# Hassan Abolhassani

## List of Publications by Citations



| | | | |
|---|---|---|---|
| 257 papers | 8,751 citations | 41 h-index | 88 g-index |
| 289 ext. papers | 16,012 ext. citations | 7.1 avg, IF | 5.86 L-index |

| # | Paper | IF | Citations |
|---|---|---|---|
| 257 | Global burden of 369 diseases and injuries in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet, The*, **2020**, 396, 1204-1222 | 40 | 1847 |
| 256 | Global burden of 87 risk factors in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet, The*, **2020**, 396, 1223-1249 | 40 | 1013 |
| 255 | The global, regional, and national burden of inflammatory bowel disease in 195 countries and territories, 1990-2017: a systematic analysis for the Global Burden of Disease Study 2017. *The Lancet Gastroenterology and Hepatology*, **2020**, 5, 17-30 | 18.8 | 448 |
| 254 | Global age-sex-specific fertility, mortality, healthy life expectancy (HALE), and population estimates in 204 countries and territories, 1950-2019: a comprehensive demographic analysis for the Global Burden of Disease Study 2019. *Lancet, The*, **2020**, 396, 1160-1203 | 40 | 228 |
| 253 | The global, regional, and national burden of stomach cancer in 195 countries, 1990-2017: a systematic analysis for the Global Burden of Disease study 2017. *The Lancet Gastroenterology and Hepatology*, **2020**, 5, 42-54 | 18.8 | 184 |
| 252 | IgA deficiency: correlation between clinical and immunological phenotypes. *Journal of Clinical Immunology*, **2009**, 29, 130-6 | 5.7 | 159 |
| 251 | Spectrum of Phenotypes Associated with Mutations in LRBA. *Journal of Clinical Immunology*, **2016**, 36, 33-45 | 5.7 | 134 |
| 250 | 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 |
| 249 | Five insights from the Global Burden of Disease Study 2019. *Lancet, The*, **2020**, 396, 1135-1159 | 40 | 113 |
| 248 | Measuring universal health coverage based on an index of effective coverage of health services in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet, The*, **2020**, 396, 1250-1284 | 40 | 112 |
| 247 | 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 |
| 246 | Different Aspects of Social Network Analysis **2006**, | | 101 |
| 245 | Selective IgA Deficiency: Epidemiology, Pathogenesis, Clinical Phenotype, Diagnosis, Prognosis and Management. *Scandinavian Journal of Immunology*, **2017**, 85, 3-12 | 3.4 | 95 |
| 244 | 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 |
| 243 | 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 |
| 242 | 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 |
| 241 | Tuberculosis and impaired IL-23-dependent IFN-γ immunity in humans homozygous for a common missense variant. *Science Immunology*, **2018**, 3, | 28 | 88 |

| | | | |
|---|---|---|---|
| 240 | 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 |
| 239 | Harmony K-means algorithm for document clustering. *Data Mining and Knowledge Discovery*, **2009**, 18, 370-391 | 5.6 | 81 |
| 238 | Persistence of SARS-CoV-2-specific B and T´cell responses in convalescent COVID-19 patients 6-8´months after the infection. *Med*, **2021**, 2, 281-295.e4 | 31.7 | 74 |
| 237 | 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 |
| 236 | 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 |
| 235 | The global, regional, and national burden of oesophageal cancer and its attributable risk factors in 195 countries and territories, 1990-2017: a systematic analysis for the Global Burden of Disease Study 2017. *The Lancet Gastroenterology and Hepatology*, **2020**, 5, 582-597 | 18.8 | 71 |
| 234 | Ataxia-telangiectasia: A review of clinical features and molecular pathology. *Pediatric Allergy and Immunology*, **2019**, 30, 277-288 | 4.2 | 70 |
| 233 | X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. *Science Immunology*, **2021**, 6, | 28 | 67 |
| 232 | Clinical implications of systematic phenotyping and exome sequencing in patients with primary antibody deficiency. *Genetics in Medicine*, **2019**, 21, 243-251 | 8.1 | 64 |
| 231 | 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 |
| 230 | 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 |
| 229 | 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 |
| 228 | Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2018**, 141, 1450-1458 | 11.5 | 56 |
| 227 | 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 |
| 226 | Cancer Incidence, Mortality, Years of Life Lost, Years Lived With Disability, and Disability-Adjusted Life Years for 29 Cancer Groups From 2010 to 2019: A Systematic Analysis for the Global Burden of Disease Study 2019.. *JAMA Oncology*, **2021**, | 13.4 | 51 |
| 225 | Impact of SARS-CoV-2 Pandemic on Patients with Primary Immunodeficiency. *Journal of Clinical Immunology*, **2021**, 41, 345-355 | 5.7 | 51 |
| 224 | Comparison of pulmonary diseases in common variable immunodeficiency and X-linked agammaglobulinaemia. *Respirology*, **2010**, 15, 289-95 | 3.6 | 49 |
| 223 | 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 |

