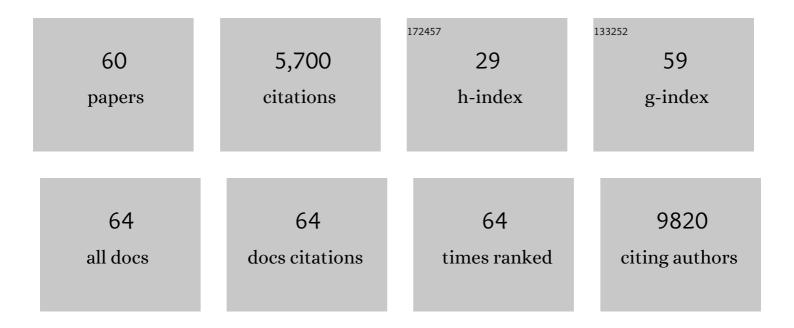
Sophie Postel-Vinay

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	Hyperprogressive Disease Is a New Pattern of Progression in Cancer Patients Treated by Anti-PD-1/PD-L1. Clinical Cancer Research, 2017, 23, 1920-1928.	7.0	960
2	High-Throughput Genomics and Clinical Outcome in Hard-to-Treat Advanced Cancers: Results of the MOSCATO 01 Trial. Cancer Discovery, 2017, 7, 586-595.	9.4	554
3	Lorlatinib in non-small-cell lung cancer with ALK or ROS1 rearrangement: an international, multicentre, open-label, single-arm first-in-man phase 1 trial. Lancet Oncology, The, 2017, 18, 1590-1599.	10.7	535
4	Targeting FGFR Signaling in Cancer. Clinical Cancer Research, 2015, 21, 2684-2694.	7.0	399
5	Safety, pharmacokinetics, and preliminary activity of the anti-IGF-1R antibody figitumumab (CP-751,871) in patients with sarcoma and Ewing's sarcoma: a phase 1 expansion cohort study. Lancet Oncology, The, 2010, 11, 129-135.	10.7	334
6	Combining epigenetic drugs with other therapies for solid tumours — past lessons and future promise. Nature Reviews Clinical Oncology, 2020, 17, 91-107.	27.6	283
7	Olaparib and durvalumab in patients with germline BRCA-mutated metastatic breast cancer (MEDIOLA): an open-label, multicentre, phase 1/2, basket study. Lancet Oncology, The, 2020, 21, 1155-1164.	10.7	274
8	PARP inhibition enhances tumor cell–intrinsic immunity in ERCC1-deficient non–small cell lung cancer. Journal of Clinical Investigation, 2019, 129, 1211-1228.	8.2	222
9	Mutational Landscape and Sensitivity to Immune Checkpoint Blockers. Clinical Cancer Research, 2016, 22, 4309-4321.	7.0	182
10	Common variants near TARDBP and EGR2 are associated with susceptibility to Ewing sarcoma. Nature Genetics, 2012, 44, 323-327.	21.4	160
11	Targeting the DNA damage response in immuno-oncology: developments and opportunities. Nature Reviews Cancer, 2021, 21, 701-717.	28.4	150
12	Phase I Trials of Molecularly Targeted Agents: Should We Pay More Attention to Late Toxicities?. Journal of Clinical Oncology, 2011, 29, 1728-1735.	1.6	120
13	Towards new methods for the determination of dose limiting toxicities and the assessment of the recommended dose for further studies of molecularly targeted agents – Dose-Limiting Toxicity and Toxicity Assessment Recommendation Group for Early Trials of Targeted therapies, an European Organisation for Research and Treatment of Cancer-led study. European Journal of Cancer, 2014, 50,	2.8	104
14	2010 2019. Circulating Cell-Free Tumor DNA Analysis of 50 Genes by Next-Generation Sequencing in the Prospective MOSCATO Trial. Clinical Cancer Research, 2016, 22, 2960-2968.	7.0	103
15	The potential of exploiting DNA-repair defects for optimizing lung cancer treatment. Nature Reviews Clinical Oncology, 2012, 9, 144-155.	27.6	96
16	Long-Term Survival in Patients Responding to Anti–PD-1/PD-L1 Therapy and Disease Outcome upon Treatment Discontinuation. Clinical Cancer Research, 2019, 25, 946-956.	7.0	96
17	Challenges of phase 1 clinical trials evaluating immune checkpoint-targeted antibodies. Annals of Oncology, 2016, 27, 214-224.	1.2	86
18	First-in-human phase I study of the bromodomain and extraterminal motif inhibitor BAY 1238097: emerging pharmacokinetic/pharmacodynamic relationship and early termination due to unexpected toxicity. European Journal of Cancer, 2019, 109, 103-110.	2.8	76

