Lukasz Kuszel

List of Publications by Year
in descending order

11	127	1478505 6	1372567 10
papers	citations	h-index	g-index

12	12	12	248
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Osteoarthritis and telomere shortening. Journal of Applied Genetics, 2015, 56, 169-176.	1.9	46
2	11p13 deletions can be more frequent than the PAX6 gene point mutations in Polish patients with aniridia. Journal of Applied Genetics, 2013, 54, 345-351.	1.9	15
3	EXOME REPORT: Novel mutation in ATP6V1B2 segregating with autosomal dominant epilepsy, intellectual disability and mild gingival and nail abnormalities. European Journal of Medical Genetics, 2020, 63, 103799.	1.3	14
4	Intrafamilial phenotypic variability in a Polish family with Sensenbrenner syndrome and biallelic WDR35 mutations. , 2017, 173, 1364-1368.		13
5	Case of Plasmodium knowlesi Malaria in Poland Linked to Travel in Southeast Asia. Emerging Infectious Diseases, 2019, 25, 1772-1773.	4.3	10
6	Prevalence of Anti-SARS-CoV-2 Antibodies in PoznaÅ„ Poland, after the First Wave of the COVID-19 Pandemic. Vaccines, 2021, 9, 541.	4.4	10
7	Severe neonatal spondylometaphyseal dysplasia in two siblings. American Journal of Medical Genetics, Part A, 2009, 149A, 2166-2172.	1.2	6
8	Massive Cryptosporidium infections and chronic diarrhea in HIV-negative patients. Parasitology Research, 2019, 118, 1937-1942.	1.6	5
9	Homozygous microdeletion in the 11p13 region in the patient with isolated form of aniridia: New challenges in the genetic diagnostics of aniridia. American Journal of Medical Genetics, Part A, 2021, , .	1.2	2
10	A new form or a variant of SMD type A4. Journal of Applied Genetics, 2012, 53, 289-294.	1.9	1
11	SMD Kozlowski type caused by p.Arg594His substitution in TRPV4 reveals abnormal ossification and notochordal remnants in discs and vertebrae. European Journal of Medical Genetics, 2017, 60, 509-516.	1.3	1