

# Editorial: Exercise as a Countermeasure to Human Aging

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The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest

### *Author contribution statement*

BE wrote the first draft. LH, DH and MB critically reviewed, and all authors approved the final version of this editorial

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### *Contribution to the field*

The world's population is rapidly aging and as the average age of the population continues to increase research needs to turn towards non-communicable diseases of ageing. This editorial discusses contributions to the research topic 'Exercise as a Countermeasure to Human Aging', and the positive contribution these make both individually and when considered as a group of publications. Both exercise and physical activity are capable of offsetting many of the physiological changes seen with ageing, and the collection of publications described in this editorial all make contributions to our understanding of the interactions between exercise, activity and the physiology of human ageing.

# 1 Exercise as a Countermeasure to Human Aging

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4 Unlike many of the branches of natural sciences, there are few true 'laws' of physiology. However,  
5 there are intrinsic theories about which we are reasonably certain. For example, it is a reasonable  
6 statement that exercise and physical activity, in all their forms, typically have positive effects on health  
7 and wider physiological function via multiple complex and interacting mechanisms (that we have not  
8 yet completely defined). Alternatively, the continuous process of human aging in the adult involves a  
9 gradual decline of physiological function across most tissues and systems, again in a complex and  
10 intertwining manner. At a point where the average age of humanity is greater than it has ever been,  
11 and is continuing to increase, we considered it timely to examine the crossover between these two  
12 interacting fields of physiology. Indeed, where past successes in physiology research have emerged  
13 from research on transmittable diseases, vaccinations and preventive medicines, our current  
14 approaches must now focus on non-transmittable disorders, including frailty syndromes, sarcopenia  
15 and chronic conditions that associate with aging, including heart disease, neuro-cognitive disorders  
16 and diabetes.

17

18 When we made this call for submissions, we did not expect the volume of responses we received. In  
19 these papers we presented a collection of over 30 articles that covered the interplay between exercise  
20 and aging, utilising approaches that spanned molecular, physiological, and population scale  
21 approaches, in both healthy older populations and certain disease subsets, and spanned three  
22 *Frontiers* journals (*Frontiers in Physiology*, *Frontiers in Sports and Active Living*, and *Frontiers in Aging*  
23 *Neuroscience*). It is a pleasure to note this range of fields and methodological approaches that authors  
24 have used.

25

26 It has long been known that exercise benefits human function, and that this effect may promote good  
27 health into older age. Philostratus (c. 170 - 250) wrote of individuals who exercised into older age that  
28 "They were healthy and did not get sick easily. They stayed youthful into old age, and competed in  
29 many Olympics, some in eight and others in nine" (Gymnasticus 44). Several papers in this collection  
30 examined classical exercise physiology approaches of a short-term training programme over weeks-  
31 to-months. In this vein [Kirk et al. \(2019\)](#) gave preliminary results from the LHU-SAT trial, examining 16  
32 weeks of training with or without protein supplementation in healthy over 60-year-old participants.  
33 While both groups improved with training, results suggested the protein supplementation group did  
34 not improve to a greater degree than the no protein group. However, compliance to protein  
35 supplementation beverages in this population continued to be low, an area that may need attention.  
36 In line with these results, positive outcomes from classical exercise physiology training interventions  
37 were seen by [Walker et al. \(2018\)](#) who reported on improved intermuscular coherence, [Gavin et al.](#)  
38 [\(2019\)](#) who noted resistance training improved stair climbing biomechanics in older individuals, [Tam](#)  
39 [et al. \(2018\)](#) reported on resistance training improving exercise economy, and [Saeidi et al. \(2019\)](#)  
40 findings that a proposed antioxidant altered resistance training-induced changes in circulating  
41 [adipokines](#) in postmenopausal women. Two exercise physiology interventional papers of note include  
42 [Franchi et al. \(2019\)](#), who used a novel trampoline plyometric training model in a safe and highly  
43 effective way, and [Jabbour and Majed \(2018\)](#) with the important observation that the widely used  
44 ratings of perceived exertion (RPE) scale over-estimated exercise intensity in sedentary older adults.

45 In meta-analyses reviewing exercise changes from short-term training interventions, endurance  
46 exercise decreased pro-inflammatory cytokines concentrations ([Zheng et al., 2019](#)), yet  
47 counterintuitively testosterone was not improved following training studies in older men ([Hayes and  
48 Elliott, 2018](#)), suggesting resistance training-induced benefits were not via circulating testosterone  
49 concentrations.

50

51 Updating us on recent advances in targeting mitochondria to offset sarcopenia, [Coen et al. \(2018\)](#)  
52 reviewed exercise and mitochondrial health for successful aging, reminding the reader that exercise  
53 is (for now) the only effective option for treatment of sarcopenia. Linking well to this review, [Ubaida-  
54 Mohien et al. \(2019\)](#) reported on a proteomic analysis of muscle biopsies from 60 individuals spanning  
55 20 – 87 years of age, and reported physical activity associated with alterations in proteins governing  
56 mitochondria energetics, muscle function, gene health, immunity and senescence, and these changes  
57 typically opposed those seen with aging. Mirroring these results, ambulatory older individuals  
58 presented a preservation in portions of the myostatin and IGF-I signalling pathways, as well as myocyte  
59 structures, that wheelchair bound older individuals did not show ([Naro et al., 2019](#)). Differing  
60 endurance exercise stimuli improved markers of t-cell senescence ([Philippe et al., 2019](#)), while in older  
61 rats, muscle protein synthesis responses were blunted relative to younger animals ([West et al., 2019](#)).  
62 All these results point to an environment that is capable of positively responding to anabolic stimuli,  
63 but perhaps not as well as younger muscle tissue, as well as a need for research to separate effects of  
64 aging and inactivity.

