**, 179, 231-246	3.7	7
216	Comparison of clinical and immunological features and mortality in common variable immunodeficiency and agammaglobulinemia patients. *Immunology Letters*, **2019**, 210, 55-62	4.1	9
215	The First Purine Nucleoside Phosphorylase Deficiency Patient Resembling IgA Deficiency and a Review of the Literature. *Immunological Investigations*, **2019**, 48, 410-430	2.9	6
214	The first cohort of Iranian patients with hyper immunoglobulin E syndrome: A long-term follow-up and genetic analysis. *Pediatric Allergy and Immunology*, **2019**, 30, 469-478	4.2	9

213	Clinical implications of systematic phenotyping and exome sequencing in patients with primary antibody deficiency. *Genetics in Medicine*, **2019**, 21, 243-251	8.1	64
212	Potential role of regulatory B cells in immunological diseases. *Immunology Letters*, **2019**, 215, 48-59	4.1	6
211	Clinical, Immunologic, and Molecular Spectrum of Patients with LPS-Responsive Beige-Like Anchor Protein Deficiency: A Systematic Review. *Journal of Allergy and Clinical Immunology: in Practice*, **2019**, 7, 2379-2386.e5	5.4	55
210	Vaccine-Derived Poliovirus Infection among Patients with Primary Immunodeficiency and Effect of Patient Screening on Disease Outcomes, Iran. *Emerging Infectious Diseases*, **2019**, 25, 2005-2012	10.2	4
209	The evaluation of neutropenia in common variable immune deficiency patients. *Expert Review of Clinical Immunology*, **2019**, 15, 1225-1233	5.1	3
208	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. *Immunology Letters*, **2019**, 216, 70-78	4.1	10
207	IL-10 induces TGF-β secretion, TGF-β receptor II upregulation, and IgA secretion in B cells. *European Cytokine Network*, **2019**, 30, 107-113	3.3	9
206	Antagonistic Property of G2013 (β-L-Guluronic Acid) on Gene Expression of MyD88, Tollip, and NF-κB in HEK293 TLR2 and HEK293 TLR4. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2019**, 19, 144-149	2.2	1
205	PIK3R1 Mutation Associated with Hyper IgM (APDS2 Syndrome): A Case Report and Review of the Literature. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2019**, 19, 941-958	2.2	0
204	IgG4 subclass and gamma-glutamyl transferase in children with ulcerative colitis with primary sclerosing cholangitis and without sclerosing cholangitis. *Clinical and Experimental Hepatology*, **2019**, 5, 285-288	2.2	0
203	The hyper IgM syndromes: Epidemiology, pathogenesis, clinical manifestations, diagnosis and management. *Clinical Immunology*, **2019**, 198, 19-30	9	36
202	Comparison of Common Monogenic Defects in a Large Predominantly Antibody Deficiency Cohort. *Journal of Allergy and Clinical Immunology: in Practice*, **2019**, 7, 864-878.e9	5.4	29
201	Evaluation of the TLR negative regulatory network in CVID patients. *Genes and Immunity*, **2019**, 20, 198-206	4.4	4
200	Interleukin-1β and interleukin-6 in Common Variable Immunodeficiency and their association with subtypes of B cells and response to the Pneumovax-23 vaccine. *European Cytokine Network*, **2019**, 30, 123-129	3.3	
199	New insights into physiopathology of immunodeficiency-associated vaccine-derived poliovirus infection; systematic review of over 5 decades of data. *Vaccine*, **2018**, 36, 1711-1719	4.1	22
198	Polyautoimmunity in Patients with LPS-Responsive Beige-Like Anchor (LRBA) Deficiency. *Immunological Investigations*, **2018**, 47, 457-467	2.9	13
197	Inflammation, a significant player of Ataxia-Telangiectasia pathogenesis?. *Inflammation Research*, **2018**, 67, 559-570	7.2	26
196	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2018**, 141, 1450-1458	11.5	56

195	Disease Evolution and Response to Rapamycin in Activated Phosphoinositide 3-Kinase δ Syndrome: The European Society for Immunodeficiencies-Activated Phosphoinositide 3-Kinase δ Syndrome Registry. *Frontiers in Immunology*, **2018**, 9, 543	8.4	88
194	Efficacy of an Attachment-Based Intervention Model on Health Indices in Children with Chronic Disease and Their Mothers. *Administration and Policy in Mental Health and Mental Health Services Research*, **2018**, 45, 900-910	3.1	4
193	Monogenic polyautoimmunity in primary immunodeficiency diseases. *Autoimmunity Reviews*, **2018**, 17, 1028-1039	13.6	18
192	Individual Radiosensitivity Assessment of the Families of Patients by G2-Checkpoint Abrogation. *Sultan Qaboos University Medical Journal*, **2018**, 18, e440-e446	0.9	4
