

# Shenhav Cohen

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1584512/publications.pdf>

Version: 2024-02-01

21  
papers

2,520  
citations

623734

14  
h-index

794594

19  
g-index

25  
all docs

25  
docs citations

25  
times ranked

4775  
citing authors

#	ARTICLE	IF	CITATIONS
1	Muscle wasting in disease: molecular mechanisms and promising therapies. <i>Nature Reviews Drug Discovery</i> , 2015, 14, 58-74.	46.4	792
2	During muscle atrophy, thick, but not thin, filament components are degraded by MuRF1-dependent ubiquitylation. <i>Journal of Cell Biology</i> , 2009, 185, 1083-1095.	5.2	499
3	The extracellular matrix protein agrin promotes heart regeneration in mice. <i>Nature</i> , 2017, 547, 179-184.	27.8	498
4	Ubiquitylation by Trim32 causes coupled loss of desmin, Z-bands, and thin filaments in muscle atrophy. <i>Journal of Cell Biology</i> , 2012, 198, 575-589.	5.2	165
5	Trim32 reduces PI3K/Akt/FoxO signaling in muscle atrophy by promoting plakoglobin/PI3K dissociation. <i>Journal of Cell Biology</i> , 2014, 204, 747-758.	5.2	82
6	The trans-Golgi network-associated human ubiquitin-protein ligase POSH is essential for HIV type 1 production. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 1478-1483.	7.1	71
7	USP1 deubiquitinates Akt to inhibit PI3K/Akt/FoxO signaling in muscle during prolonged starvation. <i>EMBO Reports</i> , 2020, 21, e48791.	4.5	64
8	GSK3- $\beta$ promotes calpain-mediated desmin filament depolymerization and myofibril loss in atrophy. <i>Journal of Cell Biology</i> , 2018, 217, 3698-3714.	5.2	58
9	Myofibril breakdown during atrophy is a delayed response requiring the transcription factor PAX4 and desmin depolymerization. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E1375-E1384.	7.1	50
10	New roles for desmin in the maintenance of muscle homeostasis. <i>FEBS Journal</i> , 2022, 289, 2755-2770.	4.7	50
11	Non-canonical activation of DAPK2 by AMPK constitutes a new pathway linking metabolic stress to autophagy. <i>Nature Communications</i> , 2018, 9, 1759.	12.8	33
12	A signaling hub of insulin receptor, dystrophin glycoprotein complex and plakoglobin regulates muscle size. <i>Nature Communications</i> , 2020, 11, 1381.	12.8	33
13	Expression of a PKR Dominant-Negative Mutant in Myogenic Cells Interferes with the Myogenic Process. <i>Experimental Cell Research</i> , 2000, 254, 45-54.	2.6	24
14	Role of calpains in promoting desmin filaments depolymerization and muscle atrophy. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2020, 1867, 118788.	4.1	24
15	Profiling of the muscle-specific dystroglycan interactome reveals the role of Hippo signaling in muscular dystrophy and age-dependent muscle atrophy. <i>BMC Medicine</i> , 2020, 18, 8.	5.5	20
16	A semiautomated measurement of muscle fiber size using the Imaris software. <i>American Journal of Physiology - Cell Physiology</i> , 2021, 321, C615-C631.	4.6	17
17	VWA domain of S5a restricts the ability to bind ubiquitin and Ubl to the 26S proteasome. <i>Molecular Biology of the Cell</i> , 2014, 25, 3988-3998.	2.1	15
18	Breakdown of Filamentous Myofibrils by the UPS Step by Step. <i>Biomolecules</i> , 2021, 11, 110.	4.0	13

#	ARTICLE	IF	CITATIONS
19	JAK-STAT signaling involved in phorbol 12-myristate 13-acetate- and dimethyl sulfoxide-induced $2\alpha$ -5 $\alpha$ - $2$ oligoadenylate synthetase expression in human HL-60 leukemia cells. <i>Leukemia Research</i> , 2005, 29, 923-931.	0.8	12
20	During muscle atrophy, thick, but not thin, filament components are degraded by MuRF1-dependent ubiquitylation. <i>Journal of Experimental Medicine</i> , 2009, 206, i13-i13.	8.5	0
21	Reply to Kissane and Eggington. <i>American Journal of Physiology - Cell Physiology</i> , 2021, 321, C1084-C1085.	4.6	0