| | | | |
|---|---|---|---|
| 222 | Current genetic landscape in common variable immune deficiency. *Blood*, **2020**, 135, 656-667 | 2.2 | 48 |
| 221 | Next Generation Sequencing Data Analysis in Primary Immunodeficiency Disorders - Future Directions. *Journal of Clinical Immunology*, **2016**, 36 Suppl 1, 68-75 | 5.7 | 46 |
| 220 | SARS-CoV-2-related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. *Journal of Experimental Medicine*, **2021**, 218, | 16.6 | 45 |
| 219 | Global, regional, and national progress towards Sustainable Development Goal 3.2 for neonatal and child health: all-cause and cause-specific mortality findings from the Global Burden of Disease Study 2019. *Lancet, The*, **2021**, 398, 870-905 | 40 | 43 |
| 218 | Evaluation of CD4+CD25+FOXP3+ regulatory T cells function in patients with common variable immunodeficiency. *Cellular Immunology*, **2013**, 281, 129-33 | 4.4 | 42 |
| 217 | 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 |
| 216 | 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 |
| 215 | Analysis of switched memory B cells in patients with IgA deficiency. *International Archives of Allergy and Immunology*, **2011**, 156, 462-8 | 3.7 | 41 |
| 214 | Autoimmunity in common variable immunodeficiency: epidemiology, pathophysiology and management. *Expert Review of Clinical Immunology*, **2017**, 13, 101-115 | 5.1 | 40 |
| 213 | Indications and safety of intravenous and subcutaneous immunoglobulin therapy. *Expert Review of Clinical Immunology*, **2011**, 7, 301-16 | 5.1 | 39 |
| 212 | 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 |
| 211 | Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. *Frontiers in Immunology*, **2017**, 8, 685 | 8.4 | 37 |
| 210 | Infectious and Noninfectious Pulmonary Complications in Patients With Primary Immunodeficiency Disorders. *Journal of Investigational Allergology and Clinical Immunology*, **2017**, 27, 213-224 | 2.3 | 36 |
| 209 | The hyper IgM syndromes: Epidemiology, pathogenesis, clinical manifestations, diagnosis and management. *Clinical Immunology*, **2019**, 198, 19-30 | 9 | 36 |
| 208 | Frequency and expression of inhibitory markers of CD4(+) CD25(+) FOXP3(+) regulatory T cells in patients with common variable immunodeficiency. *Scandinavian Journal of Immunology*, **2013**, 77, 405-12 | 3.4 | 35 |
| 207 | Role of apoptosis in common variable immunodeficiency and selective immunoglobulin A deficiency. *Molecular Immunology*, **2016**, 71, 1-9 | 4.3 | 34 |
| 206 | Cellular and molecular mechanisms of immune dysregulation and autoimmunity. *Cellular Immunology*, **2016**, 310, 14-26 | 4.4 | 33 |
| 205 | Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. *Blood*, **2020**, 136, 2638-2655 | 2.2 | 32 |

| | | | |
|---|---|---|---|
| 204 | Health sector spending and spending on HIV/AIDS, tuberculosis, and malaria, and development assistance for health: progress towards Sustainable Development Goal 3. *Lancet, The*, **2020**, 396, 693-724 | 40 | 32 |
| 203 | Clinical phenotype classification for selective immunoglobulin A deficiency. *Expert Review of Clinical Immunology*, **2015**, 11, 1245-54 | 5.1 | 31 |
| 202 | Autoimmunity in Primary Antibody Deficiencies. *International Archives of Allergy and Immunology*, **2016**, 171, 180-193 | 3.7 | 31 |
| 201 | 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 |
| 200 | Global systematic review of primary immunodeficiency registries. *Expert Review of Clinical Immunology*, **2020**, 16, 717-732 | 5.1 | 29 |
| 199 | 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 |
| 198 | Measuring routine childhood vaccination coverage in 204 countries and territories, 1980-2019: a systematic analysis for the Global Burden of Disease Study 2020, Release 1. *Lancet, The*, **2021**, 398, 503-521 | 40 | 29 |
| 197 | Autoimmunity in a cohort of 471 patients with primary antibody deficiencies. *Expert Review of Clinical Immunology*, **2017**, 13, 1099-1106 | 5.1 | 28 |
| 196 | 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 |
| 195 | Health-related quality of life in primary antibody deficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2011**, 10, 47-51 | 1.1 | 27 |
| 194 | Malignancy in common variable immunodeficiency: a systematic review and meta-analysis. *Expert Review of Clinical Immunology*, **2019**, 15, 1105-1113 | 5.1 | 26 |
| 193 | Inflammation, a significant player of Ataxia-Telangiectasia pathogenesis?. *Inflammation Research*, **2018**, 67, 559-570 | 7.2 | 26 |
| 192 | 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 |
| 191 | Economic burden of common variable immunodeficiency: annual cost of disease. *Expert Review of Clinical Immunology*, **2015**, 11, 681-8 | 5.1 | 25 |
| 190 | Ataxia telangiectasia syndrome: moonlighting ATM. *Expert Review of Clinical Immunology*, **2017**, 13, 1155-1172 | 5.1 | 24 |
| 189 | Comparison of various classifications for patients with common variable immunodeficiency (CVID) using measurement of B-cell subsets. *Allergologia Et Immunopathologia*, **2017**, 45, 183-192 | 1.9 | 24 |
| 188 | Clinical, Immunological, and Genetic Features in Patients with Immune Dysregulation, Polyendocrinopathy, Enteropathy, X-linked (IPEX) and IPEX-like Syndrome. *Journal of Allergy and Clinical Immunology: in Practice*, **2020**, 8, 2747-2760.e7 | 5.4 | 23 |
| 187 | Monogenic mutations associated with IgA deficiency. *Expert Review of Clinical Immunology*, **2016**, 12, 1321-1335 | 5.1 | 23 |