191	Immunomodulatory Effect of G2013 (L-Guluronic Acid) on the TLR2 and TLR4 in Human Mononuclear Cells. *Current Drug Discovery Technologies*, **2018**, 15, 123-131	1.5	14
190	The Clinical and Immunological Features of Patients with Primary Antibody Deficiencies. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2018**, 18, 537-545	2.2	8
189	Mannose-Binding Lectin Protein Deficiency Among Patients with Primary Immunodeficiency Disease Receiving IVIG Therapy. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2018**, 18, 175-183	2.2	
188	Toll-like receptors pathway in common variable immune deficiency (CVID) and X-linked agammaglobulinemia (XLA). *European Cytokine Network*, **2018**, 29, 153-158	3.3	1
187	Fourth Update on the Iranian National Registry of Primary Immunodeficiencies: Integration of Molecular Diagnosis. *Journal of Clinical Immunology*, **2018**, 38, 816-832	5.7	57
186	chromosomal radiosensitivity in patients with common variable immunodeficiency. *Central-European Journal of Immunology*, **2018**, 43, 155-161	1.6	9
185	The imbalance of circulating T helper subsets and regulatory T cells in patients with LRBA deficiency: Correlation with disease severity. *Journal of Cellular Physiology*, **2018**, 233, 8767-8777	7	14
184	Clearing Vaccine-Derived Poliovirus Infection Following Hematopoietic Stem Cell Transplantation: a Case Report and Review of Literature. *Journal of Clinical Immunology*, **2018**, 38, 610-616	5.7	4
183	The Profile of Toll-like Receptor 2 (TLR2), TLR4 and Their Cytosolic Downstream Signaling Pathway in Common Variable Immunodeficiency (CVID) Patients. *Iranian Journal of Allergy, Asthma and Immunology*, **2018**, 17, 188-200	1.1	3
182	Abnormality of regulatory T cells in common variable immunodeficiency. *Cellular Immunology*, **2017**, 315, 11-17	4.4	22
181	The clinical significance of complete class switching defect in Ataxia telangiectasia patients. *Expert Review of Clinical Immunology*, **2017**, 13, 499-505	5.1	19
180	Clinical, immunologic, molecular analyses and outcomes of iranian patients with LRBA deficiency: A longitudinal study. *Pediatric Allergy and Immunology*, **2017**, 28, 478-484	4.2	48
179	Agammaglobulinemia: comorbidities and long-term therapeutic risks. *Expert Opinion on Orphan Drugs*, **2017**, 5, 559-574	1.1	0
178	Measurement of Health-Related Quality of Life in Primary Antibody-Deficient Patients. *Immunological Investigations*, **2017**, 46, 329-340	2.9	12

177	Preference of Genetic Diagnosis of CXCR4 Mutation Compared with Clinical Diagnosis of WHIM Syndrome. *Journal of Clinical Immunology*, **2017**, 37, 282-286	5.7	8
176	Combined immunodeficiency and Epstein-Barr virus-induced B cell malignancy in humans with inherited CD70 deficiency. *Journal of Experimental Medicine*, **2017**, 214, 91-106	16.6	111
175	Predominantly Antibody Deficiencies **2017**, 183-244		2
174	Autoimmunity in a cohort of 471 patients with primary antibody deficiencies. *Expert Review of Clinical Immunology*, **2017**, 13, 1099-1106	5.1	28
173	Ataxia telangiectasia syndrome: moonlighting ATM. *Expert Review of Clinical Immunology*, **2017**, 13, 1155-1172	5.1	24
172	Cernunnos deficiency associated with BCG adenitis and autoimmunity: First case from the national Iranian registry and review of the literature. *Clinical Immunology*, **2017**, 183, 201-206	9	8
171	Novel Mutation of ZAP-70-related Combined Immunodeficiency: First Case from the National Iranian Registry and Review of the Literature. *Immunological Investigations*, **2017**, 46, 70-79	2.9	15
170	Autoimmunity in common variable immunodeficiency: epidemiology, pathophysiology and management. *Expert Review of Clinical Immunology*, **2017**, 13, 101-115	5.1	40
169	Long-term outcomes of 176 patients with X-linked hyper-IgM syndrome treated with or without hematopoietic cell transplantation. *Journal of Allergy and Clinical Immunology*, **2017**, 139, 1282-1292	11.5	88
168	Impaired Akt phosphorylation in B-cells of patients with common variable immunodeficiency. *Clinical Immunology*, **2017**, 175, 124-132	9	21
167	Evaluation of infectious and non-infectious complications in patients with primary immunodeficiency. *Central-European Journal of Immunology*, **2017**, 42, 336-341	1.6	22
166	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. *Frontiers in Immunology*, **2017**, 8, 685	8.4	37
165	Costs of Hospital Admission on Primary Immunodeficiency Diseases. *Iranian Journal of Public Health* , **2017**, 46, 342-350	0.7	2
