

Kate T Murphy

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/8135394/kate-t-murphy-publications-by-year.pdf>
Version: 2024-04-09

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.

| | | | |
|-------------------|-------------------------|---------------|-----------------|
| 35 papers | 1,519 citations | 21 h-index | 38 g-index |
| 38 ext. papers | 1,766 ext. citations | 6 avg, IF | 4.42 L-index |

| # | Paper | IF | Citations |
|----|---|------|-----------|
| 35 | Phosphorylation of ERK and dystrophin S3059 protects against inflammation-associated C2C12 myotube atrophy. <i>American Journal of Physiology - Cell Physiology</i> , 2021 , 320, C956-C965 | 5.4 | 3 |
| 34 | Sweet Syndrome in Eosinophilic Granulomatosis with Polyangiitis. <i>Journal of Rheumatology</i> , 2020 , 47, 1031-1032 | 4.1 | 2 |
| 33 | Mas Receptor Activation Slows Tumor Growth and Attenuates Muscle Wasting in Cancer. <i>Cancer Research</i> , 2019 , 79, 706-719 | 10.1 | 14 |
| 32 | Smad7 gene delivery prevents muscle wasting associated with cancer cachexia in mice. <i>Science Translational Medicine</i> , 2016 , 8, 348ra98 | 17.5 | 45 |
| 31 | BGP-15 Improves Aspects of the Dystrophic Pathology in mdx and dko Mice with Differing Efficacies in Heart and Skeletal Muscle. <i>American Journal of Pathology</i> , 2016 , 186, 3246-3260 | 5.8 | 19 |
| 30 | The pathogenesis and treatment of cardiac atrophy in cancer cachexia. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2016 , 310, H466-77 | 5.2 | 57 |
| 29 | Disease-Induced Skeletal Muscle Atrophy and Fatigue. <i>Medicine and Science in Sports and Exercise</i> , 2016 , 48, 2307-2319 | 1.2 | 79 |
| 28 | Glucose-6-phosphate dehydrogenase contributes to the regulation of glucose uptake in skeletal muscle. <i>Molecular Metabolism</i> , 2016 , 5, 1083-1091 | 8.8 | 15 |
| 27 | Targeting of Fn14 Prevents Cancer-Induced Cachexia and Prolongs Survival. <i>Cell</i> , 2015 , 162, 1365-78 | 56.2 | 82 |
| 26 | Elevated expression of activins promotes muscle wasting and cachexia. <i>FASEB Journal</i> , 2014 , 28, 1711-23 | 3.9 | 130 |
| 25 | Glycine administration attenuates skeletal muscle wasting in a mouse model of cancer cachexia. <i>Clinical Nutrition</i> , 2014 , 33, 448-58 | 5.9 | 59 |
| 24 | Phosphorylation within the cysteine-rich region of dystrophin enhances its association with Edystroglycan and identifies a potential novel therapeutic target for skeletal muscle wasting. <i>Human Molecular Genetics</i> , 2014 , 23, 6697-711 | 5.6 | 10 |
| 23 | Inhibition of the renin-angiotensin system improves physiological outcomes in mice with mild or severe cancer cachexia. <i>International Journal of Cancer</i> , 2013 , 133, 1234-46 | 7.5 | 25 |
| 22 | Physiological characterization of a mouse model of cachexia in colorectal liver metastases. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2013 , 304, R854-64 | 3.2 | 12 |
| 21 | Infusion with the antioxidant N-acetylcysteine attenuates early adaptive responses to exercise in human skeletal muscle. <i>Acta Physiologica</i> , 2012 , 204, 382-92 | 5.6 | 67 |
| 20 | Disruption of muscle renin-angiotensin system in AT1a ^{-/-} mice enhances muscle function despite reducing muscle mass but compromises repair after injury. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2012 , 303, R321-31 | 3.2 | 12 |
| 19 | Impaired exercise performance and muscle Na(+),K(+)-pump activity in renal transplantation and haemodialysis patients. <i>Nephrology Dialysis Transplantation</i> , 2012 , 27, 2036-43 | 4.3 | 13 |

