

Sarah M Senf

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/4298492/sarah-m-senf-publications-by-year.pdf>

Version: 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

28
papers

1,339
citations

19
h-index

30
g-index

30
ext. papers

1,553
ext. citations

5
avg, IF

4.53
L-index

#	Paper	IF	Citations
28	FoxP1 is a transcriptional repressor associated with cancer cachexia that induces skeletal muscle wasting and weakness. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2021 , 12, 421-442	10.3	1
27	The Florida Pancreas Collaborative Next-Generation Biobank: Infrastructure to Reduce Disparities and Improve Survival for a Diverse Cohort of Patients with Pancreatic Cancer. <i>Cancers</i> , 2021 , 13,	6.6	2
26	MEF2c-Dependent Downregulation of Myocilin Mediates Cancer-Induced Muscle Wasting and Associates with Cachexia in Patients with Cancer. <i>Cancer Research</i> , 2020 , 80, 1861-1874	10.1	8
25	Distinct cachexia profiles in response to human pancreatic tumours in mouse limb and respiratory muscle. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2020 , 11, 820-837	10.3	14
24	Nicotine Induces IL-8 Secretion from Pancreatic Cancer Stroma and Worsens Cancer-Induced Cachexia. <i>Cancers</i> , 2020 , 12,	6.6	2
23	IL-8 Released from Human Pancreatic Cancer and Tumor-Associated Stromal Cells Signals through a CXCR2-ERK1/2 Axis to Induce Muscle Atrophy. <i>Cancers</i> , 2019 , 11,	6.6	23
22	Mas Receptor Activation Slows Tumor Growth and Attenuates Muscle Wasting in Cancer. <i>Cancer Research</i> , 2019 , 79, 706-719	10.1	14
21	Skeletal Muscle Fibrosis in Pancreatic Cancer Patients with Respect to Survival. <i>JNCI Cancer Spectrum</i> , 2018 , 2, pky043	4.6	27
20	Local and Systemic Cytokine Profiling for Pancreatic Ductal Adenocarcinoma to Study Cancer Cachexia in an Era of Precision Medicine. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	11
19	Tumour-derived leukaemia inhibitory factor is a major driver of cancer cachexia and morbidity in C26 tumour-bearing mice. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2018 , 9, 1109-1120	10.3	39
18	Orthotopic Patient-Derived Pancreatic Cancer Xenografts Engraft Into the Pancreatic Parenchyma, Metastasize, and Induce Muscle Wasting to Recapitulate the Human Disease. <i>Pancreas</i> , 2017 , 46, 813-819	2.6	28
17	A clinically applicable muscular index predicts long-term survival in resectable pancreatic cancer. <i>Surgery</i> , 2017 , 161, 930-938	3.6	28
16	Human pancreatic cancer xenografts recapitulate key aspects of cancer cachexia. <i>Oncotarget</i> , 2017 , 8, 1177-1189	3.3	18
15	Inducible HSP70 is critical in preventing the aggregation and enhancing the processing of PMP22. <i>ASN Neuro</i> , 2015 , 7,	5.3	19
14	Identification of the Acetylation and Ubiquitin-Modified Proteome during the Progression of Skeletal Muscle Atrophy. <i>PLoS ONE</i> , 2015 , 10, e0136247	3.7	28
13	Transcriptional regulation of myotrophic actions by testosterone and trenbolone on androgen-responsive muscle. <i>Steroids</i> , 2014 , 87, 59-66	2.8	21
12	HDAC1 activates FoxO and is both sufficient and required for skeletal muscle atrophy. <i>Journal of Cell Science</i> , 2014 , 127, 1441-53	5.3	79

11	Genome-wide identification of FoxO-dependent gene networks in skeletal muscle during C26 cancer cachexia. <i>BMC Cancer</i> , 2014 , 14, 997	4.8	64
10	Skeletal muscle heat shock protein 70: diverse functions and therapeutic potential for wasting disorders. <i>Frontiers in Physiology</i> , 2013 , 4, 330	4.6	53
9	Loss of the inducible Hsp70 delays the inflammatory response to skeletal muscle injury and severely impairs muscle regeneration. <i>PLoS ONE</i> , 2013 , 8, e62687	3.7	76
8	Determination of gene promoter activity in skeletal muscles in vivo. <i>Methods in Molecular Biology</i> , 2012 , 798, 461-72	1.4	3
7	Inhibition of FoxO transcriptional activity prevents muscle fiber atrophy during cachexia and induces hypertrophy. <i>FASEB Journal</i> , 2012 , 26, 987-1000	0.9	140
6	Inhibition of IkappaB kinase alpha (IKK α) or IKKbeta (IKK β) plus forkhead box O (Foxo) abolishes skeletal muscle atrophy. <i>Biochemical and Biophysical Research Communications</i> , 2011 , 405, 491-6	3.4	52
5	p300 Acetyltransferase activity differentially regulates the localization and activity of the FOXO homologues in skeletal muscle. <i>American Journal of Physiology - Cell Physiology</i> , 2011 , 300, C1490-501	5.4	78
4	FOXO signaling is required for disuse muscle atrophy and is directly regulated by Hsp70. <i>American Journal of Physiology - Cell Physiology</i> , 2010 , 298, C38-45	5.4	133
3	Ros-mediated activation of NF-kappaB and Foxo during muscle disuse. <i>Muscle and Nerve</i> , 2010 , 41, 110-33.4	3.4	87
2	Hsp27 inhibits IKKbeta-induced NF-kappaB activity and skeletal muscle atrophy. <i>FASEB Journal</i> , 2009 , 23, 3415-23	0.9	66
1	Hsp70 overexpression inhibits NF-kappaB and Foxo3a transcriptional activities and prevents skeletal muscle atrophy. <i>FASEB Journal</i> , 2008 , 22, 3836-45	0.9	225