164	A Review on Defects of Dendritic Cells in Common Variable Immunodeficiency. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2017**, 17, 100-113	2.2	5
163	Role of Apoptosis in the Pathogenesis of Common Variable Immunodeficiency (CVID). *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2017**, 17, 332-340	2.2	5
162	Comparison of Bone Mineral Density in Common Variable Immunodeficiency and X-Linked Agammaglobulinaemia Patients. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2017**, 17, 134-140	2.2	2
161	Developing Inference Model to Diagnosis of Primary Immunodeficiency Diseases in Prot *Acta Medica Iranica*, **2017**, 55, 280-281		
160	Monogenic mutations associated with IgA deficiency. *Expert Review of Clinical Immunology*, **2016**, 12, 1321-1335	5.1	23

159	Toward the stratification and personalization of common variable immunodeficiency treatment. *Expert Opinion on Orphan Drugs*, **2016**, 4, 823-830	1.1	
158	The role of toll-like receptors in B-cell development and immunopathogenesis of common variable immunodeficiency. *Expert Review of Clinical Immunology*, **2016**, 12, 195-207	5.1	13
157	Role of apoptosis in common variable immunodeficiency and selective immunoglobulin A deficiency. *Molecular Immunology*, **2016**, 71, 1-9	4.3	34
156	Cohort of Iranian Patients with Congenital Agammaglobulinemia: Mutation Analysis and Novel Gene Defects. *Expert Review of Clinical Immunology*, **2016**, 12, 479-86	5.1	18
155	Evaluation of Known Defective Signaling-Associated Molecules in Patients Who Primarily Diagnosed as Common Variable Immunodeficiency. *International Reviews of Immunology*, **2016**, 35, 7-24	4.6	17
154	Spectrum of Phenotypes Associated with Mutations in LRBA. *Journal of Clinical Immunology*, **2016**, 36, 33-45	5.7	134
153	Managing patients with side effects and adverse events to immunoglobulin therapy. *Expert Review of Clinical Pharmacology*, **2016**, 9, 91-102	3.8	14
152	Primary ciliary dyskinesia in six patients with bronchiectasis. *Pneumonologia I Alergologia Polska*, **2016**, 84, 109-15		
151	Lymphocytic Interstitial Pneumonitis: An Unusual Presentation of X-Linked Hyper Ig M Syndrome. *Iranian Journal of Pediatrics*, **2016**, 26, e3656	1	2
150	Vaccine-Derived Polioviruses and Children with Primary Immunodeficiency, Iran, 1995-2014. *Emerging Infectious Diseases*, **2016**, 22, 1712-9	10.2	18
149	The use of Immunoglobulin Therapy in Primary Immunodeficiency Diseases. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2016**, 16, 80-88	2.2	23
148	Autoimmunity in Primary Antibody Deficiencies. *International Archives of Allergy and Immunology*, **2016**, 171, 180-193	3.7	31
147	Autoimmunity in primary T-cell immunodeficiencies. *Expert Review of Clinical Immunology*, **2016**, 12, 989-1006	5.1	9
146	Cellular and molecular mechanisms of immune dysregulation and autoimmunity. *Cellular Immunology*, **2016**, 310, 14-26	4.4	33
145	LPS-Responsive Beige-Like Anchor Gene Mutation Associated With Possible Bronchiolitis Obliterans Organizing Pneumonia Associated With Hypogammaglobulinemia and Normal IgM Phenotype and Low Number of B Cells. *Acta Medica Iranica*, **2016**, 54, 620-623		12
144	Impact of IgE-mediated Food Allergy on Parental Quality of Life in Iranian Patients. *Iranian Journal of Allergy, Asthma and Immunology*, **2016**, 15, 372-380	1.1	1
143	Important Factors Influencing Severity of Common Variable Immunodeficiency. *Archives of Iranian Medicine*, **2016**, 19, 544-50	2.4	9
142	Antibiotic resistance in patients with primary immunodeficiency disorders versus immunocompetent patients. *Expert Review of Clinical Immunology*, **2015**, 11, 1163-72	5.1	5

141	RAC2 loss-of-function mutation in 2 siblings with characteristics of common variable immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2015**, 135, 1380-4.e1-5	11.5	71
140	Novel mutations in TNFRSF7/CD27: Clinical, immunologic, and genetic characterization of human CD27 deficiency. *Journal of Allergy and Clinical Immunology*, **2015**, 136, 703-712.e10	11.5	90
139	Economic burden of common variable immunodeficiency: annual cost of disease. *Expert Review of Clinical Immunology*, **2015**, 11, 681-8	5.1	25
138	Clinical phenotype classification for selective immunoglobulin A deficiency. *Expert Review of Clinical Immunology*, **2015**, 11, 1245-54	5.1	31
137	Different brands of intravenous immunoglobulin for primary immunodeficiencies: how to choose the best option for the patient?. *Expert Review of Clinical Immunology*, **2015**, 11, 1229-43	5.1	19