| | | | |
|---|---|---|---|
| 186 | Epidemiology and pathophysiology of malignancy in common variable immunodeficiency?. *Allergologia Et Immunopathologia*, **2017**, 45, 602-615 | 1.9 | 23 |
| 185 | The use of Immunoglobulin Therapy in Primary Immunodeficiency Diseases. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2016**, 16, 80-88 | 2.2 | 23 |
| 184 | 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 |
| 183 | Approach to the Management of Autoimmunity in Primary Immunodeficiency. *Scandinavian Journal of Immunology*, **2017**, 85, 13-29 | 3.4 | 22 |
| 182 | Evaluation of infectious and non-infectious complications in patients with primary immunodeficiency. *Central-European Journal of Immunology*, **2017**, 42, 336-341 | 1.6 | 22 |
| 181 | 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 |
| 180 | Impaired Akt phosphorylation in B-cells of patients with common variable immunodeficiency. *Clinical Immunology*, **2017**, 175, 124-132 | 9 | 21 |
| 179 | Mapping local patterns of childhood overweight and wasting in low- and middle-income countries between 2000 and 2017. *Nature Medicine*, **2020**, 26, 750-759 | 50.5 | 21 |
| 178 | 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 |
| 177 | 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 |
| 176 | Mapping routine measles vaccination in low- and middle-income countries. *Nature*, **2021**, 589, 415-419 | 50.4 | 20 |
| 175 | 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 |
| 174 | The clinical significance of complete class switching defect in Ataxia telangiectasia patients. *Expert Review of Clinical Immunology*, **2017**, 13, 499-505 | 5.1 | 19 |
| 173 | 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 |
| 172 | hypomorphic mutation: identification of a novel pathogenic mutation in exon 8 and a review of the literature. *Allergy, Asthma and Clinical Immunology*, **2019**, 15, 2 | 3.2 | 19 |
| 171 | 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 |
| 170 | Monogenic polyautoimmunity in primary immunodeficiency diseases. *Autoimmunity Reviews*, **2018**, 17, 1028-1039 | 13.6 | 18 |
| 169 | A survey of complementary and alternative medicine in Iran. *Chinese Journal of Integrative Medicine*, **2012**, 18, 409-16 | 2.9 | 18 |

| | | | |
|---|---|---|---|
| 168 | Vaccine-Derived Polioviruses and Children with Primary Immunodeficiency, Iran, 1995-2014. *Emerging Infectious Diseases*, **2016**, 22, 1712-9 | 10.2 | 18 |
| 167 | Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. *Immunological Investigations*, **2021**, 50, 201-215 | 2.9 | 18 |
| 166 | Autoimmunity and its association with regulatory T cells and B cell subsets in patients with common variable immunodeficiency. *Allergologia Et Immunopathologia*, **2018**, 46, 127-135 | 1.9 | 18 |
| 165 | Tracking development assistance for health and for COVID-19: a review of development assistance, government, out-of-pocket, and other private spending on health for 204 countries and territories, 1990-2050. *Lancet, The*, **2021**, 398, 1317-1343 | 40 | 18 |
| 164 | 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 |
| 163 | Asthma and Allergic Diseases in a Selected Group of Patients With Common Variable Immunodeficiency. *Journal of Investigational Allergology and Clinical Immunology*, **2016**, 26, 209-11 | 2.3 | 17 |
| 162 | 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 |
| 161 | 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 |
| 160 | Bronchiectasis in common variable immunodeficiency: A systematic review and meta-analysis. *Pediatric Pulmonology*, **2020**, 55, 292-299 | 3.5 | 16 |
| 159 | Predictive markers for humoral influenza vaccine response in patients with common variable immunodeficiency. *Journal of Allergy and Clinical Immunology*, **2018**, 142, 1922-1931.e2 | 11.5 | 15 |
| 158 | Autosomal recessive agammaglobulinemia: a novel non-sense mutation in CD79a. *Journal of Clinical Immunology*, **2014**, 34, 138-41 | 5.7 | 15 |
| 157 | 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 |
| 156 | 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 |
| 155 | Immunity to SARS-CoV-2 up to 15´months after infection.. *IScience*, **2022**, 25, 103743 | 6.1 | 15 |
| 154 | Global, regional, and national burden of respiratory tract cancers and associated risk factors from 1990 to 2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet Respiratory Medicine,the*, **2021**, 9, 1030-1049 | 35.1 | 15 |
| 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 | Molecular diagnosis of primary immunodeficiency diseases in a developing country: Iran as an example. *Expert Review of Clinical Immunology*, **2014**, 10, 385-96 | 5.1 | 14 |
| 151 | Autoimmunity in X-linked agammaglobulinemia: Kawasaki disease and review of the literature. *Expert Review of Clinical Immunology*, **2012**, 8, 155-9 | 5.1 | 14 |