65

66 From a population health point of view, increased lifelong activity, not just short-term exercise  
67 interventions, are needed. Thus, there has been much recent interest in examining highly trained  
68 masters athletes, as a physiological model of successful aging (Pollock et al., 2015, Elliott et al., 2017).  
69 This special edition included five reports on lifelong exerciser cohorts. [Mancini et al. \(2019\)](#) compared  
70 lifelong football players with age matched controls, noting a positive influence of lifelong exercise on  
71 markers of auto-lysosomal and proteasomal-mediated processes, while [Piasecki et al. \(2019a\)](#) noted  
72 an interesting compensatory mechanism whereby power trained older adults showed increased  
73 motor unit size, possibly to compensate for decreased motor unit number. In older females,  
74 osteoporosis is often seen, however [Onambele-Pearson et al. \(2019\)](#) observes that simple mechanical  
75 loading is not sufficient to explain bone density, and that fuller measures of activity and inactivity  
76 should be considered. In masters athletes who were grouped as 'early' or 'late' starters to masters  
77 athletics (either lifelong training history or beginning after 50 years of age), [Piasecki et al \(2019b\)](#)  
78 reported no major differences in body composition or bone density between these early and late  
79 starters, but both groups reliably demonstrated a healthier phenotype vs inactive controls. Finally, it  
80 was of interest to note positive emotional and cognitive effects of lifelong Tai Chi participation relative  
81 to an age-matched control group, which was paired with resting-state fMRI connectively differences  
82 ([Liu et al., 2018](#)). It can be seen that lifelong activity promotes multiple physiological benefits in an  
83 aging population.

84

85 At one end of the population size spectrum, [Knechtle et al. \(2018\)](#) presented a case study on  
86 physiological responses in a 95-year-old masters athlete during a 12 hour ultra-marathon event. At  
87 the other end are population scale studies. It is of interest to note differences in the association  
88 between physical activity, as measured by accelerometry, and relative telomere lengths, with

89 positive associations seen in men but not in woman, across a population of 700 older participants  
90 [\(Stenback et al., 2019\)](#). By analysing records of ~27,000 track and field athletes, [Ganse et al. \(2018\)](#)  
91 observed decreases across maximal power, strength and endurance records throughout adult  
92 lifespan. Further, these declines in performance accelerated post 70 years of age, an observation that  
93 was seen in grip strength in the general population (Dodds et al., 2014), and occurred despite high  
94 levels of physical activity. These results, in combination, suggested that muscle function loss with age  
95 is not only inactivity-induced but has an intrinsic component.

96

97 As aging is associated with an increased risk of cardiovascular disease, diabetes and certain types of  
98 cancers, and chronic exercise associates with reduced rates of such disorders, it is important to  
99 examine exercise in such older populations with such conditions. Indeed, regular exercise training of  
100 any type improved quality of life, aerobic capacity and heart function in older heart failure patients  
101 [\(Slimani et al., 2018\)](#). [Mcleod et al. \(2019\)](#) argued for alterations in guidelines for exercise in the  
102 prevention of chronic disorders, promoting the role of resistance training in preventive medicine,  
103 interesting reading when paired with the [Campbell et al. \(2019\)](#) meta-analysis which observed  
104 insufficient evidence to recommend aerobic exercise for vascular function improvement in older  
105 sedentary adults. In rats, experimental data suggested that prior exercise training improved  
106 survivability from experimental coronary artery occlusion [\(Veiga et al., 2019\)](#), providing us humans  
107 with more motivation for maintaining lifelong exercise. This was reinforced by a cohort study of ~3,700  
108 individuals, where both physical activity and sedentary time both independently predicted mortality  
109 rates associated with pro-inflammatory conditions [\(Cabanas-Sanchez et al., 2018\)](#). Other findings  
110 suggested the improvements in post-exercise reaction time were not different between hypertensive  
111 and non-hypertensive patients [\(Lefferts et al., 2019\)](#), and the interesting observation that structural  
112 differences in skeletal muscle may underlie difference in stretch shorten cycle between COPD patients  
113 and healthy age-matched controls [\(Navarro-Cruz et al., 2019\)](#). These results reinforce the recent  
114 American Medical Association's guidelines promoting exercise wherever possible in chronic conditions  
115 (Piercy et al., 2018).

116

117 Historically, physiology research has primarily utilized the 'healthy young male' population, thus we  
118 are pleased to note that 14 of the 21 primary experimental papers presented here in human  
119 participants included male and female groups, while one specifically examined post-menopausal  
120 changes in women. Likewise, we feel the papers presented here give valuable insight concerning the  
121 range of ageing physiology, in a continuous rather than dichotomous manner. For example, [Knechtle](#)  
122 [et al. \(2018\)](#) concerned a 95-year-old masters athlete, considered the 'oldest old', whereas some  
123 papers [\(Hayes and Elliott, 2018\)](#) had a minimum age of 60, considered the 'young old'. Moreover,  
124 several investigations utilized a young comparison group or a cross sectional design, which permitted  
125 authors to study life course aging utilizing multiple research designs.

126

127 Both physical activity and structured exercise are near-uniformly positive for human longevity and  
128 well-being by multiple, complex physiological mechanisms and pathways that help maintain health,  
129 independence and quality of life. Indeed, the complexity of the aging process and the role of exercise  
130 in aging physiology were well represented by the diversity of experimental approaches witnessed in  
131 this research topic. Combined, the results of these investigations suggested that exercise and activity  
132 can offset decreases in human function that we consider 'inevitable aspects of aging' but cannot

133 prevent them completely. Our understanding of how and why exercise and activity promote healthy  
134 aging, and indeed the basic physiology of the aging process, is currently incomplete. It is our aim that  
135 this research topic makes a small contribution to the understanding of this complex field.

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