136	Adverse reactions to Mycobacterium bovis bacille Calmette-Guérin vaccination against tuberculosis in Iranian children. *Clinical and Experimental Vaccine Research*, **2015**, 4, 195-9	1.9	11
135	Class switch recombination process in ataxia telangiectasia patients with elevated serum levels of IgM. *Journal of Immunoassay and Immunochemistry*, **2015**, 36, 16-26	1.8	20
134	Primary immunodeficiencies: a decade of shifting paradigms, the current status and the emergence of cutting-edge therapies and diagnostics. *Expert Review of Clinical Immunology*, **2015**, 11, 117-39	5.1	7
133	Important differences in the diagnostic spectrum of primary immunodeficiency in adults versus children. *Expert Review of Clinical Immunology*, **2015**, 11, 289-302	5.1	15
132	Atopic Manifestations: Dermatitis, Allergic Rhinitis and Asthma in Patients With Hypogammaglobulinemia. *Iranian Journal of Pediatrics*, **2015**, 25, e2786	1	5
131	Primary Immunodeficiencies and Cancers **2015**, 343-375		
130	Vitamin D deficiency in chronic idiopathic urticaria. *Iranian Journal of Allergy, Asthma and Immunology*, **2015**, 14, 222-7	1.1	9
129	Autosomal recessive agammaglobulinemia: a novel non-sense mutation in CD79a. *Journal of Clinical Immunology*, **2014**, 34, 138-41	5.7	15
128	Cytokines in common variable immunodeficiency as signs of immune dysregulation and potential therapeutic targets - a review of the current knowledge. *Journal of Clinical Immunology*, **2014**, 34, 524-43	5.7	36
127	Ocular involvement in primary immunodeficiency diseases. *Journal of Clinical Immunology*, **2014**, 34, 23-38	5.7	5
126	Spectrum of bone marrow failures of myeloid series: new report of neutropenic patients from a referral pediatric center in Iran. *Pediatric Hematology and Oncology*, **2014**, 31, 109-16	1.7	0
125	Combined immunodeficiency presenting with vaccine-associated paralytic poliomyelitis: a case report and narrative review of literature. *Immunological Investigations*, **2014**, 43, 292-8	2.9	20
124	Immunoglobulin class switch recombination deficiency type 1 or CD40 ligand deficiency: from bedside to bench and back again. *Expert Review of Clinical Immunology*, **2014**, 10, 91-105	5.1	22

123	Primary immunodeficiency disorders in Iran: update and new insights from the third report of the national registry. *Journal of Clinical Immunology*, **2014**, 34, 478-90	5.7	82
122	A hypomorphic recombination-activating gene 1 (RAG1) mutation resulting in a phenotype resembling common variable immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2014**, 134, 1375-1380	11.5	72
121	Assessment of immune response following immunization with DTP/Td and MMR vaccines in children treated for acute lymphoblastic leukemia. *Pediatric Hematology and Oncology*, **2014**, 31, 656-63	1.7	9
120	Long-term evaluation of a historical cohort of Iranian common variable immunodeficiency patients. *Expert Review of Clinical Immunology*, **2014**, 10, 1405-17	5.1	16
119	Evaluation of antibody response to polysaccharide vaccine and switched memory B cells in pediatric patients with inflammatory bowel disease. *Gut and Liver*, **2014**, 8, 24-8	4.8	11
118	Alteration in frequency and function of CD4+CD25+FOXP3+ regulatory T cells in patients with immune thrombocytopenic purpura. *Iranian Journal of Allergy, Asthma and Immunology*, **2014**, 13, 85-92	1.1	22
117	Otological findings in pediatric patients with hypogammaglobulinemia. *Iranian Journal of Allergy, Asthma and Immunology*, **2014**, 13, 166-73	1.1	3
116	The persian version of the chronic urticaria quality of life questionnaire: factor analysis, validation, and initial clinical findings. *Iranian Journal of Allergy, Asthma and Immunology*, **2014**, 13, 278-85	1.1	10
115	Ataxia-telangiectasia or neurologic Wilson's disease: when strong family history becomes a diagnostic bias. *Acta Neurologica Belgica*, **2013**, 113, 195-6	1.5	2
114	Evaluation of class switch recombination in B lymphocytes of patients with common variable immunodeficiency. *Journal of Immunological Methods*, **2013**, 394, 94-9	2.5	16
113	Evaluation of CD4+CD25+FOXP3+ regulatory T cells function in patients with common variable immunodeficiency. *Cellular Immunology*, **2013**, 281, 129-33	4.4	42
112	A review on guidelines for management and treatment of common variable immunodeficiency. *Expert Review of Clinical Immunology*, **2013**, 9, 561-74; quiz 575	5.1	58