| | | | |
|---|---|---|---|
| 150 | Impact of delayed diagnosis in children with primary antibody deficiencies. *Journal of Microbiology, Immunology and Infection*, **2011**, 44, 229-34 | 8.5 | 14 |
| 149 | The uncommon combination of common variable immunodeficiency, macrophage activation syndrome, and cytomegalovirus retinitis. *Viral Immunology*, **2012**, 25, 161-5 | 1.7 | 14 |
| 148 | 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 |
| 147 | Physicians awareness on primary immunodeficiency disorders in Iran. *Iranian Journal of Allergy, Asthma and Immunology*, **2012**, 11, 57-64 | 1.1 | 14 |
| 146 | Polyautoimmunity in Patients with LPS-Responsive Beige-Like Anchor (LRBA) Deficiency. *Immunological Investigations*, **2018**, 47, 457-467 | 2.9 | 13 |
| 145 | Investigation of underlying primary immunodeficiencies in patients with severe atopic dermatitis. *Allergologia Et Immunopathologia*, **2014**, 42, 336-41 | 1.9 | 13 |
| 144 | Expanding Clinical Phenotype and Novel Insights into the Pathogenesis of ICOS Deficiency. *Journal of Clinical Immunology*, **2020**, 40, 277-288 | 5.7 | 13 |
| 143 | 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 |
| 142 | Two Faces of LRBA Deficiency in Siblings: Hypogammaglobulinemia and Normal Immunoglobulin Levels. *Journal of Investigational Allergology and Clinical Immunology*, **2018**, 28, 48-50 | 2.3 | 13 |
| 141 | Measurement of Health-Related Quality of Life in Primary Antibody-Deficient Patients. *Immunological Investigations*, **2017**, 46, 329-340 | 2.9 | 12 |
| 140 | Clinical, Immunological, and Genetic Features in 49 Patients With ZAP-70 Deficiency: A Systematic Review. *Frontiers in Immunology*, **2020**, 11, 831 | 8.4 | 12 |
| 139 | Ataxia-telangiectasia: epidemiology, pathogenesis, clinical phenotype, diagnosis, prognosis and management. *Expert Review of Clinical Immunology*, **2020**, 16, 859-871 | 5.1 | 12 |
| 138 | Review of local herbal compounds found in the Iranian traditional medicine known to optimise male fertility. *Andrologia*, **2016**, 48, 850-9 | 2.4 | 12 |
| 137 | Newborn Screening for Presymptomatic Diagnosis of Complement and Phagocyte Deficiencies. *Frontiers in Immunology*, **2020**, 11, 455 | 8.4 | 12 |
| 136 | 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 |
| 135 | Cutaneous granulomas in common variable immunodeficiency: case report and review of literature. *Acta Dermatovenerologica Croatica*, **2010**, 18, 107-13 | 0.5 | 12 |
| 134 | Family study of pediatric patients with primary antibody deficiencies. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 377-82 | 1.1 | 12 |
| 133 | The approach to children with recurrent infections. *Iranian Journal of Allergy, Asthma and Immunology*, **2012**, 11, 89-109 | 1.1 | 12 |

