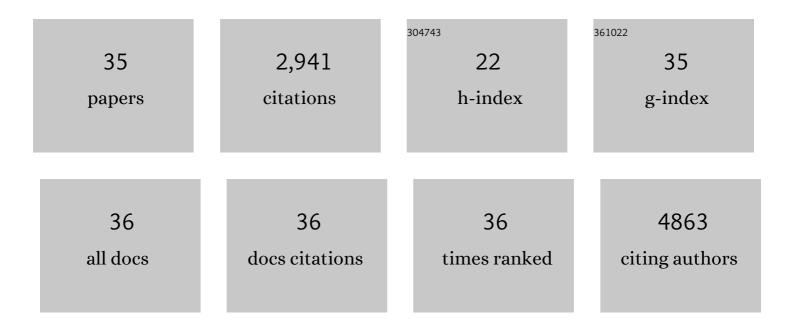
Scott J Diede

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Epacadostat plus pembrolizumab versus placebo plus pembrolizumab in patients with unresectable or metastatic melanoma (ECHO-301/KEYNOTE-252): a phase 3, randomised, double-blind study. Lancet Oncology, The, 2019, 20, 1083-1097.	10.7	611
2	Durable Complete Response After Discontinuation of Pembrolizumab in Patients With Metastatic Melanoma. Journal of Clinical Oncology, 2018, 36, 1668-1674.	1.6	360
3	Telomerase-Mediated Telomere Addition In Vivo Requires DNA Primase and DNA Polymerases α and δ. Cell, 1999, 99, 723-733.	28.9	351
4	The function of a stem-loop in telomerase RNA is linked to the DNA repair protein Ku. Nature Genetics, 2001, 27, 64-67.	21.4	205
5	Pembrolizumab in paediatric patients with advanced melanoma or a PD-L1-positive, advanced, relapsed, or refractory solid tumour or lymphoma (KEYNOTE-051): interim analysis of an open-label, single-arm, phase 1–2 trial. Lancet Oncology, The, 2020, 21, 121-133.	10.7	204
6	Exonuclease activity is required for sequence addition and Cdc13p loading at a de novo telomere. Current Biology, 2001, 11, 1336-1340.	3.9	136
7	KEYNOTE-022 part 3: a randomized, double-blind, phase 2 study of pembrolizumab, dabrafenib, and trametinib in <i>BRAF</i> -mutant melanoma. , 2020, 8, e001806.		110
8	A Phase Ib Study of Pembrolizumab as Second-Line Therapy for Chinese Patients With Advanced or Metastatic Melanoma (KEYNOTE-151). Translational Oncology, 2019, 12, 828-835.	3.7	90
9	MyoD and E-protein heterodimers switch rhabdomyosarcoma cells from an arrested myoblast phase to a differentiated state. Genes and Development, 2009, 23, 694-707.	5.9	84
10	Comparison of Genome-Wide Binding of MyoD in Normal Human Myogenic Cells and Rhabdomyosarcomas Identifies Regional and Local Suppression of Promyogenic Transcription Factors. Molecular and Cellular Biology, 2013, 33, 773-784.	2.3	62
11	DNA methylation of developmental genes in pediatric medulloblastomas identified by denaturation analysis of methylation differences. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 234-239.	7.1	59
12	Long-term safety of pembrolizumab monotherapy and relationship with clinical outcome: A landmark analysis in patients with advanced melanoma. European Journal of Cancer, 2021, 144, 182-191.	2.8	57
13	Genome-wide DNA methylation studies suggest distinct DNA methylation patterns in pediatric embryonal and alveolar rhabdomyosarcomas. Epigenetics, 2012, 7, 400-408.	2.7	56
14	ACCELERATE and European Medicines Agency Paediatric Strategy Forum for medicinal product development of checkpoint inhibitors for use in combination therapy in paediatric patients. European Journal of Cancer, 2020, 127, 52-66.	2.8	52
15	A duplication at chromosome 11q12.2-11q12.3 is associated with spinocerebellar ataxia type 20. Human Molecular Genetics, 2008, 17, 3847-3853.	2.9	50
16	Spontaneous regression of metastatic cancer: learning from neuroblastoma. Nature Reviews Cancer, 2014, 14, 71-72.	28.4	46
17	Assessment of palindromes as platforms for DNA amplification in breast cancer. Genome Research, 2012, 22, 232-245.	5.5	31
18	Phase I study of vorinostat in combination with isotretinoin in patients with refractory/recurrent neuroblastoma: A new approaches to Neuroblastoma Therapy (NANT) trial. Pediatric Blood and Cancer, 2018, 65, e27023.	1.5	31