111	Evaluation of natural regulatory T cells in subjects with selective IgA deficiency: from senior idea to novel opportunities. *International Archives of Allergy and Immunology*, **2013**, 160, 208-14	3.7	26
110	Expression of activation-induced cytidine deaminase gene in B lymphocytes of patients with common variable immunodeficiency. *Iranian Journal of Pediatrics*, **2013**, 23, 451-7	1	7
109	Recurrent Gallbladder Hydrops and Sclerosing Cholangitis in 11-Year-Old Male with Hyper IgM Syndrome. *Iranian Journal of Pediatrics*, **2013**, 23, 705-6	1	1
108	Autism in a child with common variable immunodeficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 287-9	1.1	4
107	Family study of pediatric patients with primary antibody deficiencies. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 377-82	1.1	12
106	Evaluation of humoral immune function in patients with chronic idiopathic thrombocytopenic purpura. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 50-6	1.1	3

105	Bacillus Calmette-Guﬁn (BCG) complications associated with primary immunodeficiency diseases. *Journal of Infection*, **2012**, 64, 543-54	18.9	106
104	HLA-DRB1,-DQA1 and -DQB1 allele and haplotype frequencies in female patients with early onset breast cancer. *Pathology and Oncology Research*, **2012**, 18, 49-55	2.6	15
103	Primary immunodeficiency diseases associated with neurologic manifestations. *Journal of Clinical Immunology*, **2012**, 32, 1-24	5.7	31
102	Recurrent Infections **2012**, 1-75		
101	Combined T and B Cell Immunodeficiencies **2012**, 77-112		
100	Disorders of Phagocytic Cells **2012**, 193-231		
99	Immune Dysregulation Diseases **2012**, 233-277		
98	Novel mutation of the activation-induced cytidine deaminase gene in a Tajik family: special review on hyper-immunoglobulin M syndrome. *Expert Review of Clinical Immunology*, **2012**, 8, 539-46	5.1	23
97	Home-based subcutaneous immunoglobulin versus hospital-based intravenous immunoglobulin in treatment of primary antibody deficiencies: systematic review and meta analysis. *Journal of Clinical Immunology*, **2012**, 32, 1180-92	5.7	114
96	Predominantly Antibody Deficiency **2012**, 113-192		
95	Autoimmunity in X-linked agammaglobulinemia: Kawasaki disease and review of the literature. *Expert Review of Clinical Immunology*, **2012**, 8, 155-9	5.1	14
94	Evaluation of serum IgA levels in Iranian patients with type 1 diabetes mellitus. *Acta Diabetologica*, **2012**, 49, 131-5	3.9	15
93	Deleterious mutations in LRBA are associated with a syndrome of immune deficiency and autoimmunity. *American Journal of Human Genetics*, **2012**, 90, 986-1001	11	350
92	TNF-alpha single nucleotide polymorphisms in atopic dermatitis. *European Cytokine Network*, **2012**, 23, 163-5	3.3	16
91	The uncommon combination of common variable immunodeficiency, macrophage activation syndrome, and cytomegalovirus retinitis. *Viral Immunology*, **2012**, 25, 161-5	1.7	14
90	Debilitating progressive encephalitis in a patient with BTK deficiency. *Acta Microbiologica Et Immunologica Hungarica*, **2012**, 59, 335-42	1.8	5
89	Association of HLA-DRB1, DQA1 and DQB1 Alleles and Haplotypes with Common Variable Immunodeficiency in Iranian Patients. *Avicenna Journal of Medical Biotechnology*, **2012**, 4, 103-12	1.4	7
88	Physicians awareness on primary immunodeficiency disorders in Iran. *Iranian Journal of Allergy, Asthma and Immunology*, **2012**, 11, 57-64	1.1	14

87	The approach to children with recurrent infections. *Iranian Journal of Allergy, Asthma and Immunology*, **2012**, 11, 89-109	1.1	12
86	Impact of delayed diagnosis in children with primary antibody deficiencies. *Journal of Microbiology, Immunology and Infection*, **2011**, 44, 229-34	8.5	14
85	Primary immunodeficiency diseases associated with increased susceptibility to viral infections and malignancies. *Journal of Allergy and Clinical Immunology*, **2011**, 127, 1329-41.e2; quiz 1342-3	11.5	114
84	X-linked lymphoproliferative syndrome: a genetic condition typified by the triad of infection, immunodeficiency and lymphoma. *British Journal of Haematology*, **2011**, 152, 13-30	4.5	74
83	The demographics of primary immunodeficiency diseases across the unique ethnic groups in Iran, and approaches to diagnosis and treatment. *Annals of the New York Academy of Sciences*, **2011**, 1238, 24-32	6.5	14
82	Analysis of switched memory B cells in patients with IgA deficiency. *International Archives of Allergy and Immunology*, **2011**, 156, 462-8	3.7	41