| | | | |
|---|---|---|---|
| 132 | 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 |
| 131 | Mortality and morbidity in patients with X-linked agammaglobulinaemia. *Allergologia Et Immunopathologia*, **2015**, 43, 62-6 | 1.9 | 10 |
| 130 | Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. *Immunology Letters*, **2019**, 216, 70-78 | 4.1 | 10 |
| 129 | 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 |
| 128 | Anemia prevalence in women of reproductive age in low- and middle-income countries between 2000 and 2018. *Nature Medicine*, **2021**, 27, 1761-1782 | 50.5 | 10 |
| 127 | Comparison of clinical and immunological features and mortality in common variable immunodeficiency and agammaglobulinemia patients. *Immunology Letters*, **2019**, 210, 55-62 | 4.1 | 9 |
| 126 | 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 |
| 125 | IL-10 induces TGF-□secretion, TGF-□receptor II upregulation, and IgA secretion in B cells. *European Cytokine Network*, **2019**, 30, 107-113 | 3.3 | 9 |
| 124 | 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 |
| 123 | International retrospective study of allogeneic hematopoietic cell transplantation for activated PI3K-delta syndrome. *Journal of Allergy and Clinical Immunology*, **2021**, | 11.5 | 9 |
| 122 | Autoimmunity in primary T-cell immunodeficiencies. *Expert Review of Clinical Immunology*, **2016**, 12, 989-1006 | 5.1 | 9 |
| 121 | chromosomal radiosensitivity in patients with common variable immunodeficiency. *Central-European Journal of Immunology*, **2018**, 43, 155-161 | 1.6 | 9 |
| 120 | Important Factors Influencing Severity of Common Variable Immunodeficiency. *Archives of Iranian Medicine*, **2016**, 19, 544-50 | 2.4 | 9 |
| 119 | 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 |
| 118 | Monogenic Primary Immunodeficiency Disorder Associated with Common Variable Immunodeficiency and Autoimmunity. *International Archives of Allergy and Immunology*, **2020**, 181, 706-714 | 3.7 | 8 |
| 117 | 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 |
| 116 | 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 |
| 115 | Global, regional, and national mortality among young people aged 10-24 years, 1950-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet, The*, **2021**, 398, 1593-1618 | 40 | 8 |

| | | | |
|---|---|---|---|
| 114 | Variable Abnormalities in T and B Cell Subsets in Ataxia Telangiectasia. *Journal of Clinical Immunology*, **2021**, 41, 76-88 | 5.7 | 8 |
| 113 | Heterologous immunization with inactivated vaccine followed by mRNA-booster elicits strong immunity against SARS-CoV-2 Omicron variant.. *Nature Communications*, **2022**, 13, 2670 | 17.4 | 8 |
| 112 | The Heterogeneous Pathogenesis of Selective Immunoglobulin A Deficiency. *International Archives of Allergy and Immunology*, **2019**, 179, 231-246 | 3.7 | 7 |
| 111 | The probable role of cytomegalovirus in acute myocardial infarction. *Jundishapur Journal of Microbiology*, **2014**, 7, e9253 | 1.2 | 7 |
| 110 | X-Linked TLR7 Deficiency Underlies Critical COVID-19 Pneumonia in a Male Patient with Ataxia-Telangiectasia. *Journal of Clinical Immunology*, **2021**, 42, 1 | 5.7 | 7 |
| 109 | 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 |
| 108 | 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 |
| 107 | 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 |
| 106 | Challenges in investigating patients with isolated decreased serum IgM: The SIMcal study. *Scandinavian Journal of Immunology*, **2019**, 89, e12763 | 3.4 | 6 |
| 105 | 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 |
| 104 | Identification of a novel de novo gain-of-function mutation of PIK3CD in a patient with activated phosphoinositide 3-kinase syndrome. *Clinical Immunology*, **2018**, 197, 60-67 | 9 | 6 |
| 103 | A Comparison of Clinical and Immunologic Phenotypes in Familial and Sporadic Forms of Common Variable Immunodeficiency. *Scandinavian Journal of Immunology*, **2017**, 86, 239-247 | 3.4 | 6 |
| 102 | Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 157-171 | 2.2 | 6 |
| 101 | Autoimmunity in common variable immunodeficiency: a systematic review and meta-analysis. *Expert Review of Clinical Immunology*, **2020**, 16, 1227-1235 | 5.1 | 6 |
| 100 | Consensus Middle East and North Africa Registry on Inborn Errors of Immunity. *Journal of Clinical Immunology*, **2021**, 41, 1339-1351 | 5.7 | 6 |
| 99 | Global, regional, and national sex differences in the global burden of tuberculosis by HIV status, 1990-2019: results from the Global Burden of Disease Study 2019. *Lancet Infectious Diseases, The*, **2021**, | 25.5 | 6 |
| 98 | 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 |
| 97 | 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 |