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19	Epigenetic modifiers as new immunomodulatory therapies in solid tumours. Annals of Oncology, 2018, 29, 812-824.	1.2	73
20	Can molecular biomarker-based patient selection in Phase I trials accelerate anticancer drug development?. Drug Discovery Today, 2010, 15, 88-97.	6.4	69
21	PBRM1 Deficiency Confers Synthetic Lethality to DNA Repair Inhibitors in Cancer. Cancer Research, 2021, 81, 2888-2902.	0.9	66
22	Defining dose-limiting toxicity for phase 1 trials of molecularly targeted agents: Results of a DLT-TARGETT international survey. European Journal of Cancer, 2014, 50, 2050-2056.	2.8	63
23	First-in-human Phase 1 open label study of the BET inhibitor ODM-207 in patients with selected solid tumours. British Journal of Cancer, 2020, 123, 1730-1736.	6.4	63
24	TPF induction chemotherapy increases PD-L1 expression in tumour cells and immune cells in head and neck squamous cell carcinoma. ESMO Open, 2018, 3, e000257.	4.5	62
25	Ceralasertib (AZD6738), an Oral ATR Kinase Inhibitor, in Combination with Carboplatin in Patients with Advanced Solid Tumors: A Phase I Study. Clinical Cancer Research, 2021, 27, 5213-5224.	7.0	53
26	Phase I Study of GDC-0425, a Checkpoint Kinase 1 Inhibitor, in Combination with Gemcitabine in Patients with Refractory Solid Tumors. Clinical Cancer Research, 2017, 23, 2423-2432.	7.0	50
27	ERCC1 function in nuclear excision and interstrand crosslink repair pathways is mediated exclusively by the ERCC1-202 isoform. Cell Cycle, 2013, 12, 3298-3306.	2.6	37
28	Phase 1 Study of Tazemetostat (EPZ-6438), an Inhibitor of Enhancer of Zeste-Homolog 2 (EZH2): Preliminary Safety and Activity in Relapsed or Refractory Non-Hodgkin Lymphoma (NHL) Patients. Blood, 2015, 126, 473-473.	1.4	37
29	AXL and acquired resistance to EGFR inhibitors. Nature Genetics, 2012, 44, 835-836.	21.4	31
30	Exploiting epigenetic vulnerabilities in solid tumors: Novel therapeutic opportunities in the treatment of SWI/SNF-defective cancers. Seminars in Cancer Biology, 2020, 61, 180-198.	9.6	28
31	Reovirus: Rationale and clinical trial update. Current Opinion in Molecular Therapeutics, 2009, 11, 532-9.	2.8	25
32	<i>JAK</i> Mutations as Escape Mechanisms to Anti–PD-1 Therapy. Cancer Discovery, 2017, 7, 128-130.	9.4	24
33	Immunotherapy for SMARCB1-Deficient Sarcomas: Current Evidence and Future Developments. Biomedicines, 2022, 10, 650.	3.2	24
34	Evidence of pseudoprogression in patients treated with PD1/PDL1 antibodies across tumor types. Cancer Medicine, 2020, 9, 2643-2652.	2.8	21
35	Patient-reported tolerability of adverse events in phase 1 trials. ESMO Open, 2017, 2, e000148.	4.5	20
36	In the immuno-oncology era, is anti-PD-1 or anti-PD-L1 immunotherapy modifying the sensitivity to conventional cancer therapies?. European Journal of Cancer, 2017, 87, 65-74.	2.8	19

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37	DNA repair deficiency sensitizes lung cancer cells to NAD+ biosynthesis blockade. Journal of Clinical Investigation, 2018, 128, 1671-1687.	8.2	19
38	Chromatin Regulators as a Guide for Cancer Treatment Choice. Molecular Cancer Therapeutics, 2016, 15, 1768-1777.	4.1	18