81	Indications and safety of intravenous and subcutaneous immunoglobulin therapy. *Expert Review of Clinical Immunology*, **2011**, 7, 301-16	5.1	39
80	Health-related quality of life in primary antibody deficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2011**, 10, 47-51	1.1	27
79	Oral and dental health status in patients with primary antibody deficiencies. *Iranian Journal of Allergy, Asthma and Immunology*, **2011**, 10, 289-93	1.1	6
78	Comparison of pulmonary diseases in common variable immunodeficiency and X-linked agammaglobulinaemia. *Respirology*, **2010**, 15, 289-95	3.6	49
77	A single center 14 years study of infectious complications leading to hospitalization of patients with primary antibody deficiencies. *Brazilian Journal of Infectious Diseases*, **2010**, 14, 351-355	2.8	
76	Serum bactericidal antibody response 1 year after meningococcal polysaccharide vaccination of patients with common variable immunodeficiency. *Vaccine Journal*, **2010**, 17, 524-8		21
75	Vaccine-associated paralytic poliomyelitis in immunodeficient children, Iran, 1995-2008. *Emerging Infectious Diseases*, **2010**, 16, 1133-6	10.2	36
74	B-cell-T-cell activation and interaction in common variable immunodeficiency. *Human Immunology*, **2010**, 71, 355-62	2.3	19
73	Association of IL-4 and IL-10 gene promoter polymorphisms with common variable immunodeficiency. *Immunobiology*, **2010**, 215, 81-7	3.4	29
72	Evaluation of immunoglobulin levels and infection rate in patients with common variable immunodeficiency after immunoglobulin replacement therapy. *Journal of Microbiology, Immunology and Infection*, **2010**, 43, 11-7	8.5	8
71	A single center 14 years study of infectious complications leading to hospitalization of patients with primary antibody deficiencies. *Brazilian Journal of Infectious Diseases*, **2010**, 14, 351-355	2.8	10
70	Echocardiographic abnormalities and their correlation with bronchiectasis score in primary antibody deficiencies. *Journal of Cardiovascular Medicine*, **2010**, 11, 244-9	1.9	2

69	IgA deficiency and the MHC: assessment of relative risk and microheterogeneity within the HLA A1 B8, DR3 (8.1) haplotype. *Journal of Clinical Immunology*, **2010**, 30, 138-43	5.7	43
68	BAK, BAX, and NBK/BIK proapoptotic gene alterations in Iranian patients with ataxia telangiectasia. *Journal of Clinical Immunology*, **2010**, 30, 132-7	5.7	6
67	Characterization of 11 new cases of leukocyte adhesion deficiency type 1 with seven novel mutations in the ITGB2 gene. *Journal of Clinical Immunology*, **2010**, 30, 756-60	5.7	28
66	Behavior abnormality following intravenous immunoglobulin treatment in patients with primary antibody deficiencies. *Human Psychopharmacology*, **2010**, 25, 419-22	2.3	5
65	History of primary immunodeficiency diseases in iran. *Iranian Journal of Pediatrics*, **2010**, 20, 16-34	1	13
64	Cutaneous granulomas in common variable immunodeficiency: case report and review of literature. *Acta Dermatovenerologica Croatica*, **2010**, 18, 107-13	0.5	12
63	A single center 14 years study of infectious complications leading to hospitalization of patients with primary antibody deficiencies. *Brazilian Journal of Infectious Diseases*, **2010**, 14, 351-5	2.8	5
62	Antibody response to pneumococcal capsular polysaccharide vaccination in patients with chronic kidney disease. *European Cytokine Network*, **2009**, 20, 69-74	3.3	18
61	Evaluation of liver diseases in Iranian patients with primary antibody deficiencies. *Annals of Hepatology*, **2009**, 8, 196-202	3.1	3
60	Cytokine gene polymorphisms in common variable immunodeficiency. *International Archives of Allergy and Immunology*, **2009**, 150, 1-7	3.7	17
59	Mannose-binding lectin polymorphisms in common variable immunodeficiency. *Clinical and Experimental Medicine*, **2009**, 9, 285-90	4.9	6
58	IgA deficiency: correlation between clinical and immunological phenotypes. *Journal of Clinical Immunology*, **2009**, 29, 130-6	5.7	159
57	Clinical and laboratory findings in hyper-IgM syndrome with novel CD40L and AICDA mutations. *Journal of Clinical Immunology*, **2009**, 29, 769-76	5.7	42
56	Novel mutations in TACI (TNFRSF13B) causing common variable immunodeficiency. *Journal of Clinical Immunology*, **2009**, 29, 777-85	5.7	40
55	Neutropenia and primary immunodeficiency diseases. *International Reviews of Immunology*, **2009**, 28, 335-66	4.6	40