| | | | |
|---|---|---|---|
| 96 | Compound Heterozygous Mutations of IL2-Inducible T cell Kinase in a Swedish Patient: the Importance of Early Genetic Diagnosis. *Journal of Clinical Immunology*, **2019**, 39, 131-134 | 5.7 | 5 |
| 95 | A new case of congenital ficolin-3 deficiency with primary immunodeficiency. *Expert Review of Clinical Immunology*, **2020**, 16, 733-738 | 5.1 | 5 |
| 94 | Circulating Helper T-Cell Subsets and Regulatory T Cells in Patients With Common Variable Immunodeficiency Without Known Monogenic Disease. *Journal of Investigational Allergology and Clinical Immunology*, **2018**, 28, 172-181 | 2.3 | 5 |
| 93 | IgG anti-IgA antibodies in paediatric antibody-deficient patients receiving intravenous immunoglobulin. *Allergologia Et Immunopathologia*, **2015**, 43, 403-8 | 1.9 | 5 |
| 92 | Circulating retinol-binding protein 4 concentrations in patients with coronary artery disease and patients with type 2 diabetes mellitus. *International Journal of Diabetes in Developing Countries*, **2012**, 32, 105-110 | 0.8 | 5 |
| 91 | Behavior abnormality following intravenous immunoglobulin treatment in patients with primary antibody deficiencies. *Human Psychopharmacology*, **2010**, 25, 419-22 | 2.3 | 5 |
| 90 | Heterologous immunization with inactivated vaccine followed by mRNA booster elicits strong humoral and cellular immune responses against the SARS-CoV-2 Omicron variant | | 5 |
| 89 | 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 |
| 88 | The spectrum of ATM gene mutations in Iranian patients with ataxia-telangiectasia. *Pediatric Allergy and Immunology*, **2021**, 32, 1316-1326 | 4.2 | 5 |
| 87 | Autoimmune manifestations among 461 patients with monogenic inborn errors of immunity. *Pediatric Allergy and Immunology*, **2021**, 32, 1335-1348 | 4.2 | 5 |
| 86 | The profile of IL-4, IL-5, IL-10 and GATA3 in patients with LRBA deficiency and CVID with no known monogenic disease: Association with disease severity. *Allergologia Et Immunopathologia*, **2019**, 47, 172-178 | 1.9 | 5 |
| 85 | 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 |
| 84 | Psychiatric aspects of primary immunodeficiency diseases: the parental study. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 176-81 | 1.1 | 5 |
| 83 | 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 |
| 82 | 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 |
| 81 | Inherited IFNAR1 Deficiency in a Child with Both Critical COVID-19 Pneumonia and Multisystem Inflammatory Syndrome.. *Journal of Clinical Immunology*, **2022**, 1 | 5.7 | 4 |
| 80 | Diabetes mortality and trends before 25 years of age: an analysis of the Global Burden of Disease Study 2019.. *Lancet Diabetes and Endocrinology,the*, **2022**, | 18.1 | 4 |
| 79 | The global burden of adolescent and young adult cancer in 2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet Oncology, The*, **2021**, | 21.7 | 4 |

| | | | |
|---|---|---|---|
| 78 | International Retrospective Study of Allogeneic Hematopoietic Cell Transplantation (HCT) for Activated Phosphoinositide 3-Kinase Delta (PI3K) Syndrome. *Biology of Blood and Marrow Transplantation*, **2020**, 26, S14-S15 | 4.7 | 4 |
| 77 | 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 |
| 76 | Targeted next-generation sequencing for genetic diagnosis of 160 patients with primary immunodeficiency in south China. *Pediatric Allergy and Immunology*, **2018**, 29, 863-872 | 4.2 | 4 |
| 75 | 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 |
| 74 | 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 |
| 73 | Autism in a child with common variable immunodeficiency. *Iranian Journal of Allergy, Asthma and Immunology*, **2013**, 12, 287-9 | 1.1 | 4 |
| 72 | Vitamin D Deficiency among Female Nurses of Children Medical Center Hospital and Its Related Factors. *Acta Medica Iranica*, **2016**, 54, 146-50 | | 4 |
| 71 | Burden of non-communicable diseases among adolescents aged 10-24 years in the EU, 1990-2019: a systematic analysis of the Global Burden of Diseases Study 2019.. *The Lancet Child and Adolescent Health*, **2022**, | 14.5 | 4 |
| 70 | Renal amyloidosis in common variable immunodeficiency. *Nefrologia*, **2010**, 30, 474-6 | 1.5 | 4 |
| 69 | Respiratory Complications in Patients with Hyper IgM Syndrome. *Journal of Clinical Immunology*, **2019**, 39, 557-568 | 5.7 | 3 |
| 68 | Histocompatibility Complex Status and Mendelian Randomization Analysis in Unsolved Antibody Deficiency. *Frontiers in Immunology*, **2020**, 11, 14 | 8.4 | 3 |
| 67 | Evaluation of respiratory complications in patients with X-linked and autosomal recessive agammaglobulinemia. *Pediatric Allergy and Immunology*, **2020**, 31, 405-417 | 4.2 | 3 |
| 66 | The evaluation of neutropenia in common variable immune deficiency patients. *Expert Review of Clinical Immunology*, **2019**, 15, 1225-1233 | 5.1 | 3 |
| 65 | Dystonia in Ataxia Telangiectasia: A Case Report with Novel Mutations. *Oman Medical Journal*, **2020** , 35, e93 | 1.4 | 3 |
| 64 | 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 |
| 63 | Specific Immune Response and Cytokine Production in CD70 Deficiency. *Frontiers in Pediatrics*, **2021** , 9, 615724 | 3.4 | 3 |
| 62 | 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 |
| 61 | Clinical, immunological, and genetic features in 938 patients with autoimmune polyendocrinopathy candidiasis ectodermal dystrophy (APECED): a systematic review. *Expert Review of Clinical Immunology*, **2021**, 17, 807-817 | 5.1 | 3 |

