# Victor Ef Ferraz

List of Publications by Year
in descending order



| 26 | 482 | 840776 11 | 713466 21 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 28 | 28 | 28 | 842 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Genetic services and testing in Brazil. Journal of Community Genetics, 2013, 4, 355-375. | 1.2 | 66 |
| 2 | Mucopolysaccharidoses in Brazil: What happens from birth to biochemical diagnosis?. American Journal of Medical Genetics, Part A, 2008, 146A, 1741-1747. | 1.2 | 63 |
| 3 | Recurrent Mutations in the Basic Domain of TWIST2 Cause Ablepharon Macrostomia and Barber-Say Syndromes. American Journal of Human Genetics, 2015, 97, 99-110. | 6.2 | 61 |
| 4 | Genetics and genomics in Brazil: a promising future. Molecular Genetics & Genomic Medicine, 2014, 2, 280-291. | 1.2 | 44 |
| 5 | Genetic education, knowledge and experiences between nurses and physicians in primary care in Brazil: A crossâ€sectional study. Australian Journal of Cancer Nursing, 2017, 19, 66-74. | 1.6 | 34 |
| 6 | Clinical Genetics in Developing Countries: The Case of Brazil. Public Health Genomics, 2004, 7, 95-105. | 1.0 | 23 |
| 7 | Aggravation of Eyelid and Conjunctival Malignancies Following Photodynamic Therapy in DeSanctis-Cacchione Syndrome. Ophthalmic Plastic and Reconstructive Surgery, 2006, 22, 498-499. | 0.8 | 23 |
| 8 | Ablepharon-Macrostomia syndrome: First report of familial occurrence. American Journal of Medical Genetics Part A, 2000, 94, 281-283. | 2.4 | 22 |
| 9 | TP53 p.R337H prevalence in a series of Brazilian hereditary breast cancer families. Hereditary Cancer in Clinical Practice, 2014, 12, 8. | 1.5 | 22 |
| 10 | Genotype-Phenotype Correlations in CYP1B1-Associated Primary Congenital Glaucoma Patients Representing Two Large Cohorts from India and Brazil. PLoS ONE, 2015, 10, e0127147. | 2.5 | 18 |
| 11 | Self-reported cancer family history is a useful tool for identification of individuals at risk of hereditary cancer predisposition syndrome at primary care centers in middle-income settings: a longitudinal study. Genetics and Molecular Biology, 2016, 39, 178-183. | 1.3 | 15 |
| 12 | Predictive models for mutations in mismatch repair genes: implication for genetic counseling in developing countries. BMC Cancer, 2012, 12, 64. | 2.6 | 11 |
| 13 | Perfil de CompetÃªncia em GenÃ©tica para MÃ©dicos do Brasil: uma Proposta da Sociedade Brasileira de GenÃ©tica MÃ©dica e GenÃ´mica. Revista Brasileira De Educacao Medica, 2019, 43, 440-450. | 0.2 | 9 |
| 14 | A further case of a Prader-Willi syndrome phenotype in a patient with Angelman syndrome molecular defect. Arquivos De Neuro-Psiquiatria, 2002, 60, 1011-1014. | 0.8 | 8 |
| 15 | The breast cancer immunophenotype of TP53-p.R337H carriers is different from that observed among other pathogenic TP53 mutation carriers. Familial Cancer, 2015, 14, 333-336. | 1.9 | 8 |
| 16 | Ablepharon macrostomia syndrome: A distinct genetic entity clinically related to the group of FRASâ€“FREM complex disorders. American Journal of Medical Genetics, Part A, 2013, 161, 3012-3017. | 1.2 | 7 |
| 17 | Practicability of comprehensive care in clinical genetics in the brazilian unified health system: expanding the debate. Texto E Contexto Enfermagem, 2014, 23, 1130-1135. | 0.4 | 6 |
| 18 | Molecular basis of familial adenomatous polyposis in the southeast of Brazil: identification of six novel mutations. International Journal of Biological Markers, 2019, 34, 80-89. | 1.8 | 6 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Lynch syndrome identification in a Brazilian cohort of endometrial cancer screened by a universal approach. Gynecologic Oncology, 2020, 159, 229-238. | 1.4 | 6 |
| 20 | Iris coloboma, blepharophimosis, arachnodactyly, joint contractures: Beals syndrome and Van den Endeâ€“Gupta syndrome phenotypic similarities. Clinical Dysmorphology, 2009, 18, 142-144. | 0.3 | 4 |
| 21 | I FÃ³rum Paulista de ServiÃ§os de Apoio ao Estudante de Medicina â€” Forsa Paulista â€” â€œA Carta de MarÃaâ€• Revista Brasileira De Educacao Medica, 2016, 40, 537-539. | 0.2 | 4 |
| 22 | Clinical-neurologic, cytogenetic and molecular aspects of the Prader-Willi and Angelman Syndromes. Arquivos De Neuro-Psiquiatria, 1997, 55, 199-208. | 0.8 | 3 |
| 23 | ImplementaÃ§Ã£o de um programa de mentoring para estudantes de graduaÃ§Ã£o em saÃºde: a experiÃªncia da FMRP-USP. Revista Brasileira De Educacao Medica, 2021, 45, . | 0.2 | 2 |
| 24 | First description of ultramutated endometrial cancer caused by germline loss-of-function and somatic exonuclease domain mutations in POLE gene. Genetics and Molecular Biology, 2020, 43, e20200100. | 1.3 | 2 |
| 25 | Competency Profile in Genetics for Physicians in Brazil: A Proposal of the Brazilian Society of Medical Genetics and Genomics. Revista Brasileira De Educacao Medica, 2019, 43, 440-450. | 0.2 | 2 |
| 26 | FMRP unida por um semestre melhor: narrativas de um pacto que celebra valores humanos e compromissos coletivos durante a pandemia da Covid-19. Medicina, 2021, 54, . | 0.1 | 0 |