39	Beyond DNA repair: the novel immunological potential of PARP inhibitors. Molecular and Cellular Oncology, 2019, 6, 1-4.	0.7	18
40	Chemotherapy beyond immune checkpoint inhibitors in patients with metastatic colorectal cancer. European Journal of Cancer, 2020, 137, 117-126.	2.8	16
41	Clinical benefit of early phase clinical trial participation for advanced sarcoma patients. Cancer Chemotherapy and Pharmacology, 2011, 68, 423-429.	2.3	14
42	Patterns of progression in patients treated for immuno-oncology antibodies combination. Cancer Immunology, Immunotherapy, 2021, 70, 221-232.	4.2	12
43	Efficacy of histology-agnostic and molecularly-driven HER2 inhibitors for refractory cancers. Oncotarget, 2018, 9, 9741-9750.	1.8	12
44	A Case-Control Study Brings to Light the Causes of Screen Failures in Phase 1 Cancer Clinical Trials. PLoS ONE, 2016, 11, e0154895.	2.5	10
45	Seeking the driver in tumours with apparent normal molecular profile on comparative genomic hybridization and targeted gene panel sequencing: what is the added value of whole exome sequencing?. Annals of Oncology, 2016, 27, 344-352.	1.2	9
46	Understanding genetic determinants of resistance to immune checkpoint blockers. Seminars in Cancer Biology, 2020, 65, 123-139.	9.6	9
47	Acquired EGFR Mutation as the Potential Resistance Driver to Crizotinib in a MET-Mutated Tumor. Journal of Thoracic Oncology, 2016, 11, e21-e23.	1.1	8
48	Time to progression ratio in cancer patients enrolled in early phase clinical trials: time for new guidelines?. British Journal of Cancer, 2018, 119, 937-939.	6.4	7
49	Impact of COVID-19 Pandemic on Cancer Research. Cancer Cell, 2020, 38, 591-593.	16.8	7
50	How Much Can We Bet on Activity of BET Inhibitors Beyond NUT–Midline Carcinoma?. JNCI Cancer Spectrum, 2020, 4, pkz092.	2.9	7
51	Coronavirus disease (COVID-19) outbreak and phase 1 trials: should we consider a specific patient management?. European Journal of Cancer, 2020, 137, 235-239.	2.8	7
52	3D Functional Genomics Screens Identify CREBBP as a Targetable Driver in Aggressive Triple-Negative Breast Cancer. Cancer Research, 2021, 81, 847-859.	0.9	7
53	Patients aged over 75 years enrolled in Phase I clinical trials: the <scp>G</scp> ustave <scp>R</scp> oussy experience. International Journal of Cancer, 2016, 138, 875-880.	5.1	5
54	Outcomes and prognostic factors for relapsed or refractory lymphoma patients in phase I clinical trials. Investigational New Drugs, 2018, 36, 62-74.	2.6	3

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55	A Novel Synthetic Lethal Approach to Target <i>MYC</i> -Driven Cancers. Cancer Research, 2022, 82, 969-971.	0.9	3
56	Sustained cancer clinical trial activity in a French hospital during the first wave of the COVID-19 pandemic. Cancer Cell, 2021, 39, 1039-1041.	16.8	2
57	Prognostic factors and outcome of patients with hematological malignancies in phase I trials. Anti-Cancer Drugs, 2017, 28, 540-545.	1.4	1
58	Cardiac troponin I elevation and overall survival among cancer patients receiving investigational compounds during phase I trials. International Journal of Cardiology, 2016, 214, 364-369.	1.7	0
59	Immune Therapies in Phase 1 Trials. , 2018, , 547-563.		Ο
60	You BETer be aware – learnings from a negative Phase 1 study. Oncotarget, 2019, 10, 3145-3146.	1.8	0