| | | | |
|---|---|---|---|
| 60 | Impaired respiratory burst contributes to infections in PKCδ deficient patients. *Journal of Experimental Medicine*, **2021**, 218, | 16.6 | 3 |
| 59 | Hallmarks of Cancers: Primary Antibody Deficiency Other Inborn Errors of Immunity. *Frontiers in Immunology*, **2021**, 12, 720025 | 8.4 | 3 |
| 58 | Otological findings in pediatric patients with hypogammaglobulinemia. *Iranian Journal of Allergy, Asthma and Immunology*, **2014**, 13, 166-73 | 1.1 | 3 |
| 57 | 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 |
| 56 | Graft versus host disease and microchimerism in a patient. *Allergy, Asthma and Clinical Immunology*, **2019**, 15, 47 | 3.2 | 2 |
| 55 | 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 |
| 54 | Evaluation of Radiation Sensitivity in Patients with Hyper IgM Syndrome. *Immunological Investigations*, **2021**, 50, 580-596 | 2.9 | 2 |
| 53 | Bilateral deep peroneal nerve paralysis following kerosene self-injection into external hemorrhoids. *Case Reports in Medicine*, **2010**, 2010, | 0.7 | 2 |
| 52 | Atypical Ataxia Presentation in Variant Ataxia Telangiectasia: Iranian Case-Series and Review of the Literature.. *Frontiers in Immunology*, **2021**, 12, 779502 | 8.4 | 2 |
| 51 | Infectious Complications Reporting in Common Variable Immunodeficiency: A Systematic Review and Meta-analysis. *Oman Medical Journal*, **2020**, 35, e157 | 1.4 | 2 |
| 50 | Costs of Hospital Admission on Primary Immunodeficiency Diseases. *Iranian Journal of Public Health* , **2017**, 46, 342-350 | 0.7 | 2 |
| 49 | Systematic investigation for underlying causes of recurrent infections in children: surveillance of primary immunodeficiency. *European Annals of Allergy and Clinical Immunology*, **2018**, 50, 72-80 | 1.3 | 2 |
| 48 | Persistence of SARS-CoV-2 specific B- and T-cell responses in convalescent COVID-19 patients 6-8 months after the infection | | 2 |
| 47 | Lymphocytic Interstitial Pneumonitis: An Unusual Presentation of X-Linked Hyper Ig M Syndrome. *Iranian Journal of Pediatrics*, **2016**, 26, e3656 | 1 | 2 |
| 46 | 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 |
| 45 | Evaluation of patients with primary immunodeficiency associated with Bacille Calmette-Guerin (BCG)-vaccine-derived complications. *Allergologia Et Immunopathologia*, **2020**, 48, 729-737 | 1.9 | 2 |
| 44 | 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 |
| 43 | Approach to genetic diagnosis of inborn errors of immunity through next-generation sequencing. *Molecular Immunology*, **2021**, 137, 57-66 | 4.3 | 2 |