54	Neutropenia associated with X-linked Agammaglobulinemia in an Iranian referral center. *Iranian Journal of Allergy, Asthma and Immunology*, **2009**, 8, 43-7	1.1	13
53	Severe combined immunodeficiency: a cohort of 40 patients. *Pediatric Allergy and Immunology*, **2008**, 19, 303-6	4.2	37
52	Chromosomal radiosensitivity in patients with common variable immunodeficiency. *Immunobiology*, **2008**, 213, 447-54	3.4	46

51	Immunologic evaluation of patients with recurrent ear, nose, and throat infections. *American Journal of Otolaryngology - Head and Neck Medicine and Surgery*, **2008**, 29, 385-92	2.8	19
50	Severe congenital neutropenia or hyper-IgM syndrome? A novel mutation of CD40 ligand in a patient with severe neutropenia. *International Archives of Allergy and Immunology*, **2008**, 147, 255-9	3.7	16
49	Risk factors leading to hospital admission in Iranian asthmatic children. *International Archives of Allergy and Immunology*, **2008**, 145, 244-8	3.7	6
48	Progression of selective IgA deficiency to common variable immunodeficiency. *International Archives of Allergy and Immunology*, **2008**, 147, 87-92	3.7	102
47	Serum bactericidal antibody responses to meningococcal polysaccharide vaccination as a basis for clinical classification of common variable immunodeficiency. *Vaccine Journal*, **2008**, 15, 607-11		33
46	Genotype-phenotype correlation in Bruton's tyrosine kinase deficiency. *Journal of Pediatric Hematology/Oncology*, **2008**, 30, 679-83	1.2	18
45	Increased serum levels of soluble CD30 in patients with common variable immunodeficiency and its clinical implications. *Journal of Clinical Immunology*, **2008**, 28, 78-84	5.7	26
44	Onychomadesis in a patient with immunoglobulin class switch recombination deficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 41-4	1.1	5
43	Evaluation of humoral immune function in patients with bronchiectasis. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 69-77	1.1	12
42	Selective immunoglobulin A deficiency in Iranian blood donors: prevalence, laboratory and clinical findings. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 157-62	1.1	17
41	Presence of Idiopathic Thrombocytopenic Purpura and autoimmune hemolytic anemia in the patients with common variable immunodeficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 169-75	1.1	20
40	Human leukocyte antigens (HLA) associated with selective IgA deficiency in Iran and Sweden. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 209-14	1.1	14
39	Response to polysaccharide vaccination amongst pediatric patients with common variable immunodeficiency correlates with clinical disease. *Iranian Journal of Allergy, Asthma and Immunology*, **2008**, 7, 231-4	1.1	14
38	Gastrointestinal manifestations in patients with common variable immunodeficiency. *Digestive Diseases and Sciences*, **2007**, 52, 2977-83	4	96
37	Clinical and laboratory findings in Iranian patients with leukocyte adhesion deficiency (study of 15 cases). *Journal of Clinical Immunology*, **2007**, 27, 302-7	5.7	38
36	The clinical, immunohematological, and molecular study of Iranian patients with severe congenital neutropenia. *Journal of Clinical Immunology*, **2007**, 27, 525-33	5.7	35
35	Prevalence of asthma related to BMI in adolescents in Tehran, Iran, 2004-2005. *European Journal of Pediatrics*, **2007**, 166, 453-4	4.1	10
34	Hodgkin lymphoma in two siblings with common variable immunodeficiency. *Pediatric Hematology and Oncology*, **2007**, 24, 337-42	1.7	12

33	Serum bactericidal antibody response to serogroup C polysaccharide meningococcal vaccination in children with primary antibody deficiencies. *Vaccine*, **2007**, 25, 5308-14	4.1	21
32	Ataxia-telangiectasia in Iran: clinical and laboratory features of 104 patients. *Pediatric Neurology*, **2007**, 37, 21-8	2.9	54
31	Mortality and morbidity in common variable immunodeficiency. *Journal of Tropical Pediatrics*, **2007**, 53, 32-8	1.2	34
30	Successful management of neutropenia in a patient with CD40 ligand deficiency by immunoglobulin replacement therapy. *Iranian Journal of Allergy, Asthma and Immunology*, **2007**, 6, 37-40	1.1	10
29	High production of IL-18 by dendritic cells induced by sera from patients with primary antibody deficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2007**, 6, 59-65	1.1	10
28	CD40 ligand expression on stimulated T-helper lymphocytes in patients with common variable immunodeficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2007**, 6, 129-35	1.1	11