#	Article	IF	CITATIONS
19	Treatment patterns and outcomes for patients with advanced melanoma in US oncology clinical practices. Future Oncology, 2019, 15, 459-471.	2.4	27

Real-world experience with pembrolizumab in patients with advanced melanoma. Medicine (United) Tj ETQq000 rgBT /Overlock 10 Tf 5

21	Expression of neurotrophins and the low-affinity NGF receptor in septal and hippocampal reaggregate cultures: local physiologic effects of NGF synthesized in the septal region. Developmental Brain Research, 1992, 70, 123-133.	1.7	23
22	EEL-1, a Hect E3 ubiquitin ligase, controls asymmetry and persistence of the SKN-1 transcription factor in the early C. elegans embryo. Development (Cambridge), 2007, 134, 2303-2314.	2.5	23
23	Homology-mediated end-capping as a primary step of sister chromatid fusion in the breakage-fusion-bridge cycles. Nucleic Acids Research, 2013, 41, 9732-9740.	14.5	17
24	Health-related quality of life (QoL) in patients with advanced melanoma receiving immunotherapies in real-world clinical practice settings. Quality of Life Research, 2020, 29, 2651-2660.	3.1	17
25	Fundamental differences in promoter CpG island DNA hypermethylation between human cancer and genetically engineered mouse models of cancer. Epigenetics, 2013, 8, 1254-1260.	2.7	16
26	Impediment of Replication Forks by Long Non-coding RNA Provokes Chromosomal Rearrangements by Error-Prone Restart. Cell Reports, 2017, 21, 2223-2235.	6.4	13
27	Comparison of Duration of Response vs Conventional Response Rates and Progression-Free Survival as Efficacy End Points in Simulated Immuno-oncology Clinical Trials. JAMA Network Open, 2021, 4, e218175.	5.9	13
28	Long-term outcomes in patients with advanced melanoma who had initial stable disease with pembrolizumab in KEYNOTE-001 and KEYNOTE-006. European Journal of Cancer, 2021, 157, 391-402.	2.8	13
29	Reduced Dosage of pos-1 Suppresses Mex Mutants and Reveals Complex Interactions Among CCCH Zinc-Finger Proteins During Caenorhabditis elegans Embryogenesis. Genetics, 2006, 174, 1933-1945.	2.9	12
30	TERT hypermethylation: biomarker in paediatric brain tumours. Lancet Oncology, The, 2013, 14, 447-448.	10.7	7
31	All Things Must End: Telomere Dynamics in Yeast. Cold Spring Harbor Symposia on Quantitative Biology, 2000, 65, 281-296.	1.1	7
32	Hospitalization and emergency department utilization in patients with advanced melanoma receiving pembrolizumab versus ipilimumab plus nivolumab in US academic centers. Journal of Medical Economics, 2020, 23, 132-138.	2.1	6
33	Genome-wide analysis of palindrome formation. Nature Genetics, 2010, 42, 279-279.	21.4	5
34	KEYNOTE-022: Pembrolizumab with trametinib in patients with BRAF wild-type melanoma or advanced solid tumours irrespective of BRAF mutation. European Journal of Cancer, 2022, 160, 1-11.	2.8	4
35	Factors associated with immunotherapy selection in patients with advanced melanoma. Immunotherapy, 2018, 10, 1361-1369.	2.0	2