| | | | |
|---|---|---|---|
| 42 | Ectopic decidual reaction mimicking irritable bowel syndrome: a case report. *Acta Medica Iranica*, **2014**, 52, 88-90 | | 2 |
| 41 | Immunophenotypic and functional analysis of lymphocyte subsets in common variable immunodeficiency patients without monogenic defects.. *Scandinavian Journal of Immunology*, **2022**, e13164 | 3.4 | 2 |
| 40 | Generation of a human induced pluripotent stem cell line (PHAi003) from a primary immunodeficient patient with CD70 mutation. *Stem Cell Research*, **2019**, 41, 101612 | 1.6 | 1 |
| 39 | Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency.. *Journal of Clinical Immunology*, **2022**, 1 | 5.7 | 1 |
| 38 | Evaluation of physiciansQawareness of pediatric diseases in iran. *Iranian Journal of Pediatrics*, **2014**, 24, 87-92 | 1 | 1 |
| 37 | T Cell Repertoire Abnormality in Immunodeficiency Patients with DNA Repair and Methylation Defects. *Journal of Clinical Immunology*, **2021**, 1 | 5.7 | 1 |
| 36 | Agammaglobulinemia: Epidemiology, Pathogenesis, Clinical Phenotype, Diagnosis, Prognosis and Management. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 1434-1447 | 2.2 | 1 |
| 35 | 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 |
| 34 | Evaluation of Expression of LRBA and CTLA-4 Proteins in Common Variable Immunodeficiency Patients. *Immunological Investigations*, **2020**, 1-14 | 2.9 | 1 |
| 33 | Evidence-Based Immunotherapeutic Effects of Herbal Compounds on Humoral Immunity: Ancient and New Approaches. *Chinese Journal of Integrative Medicine*, **2021**, 27, 313-320 | 2.9 | 1 |
| 32 | Activation-induced deaminase is critical for the establishment of DNA methylation patterns prior to the germinal center reaction. *Nucleic Acids Research*, **2021**, 49, 5057-5073 | 20.1 | 1 |
| 31 | The First Iranian Cohort of Pediatric Patients with Activated Phosphoinositide 3-Kinase-[PI3K] Syndrome (APDS). *Immunological Investigations*, **2021**, 1-16 | 2.9 | 1 |
| 30 | Coronavirus: Pure Infectious Disease or Genetic Predisposition. *Advances in Experimental Medicine and Biology*, **2021**, 1318, 91-107 | 3.6 | 1 |
| 29 | Inborn errors of immunity **2021**, 1-8 | | 1 |
| 28 | Comprehensive Assessment of Skin Disorders in Patients with Common Variable Immunodeficiency´(CVID).. *Journal of Clinical Immunology*, **2022**, 1 | 5.7 | 0 |
| 27 | Gene Expression Network Analysis Identifies Potential Targets for Prevention of Preeclampsia.. *International Journal of General Medicine*, **2022**, 15, 1023-1032 | 2.3 | 0 |
| 26 | 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 |
| 25 | 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 |

| | | | |
|---|---|---|---|
| 24 | Primary Immunodeficiency and Thrombocytopenia. *International Reviews of Immunology*, **2021**, 1-43 | 4.6 | 0 |
| 23 | Immunodeficiencies affecting cellular and humoral immunity **2021**, 9-39 | | 0 |
| 22 | Primary Immunodeficiency Diseases in Iran: Past, Present and Future. *Archives of Iranian Medicine*, **2021**, 24, 118-124 | 2.4 | 0 |
| 21 | Adverse reactions in a large cohort of patients with inborn errors of immunity receiving intravenous immunoglobulin. *Clinical Immunology*, **2021**, 230, 108826 | 9 | 0 |
| 20 | Acupuncture combined with TCM bonesetting in the treatment of distal radius fractures: A protocol for systematic review and meta-analysis.. *Medicine (United States)*, **2021**, 100, e28279 | 1.8 | 0 |
| 19 | Disseminated Intravascular Coagulation Associated with Large Deletion of Immunoglobulin Heavy Chain.. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 778-783 | 1.1 | 0 |
| 18 | Clinical efficacy and safety of methotrexate compared with leflunomide in the treatment of rheumatoid arthritis: A protocol for systematic review and meta-analysis.. *Medicine (United States)*, **2021**, 100, e28285 | 1.8 | 0 |
| 17 | Clinical implications of experimental analyses of AID function on predictive computational tools: Challenge of missense variants. *Clinical Genetics*, **2020**, 97, 844-856 | 4 | |
| 16 | Recurrent Infections **2012**, 1-75 | | |
| 15 | Predominantly Antibody Deficiency **2012**, 113-192 | | |
| 14 | 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 | |
| 13 | Transient increased immunoglobulin levels in a hyper-IgM syndrome patient with COVID-19 infection. *Allergologia Et Immunopathologia*, **2021**, 49, 63-66 | 1.9 | |
| 12 | Genetic Risk Variants for Class Switching Recombination Defects in Ataxia-Telangiectasia Patients. *Journal of Clinical Immunology*, **2021**, 1 | 5.7 | |
| 11 | 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 | |
| 10 | Selective IgA Deficiency. *Rare Diseases of the Immune System*, **2019**, 201-215 | 0.2 | |
| 9 | Phenocopies of inborn errors of immunity **2021**, 317-344 | | |
| 8 | Diseases of immune dysregulation **2021**, 125-153 | | |
| 7 | Defects in intrinsic and innate immunity **2021**, 219-243 | | |

6 Combined immunodeficiencies with associated or syndromic features **2021**, 41-91

5 Complement deficiencies **2021**, 291-315

4 Predominantly antibody deficiencies **2021**, 93-123

3 Management of inborn errors of immunity **2021**, 345-361

2 Congenital defects of phagocytes **2021**, 155-217

1 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