Elin TÃ¸nne

List of Publications by Year
in descending order

7 papers	132 citations	1684188 5 h-index	1720034 7 g-index
7 all docs	7 docs citations	7 times ranked	257 citing authors

#	Article	IF	Citations
1	Wholeâ€exome sequencing in syndromic craniosynostosis increases diagnostic yield and identifies candidate genes in osteogenic signaling pathways. American Journal of Medical Genetics, Part A, 2022, 188, 1464-1475.	1.2	7
2	Benefits of clinical criteria and high-throughput sequencing for diagnosing children with syndromic craniosynostosis. European Journal of Human Genetics, 2021, 29, 920-929.	2.8	13
3	Variants in the degron of AFF3 are associated with intellectual disability, mesomelic dysplasia, horseshoe kidney, and epileptic encephalopathy. American Journal of Human Genetics, 2021, 108, 857-873.	6.2	19
4	Truncating SRCAP variants outside the Floating-Harbor syndrome locus cause a distinct neurodevelopmental disorder with a specific DNA methylation signature. American Journal of Human Genetics, 2021, 108, 1053-1068.	6.2	31
5	Epidemiology of craniosynostosis in Norway. Journal of Neurosurgery: Pediatrics, 2020, 26, 68-75.	1.3	29
6	Fibromatosis in vertical rectus abdominis myocutaneous flap imitating tumor recurrence after surgery for locally advanced rectal cancer: case report. World Journal of Surgical Oncology, 2016, 14, 63.	1.9	3
7	Syndromic X-linked intellectual disability segregating with a missense variant in RLIM. European Journal of Human Genetics, 2015, 23, 1652-1656.	2.8	30