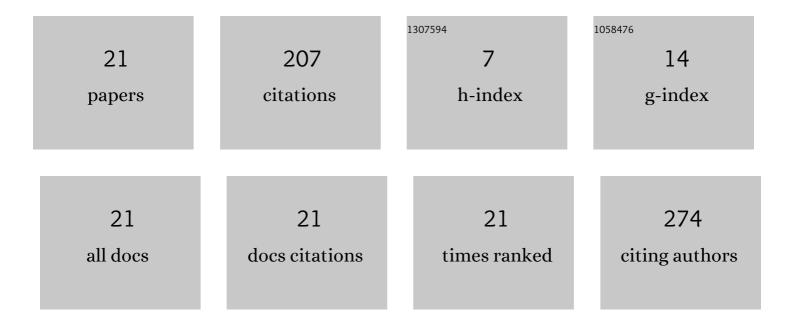
Peter J Tebben

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3423551/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Association of vitamin D deficiency with COVIDâ€19 infection severity: Systematic review and metaâ€analysis. Clinical Endocrinology, 2022, 96, 281-287.	2.4	65
2	Granulomatous Adenohypophysitis after Interferon and Ribavirin Therapy. Endocrine Practice, 2007, 13, 169-175.	2.1	26
3	Safety and efficacy of (+)â€epicatechin in subjects with Friedreich's ataxia: A phase <scp>II</scp> , openâ€label, prospective study. Journal of Inherited Metabolic Disease, 2021, 44, 502-514.	3.6	15
4	Hypercalcemia in Children Using the Ketogenic Diet: A Multicenter Study. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e485-e495.	3.6	13
5	Prevalence of Metabolic Bone Disease in Tube-Fed Children Receiving Elemental Formula. Hormone Research in Paediatrics, 2018, 90, 291-298.	1.8	12
6	Mild subclinical hypothyroidism is associated with paediatric dyslipidaemia. Clinical Endocrinology, 2018, 89, 330-335.	2.4	11
7	Patterns of amiodarone-induced thyroid dysfunction in infants and children. Heart Rhythm, 2019, 16, 1436-1442.	0.7	11
8	Kidney Cysts in Hypophosphatemic Rickets With Hypercalciuria: A Case Series. Kidney Medicine, 2022, 4, 100419.	2.0	8
9	Improved utilization of waistâ€ŧoâ€height ratio in cardiometabolic risk counselling in children: Application of DMAIC strategy. Journal of Evaluation in Clinical Practice, 2019, 25, 300-305.	1.8	7
10	Basal Ganglia Calcification Is Associated With Local and Systemic Metabolic Mechanisms in Adult Hypoparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 1900-1917.	3.6	7
11	24-Hydroxylase Deficiency Due to <i>CYP24A1</i> Sequence Variants: Comparison With Other Vitamin Dâ^'mediated Hypercalcemia Disorders. Journal of the Endocrine Society, 2021, 5, bvab119.	0.2	7
12	Variable Clinical Presentation of Children with Hereditary Hypophosphatemic Rickets with Hypercalciuria: A Case Series and Review of the Literature. Hormone Research in Paediatrics, 2021, 94, 374-389.	1.8	7
13	Congenital ichthyosis in Prader–Willi syndrome associated with maternal chromosome 15 uniparental disomy: Case report and review of autosomal recessive conditions unmasked by UPD. American Journal of Medical Genetics, Part A, 2020, 182, 2442-2449.	1.2	6
14	Lack of <i>GNAS</i> Remethylation During Oogenesis May Be a Cause of Sporadic Pseudohypoparathyroidism Type Ib. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1610-e1619.	3.6	5
15	Progression of PTH Resistance in Autosomal Dominant Pseudohypoparathyroidism Type Ib Due to Maternal <i>STX16</i> Deletions. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e681-e687.	3.6	3
16	70-Year-Old Woman With Buttock Pain and Hypercalcemia. Mayo Clinic Proceedings, 2014, 89, 1313-1317.	3.0	1
17	Onset of pituitary hormone deficiencies in optic nerve hypoplasia: a temporal trend analysis of 32 children at Mayo Clinic. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 139-145.	0.9	1
18	Growth hormone deficiency in a child with <scp>branchioâ€otoâ€renal</scp> spectrum disorder: Clinical evidence of <scp><i>EYA1</i></scp> in pituitary development and a recommendation for pituitary function surveillance. American Journal of Medical Genetics, Part A, 2021, 185, 261-266.	1.2	1

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#	Article	IF	CITATIONS
19	Skimmed breast milk for treatment of hypertriglyceridemia in an infant with congenital nephrotic syndrome. Nutrition in Clinical Practice, 2022, 37, 383-387.	2.4	1
20	Three Patient Kindred with a Novel Phenotype of Osteogenesis Imperfecta due to a Variant. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 218-224.	0.9	0
21	Three Patient Kindred with a Novel Phenotype of Osteogenesis Imperfecta due to a <i>COL1A1</i> Variant. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 218-224.	0.9	Ο