| | | | |
|----|--|-----|-----|
| 18 | Parvalbumin gene transfer impairs skeletal muscle contractility in old mice. <i>Human Gene Therapy</i> , 2012 , 23, 824-36 | 4.8 | 6 |
| 17 | Importance of functional and metabolic impairments in the characterization of the C-26 murine model of cancer cachexia. <i>DMM Disease Models and Mechanisms</i> , 2012 , 5, 533-45 | 4.1 | 80 |
| 16 | Chronic formoterol administration reduces cardiac mitochondrial protein synthesis and oxidative capacity in mice. <i>International Journal of Cardiology</i> , 2011 , 146, 270-2 | 3.2 | 11 |
| 15 | Acute antibody-directed myostatin inhibition attenuates disuse muscle atrophy and weakness in mice. <i>Journal of Applied Physiology</i> , 2011 , 110, 1065-72 | 3.7 | 39 |
| 14 | Antibody-directed myostatin inhibition enhances muscle mass and function in tumor-bearing mice. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2011 , 301, R716-26 | 3.2 | 77 |
| 13 | Cellular mechanisms underlying temporal changes in skeletal muscle protein synthesis and breakdown during chronic {beta}-adrenoceptor stimulation in mice. <i>Journal of Physiology</i> , 2010 , 588, 4811-23 | 3.9 | 50 |
| 12 | Antibody-directed myostatin inhibition in 21-mo-old mice reveals novel roles for myostatin signaling in skeletal muscle structure and function. <i>FASEB Journal</i> , 2010 , 24, 4433-42 | 0.9 | 101 |
| 11 | Antibody-directed myostatin inhibition improves diaphragm pathology in young but not adult dystrophic mdx mice. <i>American Journal of Pathology</i> , 2010 , 176, 2425-34 | 5.8 | 45 |
| 10 | Update on emerging drugs for cancer cachexia. <i>Expert Opinion on Emerging Drugs</i> , 2009 , 14, 619-32 | 3.7 | 33 |
| 9 | Analysis of exercise-induced Na ⁺ -K ⁺ exchange in rat skeletal muscle in vivo. <i>Experimental Physiology</i> , 2008 , 93, 1249-62 | 2.4 | 25 |
| 8 | Antioxidant treatment with N-acetylcysteine regulates mammalian skeletal muscle Na ⁺ -K ⁺ -ATPase alpha gene expression during repeated contractions. <i>Experimental Physiology</i> , 2008 , 93, 1239-48 | 2.4 | 16 |
| 7 | Chronic beta2-adrenoceptor stimulation impairs cardiac relaxation via reduced SR Ca ²⁺ -ATPase protein and activity. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2008 , 294, H2587-95 | 5.2 | 21 |
| 6 | Effects of endurance training status and sex differences on Na ⁺ ,K ⁺ -pump mRNA expression, content and maximal activity in human skeletal muscle. <i>Acta Physiologica</i> , 2007 , 189, 259-69 | 5.6 | 17 |
| 5 | Muscle Na ⁺ -K ⁺ -ATPase activity and isoform adaptations to intense interval exercise and training in well-trained athletes. <i>Journal of Applied Physiology</i> , 2007 , 103, 39-47 | 3.7 | 39 |
| 4 | Exercise performance falls over time in patients with chronic kidney disease despite maintenance of hemoglobin concentration. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2006 , 1, 488-95 | 6.9 | 49 |
| 3 | Beta3-adrenoceptor agonist stimulation of the Na ⁺ , K ⁺ -pump in rat skeletal muscle is mediated by beta2- rather than beta3-adrenoceptors. <i>British Journal of Pharmacology</i> , 2006 , 149, 635-46 | 8.6 | 8 |
| 2 | N-acetylcysteine attenuates the decline in muscle Na ⁺ ,K ⁺ -pump activity and delays fatigue during prolonged exercise in humans. <i>Journal of Physiology</i> , 2006 , 576, 279-88 | 3.9 | 191 |
| 1 | Intense exercise up-regulates Na ⁺ ,K ⁺ -ATPase isoform mRNA, but not protein expression in human skeletal muscle. <i>Journal of Physiology</i> , 2004 , 556, 507-19 | 3.9 | 55 |