27	Bilateral basal ganglia involvement in a patient with Griscelli syndrome. *European Journal of Paediatric Neurology*, **2006**, 10, 207-9	3.8	12
26	Lymphoma of mucosa-associated lymphoid tissue in common variable immunodeficiency. *Leukemia and Lymphoma*, **2006**, 47, 343-6	1.9	52
25	The clinical and laboratory survey of Iranian patients with hyper-IgE syndrome. *Scandinavian Journal of Infectious Diseases*, **2006**, 38, 898-903		20
24	Clinical, immunological and molecular characteristics of 37 Iranian patients with X-linked agammaglobulinemia. *International Archives of Allergy and Immunology*, **2006**, 141, 408-14	3.7	47
23	Consanguinity in primary immunodeficiency disorders; the report from Iranian Primary Immunodeficiency Registry. *American Journal of Reproductive Immunology*, **2006**, 56, 145-51	3.8	76
22	Frequency and clinical manifestations of patients with primary immunodeficiency disorders in Iran: update from the Iranian Primary Immunodeficiency Registry. *Journal of Clinical Immunology*, **2006**, 26, 519-32	5.7	130
21	Overweight and obesity and their associated factors in adolescents in Tehran, Iran, 2004-2005. *European Journal of Pediatrics*, **2006**, 165, 489-93	4.1	35
20	Effect of regular intravenous immunoglobulin therapy on prevention of pneumonia in patients with common variable immunodeficiency. *Journal of Microbiology, Immunology and Infection*, **2006**, 39, 114-20	8.5	48
19	Health-related quality of life in primary immune deficient patients. *Iranian Journal of Allergy, Asthma and Immunology*, **2006**, 5, 23-7	1.1	18
18	Congenital neutropenia and primary immunodeficiency disorders: a survey of 26 Iranian patients. *Journal of Pediatric Hematology/Oncology*, **2005**, 27, 351-6	1.2	25
17	Clinical and immunological features of 65 Iranian patients with common variable immunodeficiency. *Vaccine Journal*, **2005**, 12, 825-32		86
16	Neutropenia in Iranian patients with primary immunodeficiency disorders. *Haematologica*, **2005**, 90, 554-6	6.6	16

15	Autoimmune lymphoproliferative syndrome: meticulous care for diagnosis. *Iranian Journal of Allergy, Asthma and Immunology*, **2005**, 4, 197	1.1	2
14	Chronic granulomatous disease: a clinical survey of 41 patients from the Iranian primary immunodeficiency registry. *International Archives of Allergy and Immunology*, **2004**, 134, 253-9	3.7	70
13	Blood pressure nomograms for school children in Iran. *Pediatric Nephrology*, **2004**, 19, 164-8	3.2	21
12	Efficacy of intravenous immunoglobulin on the prevention of pneumonia in patients with agammaglobulinemia. *FEMS Immunology and Medical Microbiology*, **2004**, 40, 113-8		51
11	X-linked agammaglobulinemia: a survey of 33 Iranian patients. *Immunological Investigations*, **2004**, 33, 81-93	2.9	43
10	Adverse reactions of prophylactic intravenous immunoglobulin infusions in Iranian patients with primary immunodeficiency. *Annals of Allergy, Asthma and Immunology*, **2004**, 92, 60-4	3.2	30
9	Mortality in primary immunodeficient patients, registered in Iranian primary immunodeficiency registry. *Iranian Journal of Allergy, Asthma and Immunology*, **2004**, 3, 31-6	1.1	7
8	Clinical and laboratory findings in Iranian children with cyclic neutropenia. *Iranian Journal of Allergy, Asthma and Immunology*, **2004**, 3, 37-40	1.1	9
7	Neutropenia in patients with primary antibody deficiency disorders. *Iranian Journal of Allergy, Asthma and Immunology*, **2004**, 3, 77-81	1.1	11
6	Gastrointestinal complications in two patients with common variable immunodeficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2004**, 3, 149-52	1.1	5
5	Screening of the Bruton Tyrosine Kinase (BTK) Gene Mutations in 13 Iranian Patients with Presumed X-Linked Agammaglobulinemia. *Iranian Journal of Allergy, Asthma and Immunology*, **2004**, 3, 175-9	1.1	5
4	Identification of an SH2D1A mutation in a hypogammaglobulinemic male patient with a diagnosis of common variable immunodeficiency. *International Journal of Hematology*, **2003**, 78, 45-7	2.3	28
3	Adverse effects of intravenous immunoglobulin therapy in patients with antibody deficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2003**, 2, 121-6	1.1	3
2	Frequency of Allergic Rhinitis in School-age Children (7-18 Years) in Tehran. *Iranian Journal of Allergy, Asthma and Immunology*, **2003**, 2, 181-4	1.1	7
1	Primary immunodeficiency in Iran: first report of the National Registry of PID in Children and Adults. *Journal of Clinical Immunology*, **2002**, 22, 375-80	5.7	70